

**IN THE UNITED STATES DISTRICT COURT  
FOR THE MIDDLE DISTRICT OF TENNESSEE**

BONGO PRODUCTIONS, LLC, ROBERT  
BEINSTEIN, SANCTUARY PERFORMING  
ARTS, LLC, and KYE SAYERS,

Plaintiffs,

v.

CIVIL ACTION  
CASE NO. 3:21-cv-00490  
JUDGE TRAUGER

CARTER LAWRENCE, Tennessee State Fire  
Marshal, in his official capacity,  
CHRISTOPHER BAINBRIDGE, Director of  
Codes Enforcement, in his official capacity,  
GLENN R. FUNK, District Attorney General for  
the 20<sup>th</sup> Judicial District, in his official capacity,  
and NEAL PINKSTON, District Attorney  
General for the 11<sup>th</sup> Judicial District, in his  
official capacity,

Defendants.

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**DEFENDANTS' RESPONSE IN OPPOSITION TO PLAINTIFFS' MOTION FOR  
SUMMARY JUDGMENT**

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Pursuant to Rule 56 of the Federal Rules of Civil Procedure Defendants Carter Lawrence, Christopher Bainbridge, Glenn R. Funk, and Neal Pinkston, in their official capacities only, hereby respond in opposition to Plaintiffs'<sup>1</sup> Motion for Summary Judgment. (DE 35).

**BACKGROUND**

On March 29, 2021, and April 29, 2021, the General Assembly passed House Bill 1182/Senate Bill 1224 by overwhelming majorities in both Houses. Governor Lee signed House Bill 1182 into law on May 17, 2021, as Public Chapter 453 ("the Act"). (Ex. A.) The Act furthers the State's interests in informing persons patronizing buildings open to the public of the building

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<sup>1</sup> Plaintiffs Kye Sayers and Sanctuary Performing Arts, LLC, are no longer participating in this matter.

operator’s bathroom policy should it deviate from any existing bathroom signage designating a bathroom as available only to persons of a specific biological sex. Plaintiffs seek to enjoin the enforcement of the entire Act, which took effect on July 1, 2021.

**A. The Contested Statute.**

As pertinent here, the Act applies to all businesses and entities within the State that are open to the public and, “as a matter of formal or informal policy, allow[] a member of either biological sex to use any public restroom within the building or facility.” (Ex. A. Act, § 1(a).)

“Public restroom,” as defined by the Act, includes restrooms, locker rooms, shower facilities, dressing areas, and any similar facility that is “open to the general public,” “designated for a specific biological sex,” and constitutes a “facility or area where a person would have a reasonable expectation of privacy.” (Ex. A. Act, §§ 1(d)(2)(A)(i)-(iii).) However, the term “[p]ublic restroom” does not include single-occupancy restrooms or family restrooms “intended for use by either biological sex.” (Ex. A. Act § 1(d)(2)(B).)

Business and entities that are subject to the Act must post signage that is easily visible to people entering the restroom<sup>2</sup> and that informs the public that “THIS FACILITY MAINTAINS A POLICY OF ALLOWING THE USE OF RESTROOMS BY EITHER BIOLOGICAL SEX, REGARDLESS OF THE DESIGNATION ON THE RESTROOM.” (Ex. A. Act, § 1(b)(3).)

In short, the Act requires that any entity with multi-user, sex-designated public restrooms—and a policy of allowing all users to use either restroom, regardless of their biological sex—to post signage reflecting that policy on or near the restroom entrances. Qualifying businesses and entities that do not comply with the Act’s requirements must have received notice of their noncompliance

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<sup>2</sup> To that end, the Act also specifically describes the dimensions, coloring, and location of the required signage. (Ex. A. Act, §§ 1(b)(1)-(5).)

at least 30 days before any action is taken against them. (Ex. A. Act, § 1(c).)

## **B. Case History**

Plaintiffs—a Tennessee business and its owner filed a Complaint on June 25, 2021, challenging the constitutionality of the Act. (*See generally* DE 1; DE 1, PageID# 1.) Plaintiffs allege that they do not wish to post the signage required by the Act because they perceive that the signage required—which mirrors their bathroom-usage policies—is controversial and stigmatizing. (DE 1, PageID# 2.) They allege that by requiring them to post this signage, the Act compels speech in violation of the First Amendment to the United States Constitution. (DE 1, PageID# 18.) Contemporaneously with the filing of their Complaint, Plaintiffs moved for a preliminary injunction “enjoining enforcement of H.B. 1182/S.B. 1224, 112th Gen. Assemb., 1st Reg. Sess. (Tenn. 2021).” (*See generally* DE 6.)

Shortly thereafter, this Court granted Plaintiffs’ Motion for Preliminary Injunction and enjoined Defendants from enforcing the Act. (DE 22; DE 23). On January 31, 2022, Plaintiffs filed the instant Motion for Summary Judgment. (DE 35).

## **LEGAL STANDARD**

“The court shall grant summary judgment if the movant shows that there is no genuine dispute as to any material fact and the movant is entitled to judgment as a matter of law.” Fed. R. Civ. P. 56(a); *see also Johnson v. Karnes*, 398 F.3d 868, 873 (6th Cir. 2005). The moving party must “show that the non-moving party has failed to establish an essential element of his case upon which he would bear the ultimate burden of proof at trial.” *Guarino v. Brookfield Twp. Trustees*, 980 F.2d 399, 403 (6th Cir. 1992). After the movant makes a properly supported motion, the burden shifts to the non-moving party to demonstrate the existence of material facts in dispute. An opposing party may not simply rely on its pleadings; rather, it must “produce evidence that

results in a conflict of material fact to be resolved by a jury.” *Cox v. Ky. Dep’t of Transp.*, 53 F.3d 146, 150 (6th Cir. 1995). When the moving party has carried this burden, the nonmoving party must set forth specific facts showing that there is a genuine issue for trial. *Moldowan v. City of Warren*, 578 F.3d 351, 374 (6th Cir. 2009). In evaluating a motion for summary judgment, the Court must draw all inferences in the light most favorable to the nonmoving party. *Adickes v. S.H. Kress & Co.*, 398 U.S. 144, 158-59 (1970); *Reeves v. Sanderson Plumbing Prods., Inc.*, 530 U.S. 133, 150 (2000). However, the existence of a mere scintilla of evidence in support of the nonmoving party’s position will not be sufficient; there must be evidence on which the jury reasonably could find for the nonmoving party. *Anderson v. Liberty Lobby, Inc.*, 477 U.S. 242, 251 (1986); *Matsushita Elec. Indus. Co. v. Zenith Radio Corp.*, 475 U.S. 574, 587-88 (1986) (finding reliance upon mere allegations, conjecture, or implausible inferences to be insufficient to survive summary judgment); *Copeland v. Machulis*, 57 F.3d 476, 479 (6th Cir. 1995).

## **ARGUMENT**

Here, summary judgment is inappropriate. As a threshold matter, this Court lacks subject matter jurisdiction for two reasons: 1) Plaintiffs have failed to demonstrate standing because the enforcement of the Act that gives rise to their alleged injury-in-fact is not imminent or even likely; and 2) Plaintiffs’ First Amendment claim is not ripe as the effect of any potential enforcement is uncertain. Further, even if subject matter jurisdiction were present, the Act satisfies constitutional scrutiny. Plaintiffs’ Motion for Summary Judgment should therefore be denied.

### **I. The Court Lacks Subject Matter Jurisdiction to Consider Plaintiffs’ Claims.**

#### **A.. The Plaintiffs Lack Standing Because their Alleged Injury-in-Fact is not Imminent.**

Article III of the United States Constitution limits the jurisdiction of federal courts to cases and controversies. U.S. Const. art. 3, § 2. The doctrine of standing is “an essential and unchanging

part of the case-or-controversy requirement,” and defines the boundaries of jurisdiction under Article III by “identifying those disputes which are appropriately resolved through the judicial process.” *Lujan v. Defs. of Wildlife*, 504 U.S. 555, 560 (1992). Thus, as a threshold matter, federal courts are “under an independent obligation to examine their own jurisdiction, and standing is perhaps the most important of the jurisdictional doctrines” that a plaintiff must satisfy. *FW/PBS, Inc. v. City of Dallas*, 493 U.S. 215, 231 (1990) (internal quotations and citation omitted); see *Copas v. Lee*, 396 F.Supp.3d 777, 786 (M.D. Tenn. 2019) (“Standing is a ‘threshold determinant[ ] of the propriety of judicial intervention.’” (quoting *Warth v. Seldin*, 422 U.S. 490, 518 (1975))).

In determining whether a plaintiff has standing, courts consider whether the plaintiff has alleged “an ‘injury in fact’ that is ‘fairly traceable to the challenged action of the defendant’ and is capable of being ‘redressed’ by the court.” *McKay v. Federspiel*, 823 F.3d 862, 867 (6th Cir. 2016) (quoting *Lujan*, 504 U.S. at 560-61). And the plaintiff bears the burden of establishing the presence of all three elements. *Id.* To establish “injury-in-fact,” the plaintiff must show he suffered a *concrete* injury that is “actual or imminent, not conjectural or hypothetical.” *Lujan*, 504 U.S. at 560. Moreover, the injury alleged must be particularized to the plaintiff—“not a generalized grievance.” *Allen v. Wright*, 468 U.S. 737, 751 (1984).

Here, Plaintiffs challenge the Act under the First Amendment yet they do not allege that the Defendants have actually enforced or will enforce any provisions of the Act against them. “In a pre-enforcement challenge, whether the plaintiff has standing to sue often turns upon whether he can demonstrate an ‘injury in fact’ before the state has actually commenced an enforcement proceeding against him.” *Kiser v. Reitz*, 765 F.3d 601, 607 (6th Cir. 2014). Although Plaintiffs are not required to subject themselves to actual arrest or prosecution as a prerequisite, pre-enforcement challenges typically require “the threatened injury [to be] certainly impending or

[that] there is a substantial risk that the harm will occur.” *Susan B. Anthony List v. Driehaus*, 573 U.S. 149, 158 (2014) (quoting *Clapper v. Amnesty Int’l USA*, 568 U.S. 398, 414, n. 5 (2013)). Specifically, the Supreme Court has permitted satisfaction of the “injury-in-fact” requirement in the pre-enforcement context when plaintiffs allege “an intention to engage in a course of conduct arguably affected with a constitutional interest, but proscribed by statute, and there exists a credible threat of prosecution thereunder.” *Babbitt v. Farm Workers*, 442 U.S. 289, 298 (1979).

Under the Act, “[i]f an entity or business is notified that it is not in compliance with this section, the entity or business has thirty (30) days in which to comply before any action is taken against the entity or business.” (Ex. A. Act, § 1(c).) Plaintiffs in this case do not want to display the government-mandated warning notice required by the Act. (DE 1, PageID# 2.) But as Plaintiffs have neither received a threat of enforcement nor a notice pursuant to the statute, there mere existence of the statute alone does not amount to “a credible threat of prosecution” under the Act.

In *McKay*, the plaintiff brought a Section 1983 claim against public officials charged with enforcing a state court’s administrative order prohibiting recording devices in the courtroom. 823 F.3d at 864-65. There, the plaintiff did not allege that he requested or was denied permission to use a recording device, nor that he attempted to enter the courthouse with such a device. Instead, he merely alleged that he did not wish to be subject to contempt, confiscation of a device, a fine, or jail time pursuant to the order. *Id.* at 865-66. In holding that the plaintiff lacked standing, the Sixth Circuit considered a series of factors that, in conjunction with a plaintiff’s allegation of a subjective First Amendment chill, could establish injury-in-fact:

- (1) a history of past enforcement against the plaintiffs or others;
- (2) enforcement warning letters sent to the plaintiffs regarding their specific conduct; and/or
- (3) an attribute of the challenged statute that makes enforcement easier or more likely, such as a provision allowing any member of the public to initiate an enforcement

action...[and] a defendant's refusal to disavow enforcement of the challenged statute against a particular plaintiff.

*Id.* at 869 (internal citations omitted). Indeed, courts considering credible threats of prosecution in the context of a pre-enforcement First Amendment challenge consistently look to some indication of a threat beyond the simple possibility of enforcement. *Id.*; *see also Kiser*, 765 F.3d at 609 (finding a credible threat where the plaintiff received two letters from the defendant warning him that he was in violation of the regulation at issue); *Susan B. Anthony List*, 573 U.S. at 164 (2014) (noting that the plaintiff was previously prosecuted by the defendant for the same sort of speech); *Plunderbund Media, L.L.C v. DeWine*, 753 Fed. App'x. 362, 366-72 (6th Cir. 2018) (finding no credible threat where there was no history of enforcement against defendants or the kind of speech at issue, nothing making the statute easier or more likely to enforce, and no evidence of an intention to enforce).

Here, Plaintiffs do not allege facts that satisfy any of the *McKay* factors, and therefore do not allege a credible threat of prosecution. (*See generally*, DE 1.) Indeed, they do not allege receipt of any warning letters, any attribute of the Act that would make enforcement easier or more likely,<sup>3</sup> or that Defendants have refused to disavow enforcement against them. Just the opposite is true: Defendant District Attorney General Funk has made public statements expressing his intention to disavow enforcement of the Act in Davidson County, where Bongo's business is located. Kimberlee Kruesi, *Nashville DA won't enforce new bathroom sign law*, Associated Press (May 24, 2021), <https://apnews.com/article/nashville-laws-government-and-politics-50412b91ca33cc45c426a9b5a89b1133> (Ex. B).

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<sup>3</sup> The Act's 30-day notice requirement before Defendants could subject Plaintiffs to penalties under the Act make its enforcement *more* difficult and *less* likely.

Moreover, the Act provides a 30-day cure period for businesses before any action can be taken. (Ex. A. Act, § 1(c).) Yet Plaintiffs do not allege receipt of such notice. In other words, despite Plaintiffs' express desire not to comply with the Act, they cannot be subject to any action for at least 30 days, starting from notice of noncompliance. Thus, because Plaintiffs fail to allege a credible threat of prosecution under the Act, they fail to properly allege injury-in-fact and lack standing to bring this claim.

Lastly, because Plaintiffs do not allege enforcement of the challenged statute by Defendants, there is no injury for the court to redress in this case. In *California v. Texas*, 593 U.S. \_\_\_, No. 19-840, 2021 WL 2459255 (June 17, 2021), the Supreme Court considered the standing of several individual plaintiffs to challenge the Affordable Care Act's minimum-coverage requirement. The Court held that those plaintiffs lacked standing because "no unlawful Government action "fairly traceable" to [the challenged statute] caused the plaintiffs'... harm." *Id.* at \*5. As the Court pointed out, it has consistently required plaintiffs "to assert an injury that is the result of a statute's actual or threatened enforcement, whether today or in the future." *Id.* The Court further noted that issuing an injunction in a case where plaintiffs were not harmed by actual or threatened injury would ultimately amount to "an advisory opinion without the possibility of any judicial relief." *Id.* at \*6 (quoting *Los Angeles v. Lyons*, 461 U.S. 95, 129, (1983) (Marshall, J., dissenting)); *see also Carney v. Adams*, 141 S. Ct. 493, 498 (2020) (Article III "require[s] that a case embody a genuine, live dispute between adverse parties, thereby preventing the federal courts from issuing advisory opinions"). Here, Plaintiffs simply cannot show a relationship between the judicial relief requested and the "injury" they allege because they have not suffered an injury. In effect, Plaintiffs are requesting an advisory opinion under the guise of injunctive relief.

**B. Plaintiffs' Claims are Not Ripe.**

Like the doctrine of standing, the ripeness doctrine is rooted in Article III limitations on federal-court jurisdiction. However, ripeness distinctly aims at ensuring cases or controversies filed in federal court are timely to help the court “avoid[]...premature adjudication.” *Abbott Labs. v. Gardner*, 387 U.S. 136, 148-49 (1967). To show that the claim is ripe for review, a plaintiff must allege more than some future acts or events that “may not occur as anticipated, or at all.” *Nat’l Rifle Ass’n of Am. v. Magaw*, 132 F.3d 272, 284 (6th Cir. 1997).

Determining whether a claim is ripe for review requires a court to consider both “the fitness of the issue for judicial decision and the hardship to the parties of withholding court consideration.” *Abbott Labs*, 387, U.S. at 149. Specifically, the Sixth Circuit considers three factors in determining whether a claim is ripe: “(1) the likelihood that the harm alleged will ever come to pass; (2) whether the factual record is sufficiently developed to allow for adjudication; and (3) hardship to the parties if judicial review is denied.” *Norton v. Ashcroft*, 298 F.3d 547, 554 (6th Cir. 2002); *see also Ammex, Inc. v. Cox*, 351 F.3d 697, 706 (6th Cir. 2003). In the context of a claim that relies on the threat of injury, instead of an actual one, the standing and ripeness doctrines can be difficult to distinguish. *Airline Pros. Ass’n of Int’l Bhd. of Teamsters, Local Union No. 1224, AFL-CIO v. Airborne, Inc.*, 332 F.3d 983, 988 (6th Cir. 2003). For example, “[a] threatened or imminent injury may satisfy standing’s injury-in-fact requirement, yet the claim may still be unripe if the issues are not fit for judicial review, perhaps because future events may greatly affect the outcome of the litigation and the cost of waiting is not particularly severe.” *Id.*

Here, as discussed above at I.A., Plaintiffs fail to allege an injury beyond potential future events that may not occur as they expect or may not occur at all. Plaintiffs do not know that the Act will be enforced against them, and even assuming it were enforced against them, they can

predict neither how it would be enforced nor the effect of its enforcement. This case therefore lacks the factual development necessary for this Court to properly adjudicate Plaintiffs' claim.

Thus, due to a lack of standing and ripeness that deprives this Court of subject matter jurisdiction, summary judgment is inappropriate.

## **II. The Act is Constitutional.**

Even if the Court finds that it possesses subject matter jurisdiction, the Court should decline to grant summary judgment as the Act satisfies constitutional scrutiny.

Plaintiffs raise only one claim challenging the constitutionality of the Act: that the Act violates their First Amendment rights by compelling them, on pain of criminal penalty, to communicate a misleading and controversial government-mandated message that they would not otherwise display. (DE 1, PageID# 2.) Plaintiffs urge this Court to subject the Act to strict scrutiny. Plaintiffs, though, have conceded that, as businesses open to the public, they are subject to regulation, including regulations that require them to post certain signage. (DE 7, PageID# 51). Plaintiffs also conceded that “notices [that] communicate purely factual and non-controversial speech [] do[] not offend the First Amendment.” (DE 7, PageID# 51.)

Plaintiffs' prior concessions confirm that strict scrutiny is inappropriate. It is well settled, of course, that a State may not “constitutionally require an individual to participate in the dissemination of an ideological message by displaying it on his private property in a manner and for the express purpose that it be observed and read by the public.” *Wooley v. Maynard*, 430 U.S. 705, 713 (1977).

But while the State has “no power to restrict expression because of its message, its ideas, its subject matter, or its content,” “[u]nder the First Amendment,” it may still “regulate certain aspects of speech.” *Thomas v. Bright*, 937 F.3d 721, 729 (6th Cir. 2019) (quoting *Police Dep't of*

*City of Chi. v. Mosely*, 408 U.S. 92, 95 (1972)). And when a statute does not regulate or compel expressive or ideological speech, strict scrutiny is not the applicable constitutional test. Indeed, the federal appellate courts have repeatedly declined to apply strict scrutiny to non-expressive, non-ideological disclosure requirements in the face of First Amendment challenges. *See, e.g., Zauderer v. Off. of Disciplinary Couns. of Sup. Ct. of Ohio*, 471 U.S. 626, 651 (1985) (declining to apply strict scrutiny to commercial speech); *see also Nat'l Electric Mfrs. Ass'n v. Sorrell*, 272 F.3d 104 (2d Cir. 2001) (upholding a labeling requirement containing purely factual and uncontroversial speech); *Conn. Bar Ass'n v. United States*, 620 F.3d 81 (2d Cir. 2010) (subjecting disclosure requirements to rational basis review); *N. Y. State Rest. Ass'n v. N. Y. C. Bd. of Health*, 556 F.3d 114 (2d Cir. 2009) (applying rational basis review to government mandated caloric disclosure requirements); *Scope Pictures, of Mo, Inc. v. City of Kan. City*, 140 F.3d 1201 (8th Cir. 1998) (upholding a signage requirement regarding venereal disease where the signage conveyed no political or ideological message); *United States v. Sindel*, 53 F.3d 874, 878 (8th Cir. 1995) (“First Amendment protection against compelled speech. . . has only been found in the context of governmental compulsion to disseminate a particular political or ideological message.”).

The signage required by the Act is neither ideological nor expressive speech. Ideological speech is speech which conveys a “point of view.” *Wooley*, 430 U.S. at 715. An accurate statement of fact—such as the bathroom-usage policy chosen by Plaintiffs—does not communicate an ideological or political message. *See, e.g., Dutchess/Putnam Rest. & Tavern Ass'n, Inc. v. Putnam Cnty. Dep't of Health*, 178 F.Supp.2d 396, 406 (S.D.N.Y. 2001) (holding that state-

mandated signs informing patrons of the risk of smoking did not constitute ideological speech and therefore did not violate the First Amendment).

Plaintiffs either misapprehend or have mischaracterized what the Act does. It does not compel that a business adopt a particular bathroom policy. Nor does it require exclusion of transgendered persons from the bathroom of their choice. Nor does it stigmatize businesses or patrons. Instead, the law requires only one thing: if a business's bathroom-use policy is different from its existing bathroom signage, it needs to inform its patrons as such. (Ex. A. Act.). Plaintiffs have imagined an idiosyncratic, hidden undertone to the signage that is not reflected in the Act's plain language. But Plaintiffs' projections cannot transform a neutral statute requiring a simple truthful statement of fact—aimed at informing the public without compelling the business or entity's ultimate choice of bathroom-usage policy—into expressive speech.

Plaintiffs' evidence in support of their idiosyncratic interpretation of the Act is a disputed fact rendering summary judgment inappropriate. In their statement of undisputed material facts, Plaintiffs assert that the term "biological sex" is a recent one without a uniform definition. Not so. The phrase "biological sex" is a common phrase that is also used throughout scientific literature and possesses a fixed and uniform definition. *See* Am. Psychiatric Assoc., *The Diagnostic and Statistical Manual of Mental Disorders*, 15 (5th ed. 2013) (Ex. C); Riittakerttu Kaltiala-Heino et al., *Gender dysphoria in adolescence: current perspectives*, *ADOLESCENT HEALTH, MEDICINE AND THERAPEUTICS* 2018: 9, 21, 21 (2018) (Ex. D); L.A. Walter and A.J. McGregor, *Sex- and Gender-specific Observations and Implications for COVID-19*, *WEST J EMERG MED.* 21(3): 507-509 (2020) (Ex. E); E.P. Scully et al., *Considering how biological sex impacts immune responses and COVID-19 outcomes*, *NAT REV IMMUNOL* 20, 442-447 (2020) (Ex. F); S.L. Klein et al., *Biological sex impacts COVID-19 outcomes*, *PLOS PATHOG* 16:6 (2020) (Ex. G). Further, "biological sex"

has the same meaning as the longstanding and broadly accepted definition of “sex.” *See, e.g., Sex*, New Oxford American Dictionary (3d ed. 2010) (“either of the two main categories (male and female) into which humans and many other living things are divided on the basis of their reproductive functions”).

Plaintiffs also argue that the term “biological sex” is stigmatizing and inherently discriminatory. Again, not so. The term “biological sex” is a neutral term that conveys no stigma or viewpoint. Nor is the term necessarily indicative of attempts to limit or eliminate the legal recognition, protection, and rights of transgender people. *See; Doe 2 v. Shanahan*, 917 F.3d 694, 698 (D.C. Cir. 2019); *Able v. United States*, 88 F.3d 1280, 1286 (2d Cir. 1996); *Grimm v. Gloucester Cnty. Sch. Bd.*, 972 F.3d 586, 614 (4th Cir. 2020); *Parents for Priv. v. Barr*, 949 F.3d 1210, 1217 (9th Cir. 2020); *Doe by and through Doe v. Boyertown Area Sch. Dist.*, 897 F.3d 518, 529 (3d Cir. 2018); *Hively v. Ivy Tech Cmty. Coll. of Ind.*, 853 F.3d 339, 347 (7th Cir. 2017); *Cruzan v. Special Sch. Dist, No. 1*, 294 F.3d 981, 983 (8th Cir. 2002); *Jackson v. Valdez*, --- Fed.App’x ---, 2021 WL 1990788, at \*5 (5th Cir. 2021); *Etsitty v. Utah Transit Auth.*, 502 F.3d 1215, 1225 (10th Cir. 2007); *E.E.O.C. v. R.G. & G.R. Harris Funeral Homes, Inc.*, 884 F.3d 560, 578 (6th Cir. 2018) (affirmed by *Bostock v. Clayton Cnty., Ga.*, 140 S.Ct. 1731 (2020)); *Doe v. Hamilton Cnty. Bd. of Educ.*, 329 F.Supp.3d 543, 580 (E.D. Tenn. 2018); *Farmer v. Brennan*, 511 U.S. 825, 829 (1994); *Bostock*, 140 S.Ct. at 1752. The challenged Act speaks for itself, and like the cited judicial opinions, uses the term “biological sex” objectively and neutrally. *See also* Taylor Depo. at 79: 17-22 (“I think the intention of both of those terms [“biological sex” and “sex assigned at birth”] is the same”). (Attachment A).

Plaintiffs’ preferred term “gender identity” is the newcomer. “Gender identity” is a term popularized by Robert Stoller, a UCLA psychoanalyst. According to Stoller, “sex was biological

but gender was social.” David Haig, *The Inexorable Rise of Gender and the Decline of Sex: Social Change in Academic Titles, 1945-2001*, Archives of Sexual Behavior, Apr. 2004, at 93 (Ex. H). The term “gender”—previously a grammatical term only—was itself introduced into scientific discourse in the 1950s by John Money, a psychologist at Johns Hopkins University. Joanne Meyerowitz, *A History of “Gender,”* 113 *The American Historical Review* 1346, 1354 (2008) (Ex. I). Research has not established “a strong biological basis” connecting transgender individuals’ gender identity with the biological reality of how their bodies’ reproductive systems are organized. Accordingly, many purported gender identities are unmoored from the male/female binary. *Cf. United States v. Varner*, 948 F.3d 250, 256-57 (5th Cir. 2020). Thus, gender identity plays no role in determining sex. *See* Bachtrog et al., *Sex Determination, Why So Many Ways of Doing It?* PLoS Biol, 2014 Jul; 12(7) (Ex. J).

Nor is their expert testimony on the alleged stigmatizing effect of the term “biological sex” undisputed. *See* Taylor Depo. 108; 8-13; 109; 24-25; 110; 1-7 (Attachment A). Plaintiffs’ own expert testified that the term “biological sex” does not, in her experience, worsen gender dysphoria. *Id.* And Plaintiffs’ attempts to call the term “biological sex” into question using rare syndromes likewise fails. Humans with disorders of sexual development still have a sex. For example, humans with Klinefelter syndrome (47XXY) are still men, and humans with Turner syndrome (45XO) are still women. Despite departures from the normal development expected for humans with 46XY or 46XX chromosomes, those with Klinefelter syndrome still have a male reproductive system while those with Turner system still have a female reproductive system. These two chromosomal disorders occur in a small proportion of the population. *See* NIH, *Klinefelter Syndrome*, Eunice Kennedy Shriver National Institute of Child Health and Human Development -

NICHD (nih.gov) (Ex. K); NIH, *Turner Syndrome*, Eunice Kennedy Shriver National Institute of Child Health and Human Development - NICHD (nih.gov) (Ex. L).

Nor do Plaintiffs demonstrate that the Act's required language would be misleading or untruthful. By the Act's plain language, Plaintiffs need only post the required signage if they *agree* with the language set forth by the Act—that “this facility maintains a policy of allowing the use of restrooms by either biological sex, regardless of the designation on the restroom.” (Ex. A, Act. § 1(a) & (b)(3) (capitalization omitted)).

This distinction is critical. Because Plaintiffs need only comply with the Act if the signage language matches their bathroom-usage policy, the Act can only compel speech that is necessarily accurate. Accordingly, as the Act only requires the disclosure of accurate, non-ideological, and non-expressive speech, strict scrutiny does not apply and the statute need only satisfy the requirements of rational-basis review.

Thus, because the Act does not require expressive or ideological speech, the deferential standard of rational-basis review applies. Under rational-basis review, a law is presumed constitutional, and “[t]he burden is on the one attacking the legislative arrangement to negate every conceivable basis which might support it.” *Heller v. Doe*, 509 U.S. 312, 320 (1993) (internal quotations omitted); *see also Walker v. Bain*, 257 F.3d 660, 668 (6th Cir. 2001) (stating that a statute is subject to a “strong presumption of validity” under rational-basis review and will be upheld “if there is any reasonably conceivable state of facts that could provide a rational basis”).

A court conducting a rational-basis review does not sit “as a super legislature to judge the wisdom or desirability of legislative policy determinations” but asks only whether there is some conceivable rational basis for the challenged statute. *Heller*, 509 U.S. at 319. This means that under rational-basis review, it is “constitutionally irrelevant [what] reasoning in fact underlays

the legislative decision.” *R.R. Ret. Bd. v. Fritz*, 449 U.S. 166, 179 (1980) (quoting *Flemming v. Nestor*, 363 U.S. 603, 612 (1960)).

In enacting the informed-consent provision, the General Assembly and the citizens of Tennessee had “absolutely no obligation to select the scheme” that a court might later conclude was best. *Nat’l R.R. Passenger Corp. v. A.T. & S.F.R. Co.*, 470 U.S. 451, 477 (1985); see *McGowan v. Maryland*, 366 U.S. 420, 425-426 (1961) (“State legislatures are presumed to have acted within their constitutional power despite the fact that in practice, their laws result in some inequality.”). And Tennessee “has no obligation to produce evidence to sustain the rationality of its action; its choice is presumptively valid and ‘may be based on rational speculation unsupported by evidence or empirical data.’” *TriHealth, Inc. v. Bd. of Comm’rs*, 430 F.3d 783, 790 (6th Cir. 2005) (quoting *FCC v. Beach Commc’ns, Inc.*, 508 U.S. 307, 315 (1993)).

Here, the rational basis for the Act is readily apparent. Tennessee certainly has a compelling interest in ensuring that patrons are informed of the bathroom-use policy at businesses they frequent—especially when the bathroom-usage policy differs in practice from the existing bathroom signage used by business owners. Many Tennesseans would agree that being “forced to share changing, shower, and bathroom space with members of the opposite sex” does not provide the same “level of privacy and comfort that” a patron could “expect” in facilities separated based on biological sex. *Stuart v. Metro. Gov’t of Nashville & Davidson Cnty.*, 679 F. Supp. 2d 851, 854, 859 (M.D. Tenn. 2009) (Trauger, J.), *vacated after settlement*.

And, although not required by rational-basis review, the statute is also narrowly tailored. The Act simply ensures that Tennesseans are informed of a company’s policy before they enter a locker room or bathroom. It does not require Plaintiffs to adopt any specific bathroom-usage policy, nor does it blanketly prohibit patrons from using the bathroom contrary to their biological

sex. And the Act does not prohibit Plaintiffs from posting additional signs expressing their particular political or social views. Put another way, the law *does* allow Plaintiffs to permit any individual to use whatever restroom they choose, and it *does not* require using only the restroom that corresponds to biological sex. All that is required is that Plaintiffs inform their patrons of their bathroom-usage policy in the event that Plaintiffs use bathroom signage inconsistent with their chosen usage policy.

As the Act does not require—or inhibit—expressive, untruthful, or ideological speech, it easily satisfies the applicable constitutional standard.

### **CONCLUSION**

For the foregoing reasons, Plaintiffs’ Motion for Summary Judgment should be denied.

Respectfully submitted,

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**CERTIFICATE OF SERVICE**

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**In The Matter Of:**

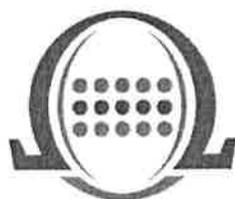
*Bongo Productions, LLC, et al. vs.  
Carter Lawrence, et al.*

---

*Shayne Taylor, M.D.  
December 22, 2021*

---

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IN THE UNITED STATES DISTRICT COURT  
FOR THE MIDDLE DISTRICT OF TENNESSEE  
NASHVILLE DIVISION

BONGO PRODUCTIONS, LLC, ROBERT )  
BERNSTEIN, SANCTUARY PERFORMING )  
ARTS LLC, and KYE SAYERS, )  
)

Plaintiffs, )

NO. 3:32-cv-00490

VS. )

JUDGE TRAUGER

CARTER LAWRENCE, Tennessee State )  
Fire Marshal, in his official )  
capacity, CHRISTOPHER BAINBRIDGE, )  
Director of Codes Enforcement, in )  
his official capacity, GLENN R. )  
FUNK, District Attorney General for )  
the 20th Judicial District, in his )  
official capacity, and NEAL )  
PINKSTON, District Attorney General )  
for 11th Judicial District, in his )  
official capacity, )

Defendants. )

WEB CONFERENCE/REMOTE DEPOSITION OF

SHAYNE TAYLOR, M.D.

December 22, 2021

LYNETTE L. MUELLER, LCR, RDR, CRR, FAPR  
LCR No. 351

Omega Reporting  
901.827.8671

1           The web conference/remote deposition of  
2 SHAYNE TAYLOR, M.D. is taken on December 22, 2021, on  
3 behalf of the Defendants, pursuant to notice and  
4 consent of counsel, beginning at approximately  
5 11:00 a.m.

6           This web conference/remote deposition is  
7 taken pursuant to the terms and provisions of the  
8 Federal Rules of Civil Procedure.

9           The right to read and sign was requested.

10

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25

## A P P E A R A N C E S

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1 SHAYNE TAYLOR, M.D.,  
2 having been first duly sworn, was examined and  
3 testified as follows:

4 EXAMINATION

5 BY MR. RIEGER:

6 Q. Hi, Dr. Taylor. Thanks for -- thanks for  
7 sitting down with us to have this deposition right  
8 before the holidays. We do appreciate it.

9 My name is Alex Rieger, and I am with the  
10 Attorney General's office. And I am lead counsel for  
11 the Defendants in this case.

12 For the record, could I get you to  
13 introduce yourself and spell your last name for the  
14 record.

15 A. Sure. My name is Dr. Shayne Taylor. My last  
16 name is Taylor. T, as in Tom, A-Y-L-O-R.

17 Q. All right. Well, thank you.

18 So in order to ensure that the deposition  
19 goes as smoothly as possible, I like to start with some  
20 rules of the road. So I know, from your preliminary  
21 statement in your expert report, that you have given a  
22 deposition before; is that correct?

23 A. Just one. Yes.

24 Q. Okay. And what was the -- what was your role  
25 in that case? Were you an expert witness in that case?

1 Q. Okay. Is there anything else you do to stay  
2 current on medical research and literature as described  
3 in paragraph 9?

4 A. That's generally most of what I do.

5 Q. Okay.

6 A. The occasional just literature search to make  
7 sure I'm not missing anything big.

8 Q. Okay. All right.

9 Paragraph No. 14, when you say "sex" in  
10 that paragraph, what are you referring to? Are you  
11 referring to biological sex?

12 MS. PICASSO: Object to form.

13 But go ahead and answer, Dr. Taylor.

14 A. I'm describing the sex as assigned at birth by  
15 the visual appearance of an infant's genitals, external  
16 genitals.

17 Q. Is the concept of sex assigned at birth  
18 different from the concept of biological sex?

19 MS. PICASSO: Object to form.

20 But go ahead and answer, Dr. Taylor.

21 A. I think the intention of both of those terms  
22 is the same. I think the difference in the language is  
23 just a bit more nuanced.

24 Q. Can you describe the nuance for me.

25 A. I think "biological sex" can mean any number

1 just in general.

2 A. I think -- I think both. I think it's  
3 problematic for the person who is faced with the  
4 decision to go to the bathroom and being faced with  
5 this sign. I think it's also a larger scale as being a  
6 member of a community whose state legislature feels  
7 that it's appropriate to put up that sign.

8 Q. Okay. Now, is it possible that someone who  
9 has gender dysphoria could see the sign that's proposed  
10 by the challenged law and not think -- not risk  
11 worsening their gender dysphoria or having any -- or  
12 not thinking that it's dangerous and distressing?

13 A. Sure. That's certainly possible.

14 Q. So is there any way to know in advance who  
15 would have that issue of thinking the sign is dangerous  
16 and distressing and -- you know, someone with gender  
17 dysphoria, is there any way to tell whether or not, in  
18 advance of them seeing the sign, that seeing that sign  
19 would worsen their gender dysphoria or be dangerous or  
20 distressing to them?

21 A. No, not that I'm aware of.

22 Q. Okay. Would it just be in -- you know, does  
23 it depend on what they think it represents? What  
24 someone with gender dysphoria, if they saw the sign, is  
25 the harm that you described, that it's dangerous and

1 distressing and runs the risk of worsening gender  
2 dysphoria, is that based upon their perception of what  
3 the sign and use of the term "biological sex"  
4 represents?

5 A. Again, I think that that's part of it, for  
6 sure.

7 Q. Okay. And the other part being the sort of  
8 penumbral societal issue that you've talked about; is  
9 that right?

10 A. Yeah.

11 Q. Okay. Are there any other -- are there any  
12 other facets of harm besides that, you know, person by  
13 person, you know, it's dangerous, distressing,  
14 worsening gender dysphoria, and the societal aspect,  
15 are there any other facets of harm that you think would  
16 be caused by the sign referenced in the challenged law?

17 A. I'm sure --

18 MS. PICASSO: Object to form.

19 But go ahead and answer, Dr. Taylor.

20 A. I'm sure there are plenty. And if -- you  
21 know, given time, I could maybe think of others. But  
22 those are the overarching ones that I'm seeing right  
23 now.

24 Q. Okay. Have you had any -- without going into  
25 patient identity, without broaching any sort of

1 confidentiality, have you had a patient who has -- who  
2 has had worsening gender dysphoria for seeing the  
3 term or having the term said to them "biological sex"?

4 A. No, not specifically that --

5 Q. Okay.

6 A. Nobody has come to my office and said, "I read  
7 this term and that made my dysphoria worse."

8 Q. Okay. Are there common -- this is -- I'm not  
9 a doctor. I'm a lawyer. So forgive me if this is --  
10 this is imprecise or a problem. Feel free to correct  
11 me.

12 Are there common triggers for worsening  
13 gender dysphoria that are common in your practice?

14 A. Yes.

15 Q. What are those?

16 A. There are many upon many upon many. But one  
17 of them could be discrimination that patients are  
18 facing in their communities or at their schools or in  
19 their families. You know, they can be -- I mean, there  
20 are countless reasons why people come to me or reasons  
21 to tell me that their dysphoria is worsening.

22 For fear -- like, in my young -- my young  
23 patients, fear that they will no longer be able to  
24 legally access hormonal treatment; that that could  
25 potentially be taken away from them. They lose

1 insurance. They can potentially lose access to their  
2 treatment.

3 There are many things that -- I mean,  
4 hundreds -- of why somebody would come to my office and  
5 be feeling more dysphoric than prior.

6 Q. But in your opinion, in your years of  
7 practice, the term "biological sex" isn't one of those  
8 common triggers for worsening gender dysphoria?

9 A. No. That has not been something that somebody  
10 has come to my office and complained about  
11 specifically.

12 Q. Okay. Okay.

13 MR. RIEGER: Well, that will do it for me. I  
14 believe that your counsel wants to take a break and  
15 then ask you a couple more questions. I can't promise  
16 that I'm done. Because, you know, I may, depending on  
17 the cross, have one or two.

18 But I'm ready to take that break, Malita,  
19 if you are to go from there.

20 MS. PICASSO: Yeah. Is there any way I could  
21 just have like a ten-minute break? I just have to run  
22 to the restroom. Is that all right?

23 MR. RIEGER: Of course.

24 MS. PICASSO: All right. Cool.

25 MR. RIEGER: Thank you, Dr. Taylor, for

AMENDMENT SHEET

I, the undersigned, SHAYNE TAYLOR, M.D., do hereby certify that I have read the foregoing deposition in the case of BONGO PRODUCTIONS vs. CARTER LAWRENCE and that, to the best of my knowledge, said deposition is true and accurate with the exception of the following corrections listed below:

PAGE/LINE/REASON

Multiple horizontal lines for listing corrections.

Date Signature of Witness

Sworn to and Subscribed before me, this \_\_\_\_ day of \_\_\_\_\_, 2022.

Notary Public My Commission Expires

1 REPORTER'S CERTIFICATE

2 STATE OF TENNESSEE )

3 COUNTY OF SHELBY )

4 I, LYNETTE L. MUELLER, LCR #351, RDR, CRR,  
5 FAPR, and Notary Public for the State of Tennessee, do  
6 hereby certify that the above transcript of proceedings  
7 was reported by me and that the foregoing transcript,  
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9 forth in the caption thereof, were stenographically  
reported by me; constitute a true and correct  
transcript of said proceedings to the best of my  
knowledge, skills, and ability.

10 I FURTHER CERTIFY that I am not related to  
11 any of the parties named herein, nor their counsel, and  
12 have no interest, financial or otherwise, in the  
13 outcome or events of this action.

14 I FURTHER CERTIFY that I am duly licensed  
15 by the Tennessee Board of Court Reporting as a Licensed  
16 Court Reporter as evidenced by the LCR number and  
17 expiration date following my name below.

18 I FURTHER CERTIFY that the right to read  
19 and sign was requested.

20 IN WITNESS WHEREOF, I have hereunto affixed  
21 my official signature and seal of office on 6th of  
22 January, 2022.

23 



25 LYNETTE L. MUELLER, LCR, RDR, CRR, FAPR  
LCR NO. 351, Expires June 30, 2022  
Tennessee LCR No. 351  
Mississippi CSR No. 1794  
Arkansas CCR No. 788  
Notary Public at Large  
For the State of Tennessee  
Commission Expires August 17, 2025

# EXHIBIT A



# State of Tennessee

## PUBLIC CHAPTER NO. 453

HOUSE BILL NO. 1182

By Representatives Rudd, Cepicky, Griffey, Sherrell, Moody, Todd

Substituted for: Senate Bill No. 1224

By Senators Rose, Hensley, Pody

AN ACT to amend Tennessee Code Annotated, Title 4; Title 5; Title 6; Title 7; Title 49 and Title 68, relative to public facilities.

BE IT ENACTED BY THE GENERAL ASSEMBLY OF THE STATE OF TENNESSEE:

SECTION 1. Tennessee Code Annotated, Title 68, Chapter 120, Part 1, is amended by adding the following as a new section:

(a) A public or private entity or business that operates a building or facility open to the general public and that, as a matter of formal or informal policy, allows a member of either biological sex to use any public restroom within the building or facility shall post notice of the policy at the entrance of each public restroom in the building or facility.

(b) Signage of the notice must be posted in a manner that is easily visible to a person entering the public restroom and must meet the following requirements:

(1) Be at least eight inches (8") wide and six inches (6") tall;

(2) The top one-third (1/3) of the sign must have a background color of red and state "NOTICE" in yellow text, centered in that portion of the sign;

(3) The bottom two-thirds (2/3) of the sign must contain in boldface, block letters the following statement centered on that portion of the sign:

THIS FACILITY MAINTAINS A POLICY OF ALLOWING THE USE  
OF RESTROOMS BY EITHER BIOLOGICAL SEX,  
REGARDLESS OF THE DESIGNATION ON THE RESTROOM

(4) Except as provided in subdivision (b)(2), have a background color of white with type in black; and

(5) Be located on a door to which the sign must be affixed or have its leading edge located not more than one foot (1') from the outside edge of the frame of a door to which the sign must be affixed.

(c) If an entity or business is notified that it is not in compliance with this section, the entity or business has thirty (30) days in which to comply before any action is taken against the entity or business.

(d) As used in this section:

(1) "Policy" means the internal policy of a public or private entity or such policy as the result of a rule, ordinance, or resolution adopted by an agency or political subdivision of this state; and

(2) "Public restroom":

(A) Includes a locker room, shower facility, dressing area, or other facility or area that is:

(i) Open to the general public;

**HB1182**

(ii) Designated for a specific biological sex; and

(iii) A facility or area where a person would have a reasonable expectation of privacy; and

(B) Excludes a unisex, single-occupant restroom or family restroom intended for use by either biological sex.

SECTION 2. This act takes effect July 1, 2021, the public welfare requiring it.

HOUSE BILL NO. 1182

PASSED: April 29, 2021



CAMERON SEXTON, SPEAKER  
HOUSE OF REPRESENTATIVES



RANDY MCNALLY  
SPEAKER OF THE SENATE

APPROVED this 17<sup>th</sup> day of May 2021



BILL LEE, GOVERNOR

# Nashville DA won't enforce new bathroom sign law

AP [apnews.com/article/nashville-laws-government-and-politics-50412b91ca33cc45c426a9b5a89b1133](https://apnews.com/article/nashville-laws-government-and-politics-50412b91ca33cc45c426a9b5a89b1133)

May 24, 2021

By KIMBERLEE KRUESI May 24, 2021



Amy Allen, the mother of an 8th grade transgender son, speaks after a Human Rights Campaign round table discussion on anti-transgender laws Friday, May 21, 2021, in Nashville, Tenn. Conservative lawmakers nationwide introduced a flurry of anti-LGBTQ bills this year, but no state's political leaders have gone further than Tennessee in enacting new laws targeting transgender people. (AP Photo/Mark Humphrey)

NASHVILLE, Tenn. (AP) — Nashville's top prosecutor said Monday that he will not enforce a newly enacted law that requires businesses and government facilities open to the public to post a sign if they let transgender people use multiperson bathrooms and other facilities associated with their gender identity.

"I believe every person is welcome and valued in Nashville," Nashville District Attorney General Glenn Funk said in a statement. "Enforcement of transphobic or homophobic laws is contrary to those values. My office will not promote hate."

Funk's office clarified that this refusal to enforce "transphobic or homophobic laws" specifically included the first-of-its kind measure signed by Republican Gov. Bill Lee earlier this month.

The move, along with the flurry of other anti-transgender laws approved by Lee, has sparked alarm among LGBTQ advocates. Many have decried the latest measure as discriminatory and said the required signs are "offensive and humiliating." The law will go into effect July 1.

However, questions have remained about how specifically it will be enforced.

Republican Rep. Tim Rudd, who sponsored the legislation, told a legislative committee in March that the bill "does not provide any fines or penalties at this point," and the amended version passed by that committee became law. Rudd has also said that the law could be enforced by people filing lawsuits or district attorneys asking a judge to force businesses to comply.

Yet Tennessee District Attorneys General Conference President Amy Weirich argued that the language in the new law "doesn't speak to anything having to do with enforcement."

"The way it's written, I don't see anything that allows or provides me the responsibility or right to go to civil court and ask a judge to enforce it," said Weirich, Shelby County's district attorney.

Lee did not have a strong reaction when pressed by reporters Monday on Funk's refusal to enforce the bathroom sign law.

"I think his decision will be his own," he said. "I signed the law; it's his decision how he wants to respond to it."

Lee's response was markedly different than when Funk announced in September he would not enforce a new law that required abortion providers to tell their patients it may be possible to reverse the action of abortion medication half-way through the procedure. Funk said at the time he believed the law was unconstitutional.

Without naming Funk, Lee's office tweeted that, "A district attorney purposefully disregarding current, duly enacted laws by the legislature is a grave matter that threatens our justice system and has serious consequences."

---

Associated Press writer Jonathan Mattise contributed to this report.

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DIAGNOSTIC AND STATISTICAL  
MANUAL OF  
MENTAL DISORDERS

FIFTH EDITION

DSM-5®



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AMERICAN  
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ASSOCIATION  
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These three concepts (for which discussion and examples are provided in Section III and the Appendix) suggest cultural ways of understanding and describing illness experiences that can be elicited in the clinical encounter. They influence symptomatology, help seeking, clinical presentations, expectations of treatment, illness adaptation, and treatment response. The same cultural term often serves more than one of these functions.

## Gender Differences

Sex and gender differences as they relate to the causes and expression of medical conditions are established for a number of diseases, including selected mental disorders. Revisions to DSM-5 included review of potential differences between men and women in the expression of mental illness. In terms of nomenclature, *sex differences* are variations attributable to an individual's reproductive organs and XX or XY chromosomal complement. *Gender differences* are variations that result from biological sex as well as an individual's self-representation that includes the psychological, behavioral, and social consequences of one's perceived gender. The term *gender differences* is used in DSM-5 because, more commonly, the differences between men and women are a result of both biological sex and individual self-representation. However, some of the differences are based on only biological sex.

Gender can influence illness in a variety of ways. First, it may exclusively determine whether an individual is at risk for a disorder (e.g., as in premenstrual dysphoric disorder). Second, gender may moderate the overall risk for development of a disorder as shown by marked gender differences in the prevalence and incidence rates for selected mental disorders. Third, gender may influence the likelihood that particular symptoms of a disorder are experienced by an individual. Attention-deficit/hyperactivity disorder is an example of a disorder with differences in presentation that are most commonly experienced by boys or girls. Gender likely has other effects on the experience of a disorder that are indirectly relevant to psychiatric diagnosis. It may be that certain symptoms are more readily endorsed by men or women, and that this contributes to differences in service provision (e.g., women may be more likely to recognize a depressive, bipolar, or anxiety disorder and endorse a more comprehensive list of symptoms than men).

Reproductive life cycle events, including estrogen variations, also contribute to gender differences in risk and expression of illness. Thus, a specifier for postpartum onset of mania or major depressive episode denotes a time frame wherein women may be at increased risk for the onset of an illness episode. In the case of sleep and energy, alterations are often normative postpartum and thus may have lower diagnostic reliability in postpartum women.

The manual is configured to include information on gender at multiple levels. If there are gender-specific symptoms, they have been added to the diagnostic criteria. A gender-related specifier, such as perinatal onset of a mood episode, provides additional information on gender and diagnosis. Finally, other issues that are pertinent to diagnosis and gender considerations can be found in the section "Gender-Related Diagnostic Issues."

## Use of Other Specified and Unspecified Disorders

To enhance diagnostic specificity, DSM-5 replaces the previous NOS designation with two options for clinical use: *other specified disorder* and *unspecified disorder*. The other specified disorder category is provided to allow the clinician to communicate the specific reason that the presentation does not meet the criteria for any specific category within a diagnostic class. This is done by recording the name of the category, followed by the specific reason. For example, for an individual with clinically significant depressive symptoms lasting 4 weeks but whose symptomatology falls short of the diagnostic threshold for a major depressive episode, the clinician would record "other specified depressive disorder, depressive episode with insufficient symptoms." If the clinician chooses not to specify the

# Gender dysphoria in adolescence: current perspectives

This article was published in the following Dove Press journal:  
Adolescent Health, Medicine and Therapeutics

Riittakerthu  
Kaltiala-Heino<sup>1-3</sup>  
Hannah Bergman<sup>4</sup>  
Marja Työläjärvä<sup>2</sup>  
Louise Frisén<sup>4</sup>

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**Abstract:** Increasing numbers of adolescents are seeking treatment at gender identity services in Western countries. An increasingly accepted treatment model that includes puberty suppression with gonadotropin-releasing hormone analogs starting during the early stages of puberty, cross-sex hormonal treatment starting at ~16 years of age and possibly surgical treatments in legal adulthood, is often indicated for adolescents with childhood gender dysphoria (GD) that intensifies during puberty. However, virtually nothing is known regarding adolescent-onset GD, its progression and factors that influence the completion of the developmental tasks of adolescence among young people with GD and/or transgender identity. Consolidation of identity development is a central developmental goal of adolescence, but we still do not know enough about how gender identity and gender variance actually evolve. Treatment-seeking adolescents with GD present with considerable psychiatric comorbidity. There is little research on how GD and/or transgender identity are associated with completion of developmental tasks of adolescence. **Keywords:** gender dysphoria, gender identity, adolescence, developmental tasks

## Gender dysphoria and related concepts

The fifth edition of the Diagnostic and Statistical Manual of Mental Disorders (DSM-5)<sup>1</sup> defines gender dysphoria (GD) as a condition in which a person has marked incongruence between the expressed or experienced gender and the biological sex at birth. This causes clinically significant distress or impairment in social, occupational or other important areas of functioning. Individuals with GD experience a strong desire to be treated as the other gender (or some alternative gender different from their assigned gender) and/or to be rid of their sex characteristics, and/or the strong conviction of having feelings and reactions typical of the other gender (or some alternative gender). The previous diagnostic term, gender identity disorder, was rejected in the DSM-5 to avoid pathologizing identity.

According to the International Classification of Diseases (ICD)-10,<sup>2</sup> transsexualism is defined as a desire to live and be accepted as a member of the opposite sex, usually accompanied by a sense of discomfort with or the inappropriateness of one's anatomical sex and a wish to undergo surgery and hormonal treatment to make the body as congruent as possible with the individual's preferred sex. The forthcoming ICD-11 will reconceptualize gender identity-related diagnoses using gender incongruence as the main term.<sup>3</sup>

In addition to the DSM-5 diagnostic term, gender dysphoria can also refer to anxiety and distress about gender features at large. Gender nonconformity refers to

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behaviors and an appearance that are considered atypical of an individual's assigned gender. Gender variance refers to a spectrum of gender experience, in contrast to the dichotomized conception of gender. The term "transgender" is used as an umbrella term to refer to wider variation of gender identities. Not all who identify as transgender or display gender nonconformity or gender variance suffer from dysphoria.

In this article, we use the DSM-5 and ICD-10 terms "gender dysphoria (GD)" and "transsexualism/transsexual", respectively, when referring to diagnosed clinical samples, and also when referring to the literature published when earlier versions of the DSM classification were in use. We use "transgender" to refer to self-identified population samples and "gender dysphoria" to refer to those who present clinical symptoms.

## How common are GD and transgender identity among adults and adolescents?

The number of people who seek treatment suggest that male-to-female transsexualism has a prevalence of 6.8/100,000 and female-to-male transsexualism has a prevalence of 2.6/100,000 among adults.<sup>4,5</sup> In the Netherlands, 0.6% of men and 0.2% of women (aged 15–70 years) reported incongruent gender identity and a desire to undergo sex reassignment (SR).<sup>6</sup> Population surveys have suggested that about 0.5% of adults in the general population identify as transgender.<sup>5,7</sup>

The number of adolescents contacting specialized gender identity services has risen considerably over the past decade across Europe and North America.<sup>8,9</sup> No conclusions regarding the prevalence of GD in general or of GD/transsexualism specifically can be drawn based on these increases. Studies using short (one to three item) self-reports of gender identity and its variance suggest that 0.17%–1.3% of adolescents and young adults identify as transgender.<sup>5,10</sup> A school-based survey eliciting gender experiences with scales commonly used at gender identity services suggested that 1.3% of 16–19 year olds had potentially clinically significant gender dysphoria.<sup>11</sup>

## Gender identity

Identity is the way one understands, describes and expresses oneself and the reflection of those entities to others. Identity consists of many integrated aspects such as gender, nationality, language, academic and occupational endeavors, and religious and political convictions. It is affected by interpersonal relationships, society and different events throughout the life course.<sup>12</sup> Adolescence is an important period of identity formation and integration.<sup>12,13</sup> Adolescents and young adults

establish their identity by actively exploring identity-related choices and making identity commitments in their chosen directions.<sup>12,13</sup>

Gender identity concerns the individual's core sense of being "female", "male" or another gender. The development of gender identity is a complex process affected by multiple factors.<sup>14,15</sup> In the research tradition of gender identity, the broad focus has been on the theme "sex differences", and two major topics have received the most attention: the description and measurement of sex differences and the etiology of these differences.<sup>16</sup> Several theories have been proposed. According to early psychodynamic theories, gender variant behavior was hypothesized to derive from parent–infant interpersonal issues or trauma (see Gray et al<sup>15</sup>). However, these theories have not undergone adequate scientific testing. Gender identity development has mostly been described from the viewpoint of cognitive and social learning theories, which argue that human beings are active constructors of cognitive schemas, including gender, in continuous interaction with the environment.<sup>14</sup> Other theories on processes of gender typing have focused on proximal and distal biological influences, genetic and epigenetic or hormonal and neural mechanisms as well as brain anatomy differences in the etiology of gendered behavior and gender variance.<sup>17–19</sup> There are structural and functional sex differences in the brain, some of them observable across the life span and others only during specific developmental phases. Sex differences in the brain are largely determined by steroid hormone exposure during a perinatal sensitive period that alters subsequent hormonal and non-hormonal responses throughout the lifespan, but they also depend on genes on sex chromosomes. Moreover, there is continuous interaction between genes and experiences, "epigenetics", which changes the expression of genes without any change in the underlying DNA sequence. Research suggests that, for example, early social experiences may act as such epigenetic influence that they ultimately shape lasting sex differences in brain and behavior,<sup>20–22</sup> but a lot more research is needed in this field to obtain solid knowledge relevant for understanding GD.

While the theories proposed in the past have generally been either essentialist or social cognitive/constructivist in nature, researchers today are expanding the focus to include the bio-psycho-social processes that probably occur across development.<sup>14,15,23</sup>

## Childhood GD and puberty development

GD in childhood (GDC)<sup>1</sup> describes a feeling of incongruence between the experienced (psychological) gender and the sex

assigned at birth. A corresponding diagnosis is included in the ICD-10.<sup>2</sup> Healthy children vary considerably in their gendered behaviors.<sup>15</sup> The diagnosis of GD in prepubescent children has been widely discussed throughout the history of gender identity research, mainly in terms of weighing the risk of stigmatization against diagnosis as a means of access to publicly funded or insurance-covered health care.<sup>3,24</sup> The prevalence of GDC is not known.<sup>5</sup>

Adolescence is a crucial time for identity and psychosexual development in young people with gender identity concerns.<sup>25</sup> The outcomes of GDC have been discussed in terms of its persistence and desistance. For most children with GDC, whether GD will persist or desist will probably be determined between the ages of 10 and 13 years,<sup>26</sup> although some may need more time.<sup>27</sup> Evidence from the 10 available prospective follow-up studies from childhood to adolescence (reviewed in the study by Ristori and Steensma<sup>28</sup>) indicates that for ~80% of children who meet the criteria for GDC, the GD recedes with puberty. Instead, many of these adolescents will identify as non-heterosexual.<sup>17,29</sup> Steensma et al<sup>26</sup> interviewed adolescents with different outcomes of GDC (persistence or desistance). The adolescents mentioned social environment, the anticipated results of bodily changes and first romantic and/or sexual experiences as central factors in the desistance or persistence of GD.

## Treatment of GD intensifying in puberty: the Dutch model

The most commonly used guidelines for the treatment of GD in children and adolescents are those of The Endocrine Society<sup>30</sup> and the Standard of Care from the World Professional Association for Transgender Health,<sup>31</sup> which are based on the so-called Dutch Model protocols published and practiced at the Amsterdam Gender Clinic in the Netherlands.<sup>32</sup>

The Dutch protocol recommends medical treatment if GD intensifies in puberty, while the care for children with GD and their families consists of providing information, psychological support, parental or/and family counseling. In adolescents, medical treatment is recommended at age 12 years and older for those who are in or beyond the early stages (Tanner II–III) of puberty and are still experiencing persistent GD. Puberty suppression with gonadotropin-releasing hormone analogs is part of the protocol for these patients. The purpose of puberty suppression is to relieve the psychological suffering caused by the development of secondary sex characteristics, to give the adolescent time to make a balanced decision regarding whether to undergo actual medical gender-confirming treatment (with cross-sex

hormones and surgery) and to make social “passing” in the experienced gender easier. Cross-sex hormones are used for adolescents aged 16 years and older who continue to experience persistent GD. People aged 18 years and older with a diagnosis of GD may undergo SR surgery.<sup>32</sup>

## Outcome of and ethical debates around medical interventions for GD in adolescence

The Dutch protocol is largely used, but it has its critics.<sup>33–35</sup> Controversy regarding the use of drugs for puberty touches on fundamental ethical concepts in pediatrics: the best interests of the minor, autonomy and the role of social context. Professionals recognize the distress of young people with GD and feel an urge to treat them. At the same time, most of these professionals have doubts because of the lack of data regarding long-term physical and psychological outcomes.<sup>36,37</sup>

Reports of the outcomes of puberty suppression treatment in adolescents have shown reasonable safety and good outcomes regarding patient satisfaction and psychosocial functioning, but research is still scarce. Nevertheless, puberty suppression is not indicated in a considerable proportion of gender dysphoric minors because of several reasons, for example, severe psychiatric comorbidity, considerable instability of psychosocial support or onset of GD later during puberty and diagnostic uncertainty,<sup>38–40</sup> nevertheless, more follow-up data even from patients who are fulfilling the criteria for “the Dutch model” are still needed.<sup>37</sup>

## Psychiatric disorders among adolescents with GD

Descriptive studies of adolescents referred to specialized gender identity services at different centers in Europe and North America have mainly suggested that ~40%–45% of these young people present with clinically significant psychopathology.<sup>38,39,41–50</sup> The lowest figures for psychiatric comorbidity (one-third of the presenting population) were reported in the Netherlands,<sup>41</sup> and the highest (up to three quarters) was reported in Finland and Canada.<sup>39,50</sup> Gender-referred adolescents actually appear to display clinically significant psychopathology to the same extent as adolescents referred to mental health services due to other reasons.<sup>48,50</sup> The most commonly reported disorders are depression and anxiety disorders. Self-harm and suicidal ideation/behavior are also common, whereas conduct disorder and antisocial development do not appear central in this population.

Likewise, community-level information suggests that transgender-identifying youth present four to six times more often with depression and three to four times more often with self-harm and/or suicidal behavior compared with cisgender adolescents.<sup>10,51</sup> Clinical and population data, though scarce, also suggest an overrepresentation of eating pathology among adolescents with GD or transgender identity.<sup>46,52</sup>

An increased prevalence of autism spectrum disorders (ASDs), varying from ~6% to over 20%, has been reported among samples of adolescents referred to gender identity services.<sup>39,42,46,53</sup> This vastly exceeds the estimated prevalence of 0.6%–0.7%<sup>54</sup> in the general population. In comparison, among children and early adolescents with ASDs, gender variance is >7-fold more common than among non-referred controls.<sup>53,55</sup>

Hypotheses to explain this are manifold. The theory of the extreme male brain suggests that individuals with ASD demonstrate an extreme of the typical male pattern of behaviors and cognitions originating from high levels of fetal testosterone. High fetal levels could likewise contribute to GD in natal girls, explaining their male identity and behavior. However, this theory cannot explain the association between ASD and GD in natal boys. Social factors-related hypotheses propose that the social perception and communication difficulties typical of autism could make a child more likely to miss social cues regarding how to conform to gender norms or to identify with the opposite sex when he/she faces difficulties joining the peer group of her/his own sex. Hypotheses focusing on individual psychological characteristics suggest, firstly, that gender could be among the preoccupations or obsessions often seen in ASDs. On the other hand, the development of atypical gender identity in autism could relate to the developmental rigidity typical of autism. Individuals with ASD might not reach normative flexibility in gender development necessary to deal with gender variant feelings, which might lead to the overrepresentation of ASD in GD.<sup>53,56</sup> The suggested causes, however, remain speculative. In a recent study, both boys and girls with GD displayed elevated levels of autistic symptomatology in all subdomains of autism, which did not exclusively support any of the suggested hypotheses.<sup>56</sup> Nevertheless, ASDs pose particular challenges for the diagnosis and treatment of GD in adolescents.

## GD and the developmental tasks of adolescence

“Developmental tasks” refer to the normative developmental milestones that should be reached during a given

developmental stage.<sup>57,58</sup> They arise from interactions among physical development, personal attributes and societal expectations. Favorable completion of the developmental tasks of a given stage is a prerequisite for success in the subsequent stages. The developmental tasks of adolescence were first formulated by Havighurst<sup>57</sup> and comprise accepting one’s body, adopting a masculine or feminine social role, achieving emotional independence from parents, developing close relationships with peers of the same and opposite genders, preparing for an occupation, preparing for marriage and family life, establishing a personal value or an ethical system and achieving socially responsible behavior. Although puberty now occurs earlier and the transition to adulthood occurs later than they did when these developmental tasks were initially proposed, they remain relevant.<sup>58</sup> The relationship with one’s own body and the acquisition of a gendered social role – not necessarily binary – are by definition challenging for adolescents with GD. In the following sections, we discuss the available information on GD/transgender identity and the other developmental tasks of adolescence.

## GD in adolescence and relationships with parents

Parents of adolescents with GD and/or transgender identity may face special challenges that are shaped by a variety of factors, such as ethnicity, religious background, social class and the prevailing attitudes in their community and society.<sup>59,60</sup> These challenges likely shape the support that a nonconforming adolescent can receive. Adverse parental reactions toward an adolescent’s gender nonconformity have been noted as a special risk,<sup>61</sup> but parents of sexual- and gender-minority offspring have also reported particular positive aspects of being a parent in this situation, such as personal growth, unconditional love, activism, social connection and closer relationships.<sup>62</sup> However, few studies have empirically explored the parental reactions and support among youth with GD and/or transgender identity.

In a Canadian community study of transgender-identifying youth,<sup>63</sup> of those who had disclosed their gender identity to their parents, 34% considered their parents “very” supportive and 25% considered their parents “somewhat” supportive. Forty-two percent reported that their parents were “not very” or “not at all” supportive. However, the study was based on a nonrandom sample and solely adolescent self-reports, so findings need to be interpreted with caution and causalities cannot be concluded. Strong perceived parental support was, nevertheless, associated with many positive mental health outcomes. Lack of parental support was associated

with inadequate housing and homelessness in addition to negative psychological outcomes. Better parental support has also been associated with fewer risk-taking sexual behaviors among transgender youth.<sup>64</sup>

In a community study of trans female adolescents and young adults,<sup>65</sup> more than half of the participants reported that their parents supported their gender identity, showed their support in many ways and believed the respondent could have a happy future as a trans adult. However, approximately two in five respondents had not experienced parental acceptance. Parental acceptance was associated with perceiving parents as the primary source of social support.

In a school-based survey<sup>51</sup> transgender-identifying adolescents felt less often (odds Ratio 0.3) than their cis-gender peers that at least one parent cared for them.

Studies of clinically referred gender dysphoric youth have rarely addressed parent-related issues. Simons et al<sup>66</sup> reported that in a clinical sample of adolescents with GD, parental support was significantly associated with higher life satisfaction, lower perceived burden of being transgender and fewer depressive symptoms. In a Finnish study comparing childhood gender identity in community and clinical samples, a smaller proportion of adolescents with GD than of non-referred adolescents in the population agreed with the statement "I always felt that my parents cared for me."<sup>11</sup> It was also noticed that the clinically referred adolescents with GD less commonly lived with both their parents than the adolescents in the normal population (48% vs. 78%).<sup>67</sup> In British and Spanish samples of gender-referred adolescents, parental divorce was observed in the background of approximately three in five participants, but the authors did not compare this finding with a corresponding rate in the general population.<sup>46,49</sup>

## Gender nonconformance and peer relationships in adolescence

During adolescence, peer relationships are critical for psychological well-being.<sup>68,69</sup> Peer relationships also shape development, including aspects of gender identity consolidation.<sup>70</sup> Loneliness and social isolation from peer relationships is associated with developmental difficulties and impaired mental health.<sup>71,72</sup> An important peer network-related risk factor is bullying.<sup>73</sup>

Observations in referred samples of adolescents with GD suggest considerable peer relationship difficulties. In both the UK<sup>46</sup> and in Finland,<sup>39</sup> approximately half of adolescents who presented at a specialized gender identity service reported significant experiences of being bullied. In the Finland study, 45% of the referred adolescents also had a history of marked

periods of social isolation in childhood and/or adolescence. In the Netherlands and in Canada, self-, parent and teacher ratings indicated poorer peer relationships among adolescents referred for GD than in the same-aged population<sup>47,48</sup> and poor peer relationships were an important correlate of mental health problems in this group. Similarly, in another Canadian comparison among gender-referred, mental health-referred and general population adolescents bullying was reported by the GD group more commonly than by population controls, and to the same extent as by those referred due to mental health issues. Gender-related bullying was most common among the GD group.<sup>74</sup>

On the population level, Clark et al<sup>51</sup> found that transgender-identifying adolescents had 4.5-fold increased odds of being bullied and were approximately twice as likely to report being afraid for their personal safety, having been in a serious physical fight and having been hit or otherwise harmed by others, compared with their cisgender-identifying peers. They also less commonly felt that their friends cared about them and that school was okay.

Gender-nonconforming behavior is characteristic of both sexual- and gender-minority youth and has been associated with an increased likelihood of experiencing bullying and harassment in peer groups.<sup>75,76</sup> Adolescents with GD likely represent the extreme end of gender nonconformity, and this may strongly contribute to their experiences of being bullied. Bullying and stigmatization have also been suggested to (partially) mediate the association between gender nonconformity and lower mental well-being across adolescence.<sup>74,77-79</sup>

However, not all the difficulties the gender dysphoric adolescents face in peer relationships can be explained by gender expression-related victimization or discrimination. In the Finnish clinical sample, of the gender identity-referred adolescents who had experienced severe bullying at school, three quarters had been bullied before they ever questioned their gender. Likewise, three-quarters of them reported that the bullying had not been related to gender expression or sexual identity, but to other factors such as not being slim, being successful at school or having unfashionable hobbies and interests.<sup>39</sup> Bullying is a severe problem regardless of the reported reasons for it, but it is important to acknowledge that adolescents who develop GD also have unrelated difficulties that may need attention.

## GD, transgender identity and sexuality in adolescence

Sexual orientation and gender identity are different entities, and transgender people present with a variety of sexual orientations. Nevertheless, sexual orientation has long been used

to subtype GD/transsexualism.<sup>80</sup> During adolescence, the different facets of sexual orientation – attraction, behavior and identity – may still be developing. It may be more important to determine whether adolescents with GD or transgender identity display developmentally appropriate and favorable involvement in romantic and erotic relationships.

In adolescence, sexual development accelerates. Young people's experiences of a maturing and changing body, sexuality and their developing gender identity affect intrapersonal, interpersonal and societal interactions.<sup>81</sup> In Western countries, between one-tenth and one-third of adolescents first experience sexual intercourse by the age of 15, and the vast majority experience it by age 20.<sup>82,83</sup> Various practices of kissing and petting typically precede first sexual intercourse by several years. Early sexual activity has been viewed as a problem behavior associated with risky sexual behaviors, psychosocial difficulties and emotional and behavioral disorders.<sup>82,83</sup> In contrast, in the late stages of adolescent development, a lack of experiences might suggest developmental difficulties.

GD and/or transgender identification could be expected to be associated with delayed sexual development, given that it is the sexual body, in particular, that is the source of distress in GD and that differing from the mainstream may increase the adolescent's risk of problems in social relationships, including dating, and sexual encounters. Sexual- and gender-minority adolescents may also have a reduced availability of potential partners and increased challenges in finding potential partners than their heterosexual peers.<sup>84</sup> However, developmental challenges have also been associated with premature and risky sexual behavior.<sup>82,85</sup> Adolescents with GD and/or transgender identification could engage in risky sexual behaviors due to identity experiments or because associated mental health problems could increase their search for comfort in intimacy or decrease their self-protection skills.

To the best of our knowledge, the only study focusing on the sexual experiences of treatment-seeking adolescents with GD is that of Bungener et al<sup>86</sup> from the Netherlands. They compared the sexual experiences of 137 transsexual adolescents (mean [SD] age 14.69 [2.2] years) with those of a same-aged adolescent population. Transsexual adolescents had fewer sexual experiences than the same-aged population in all areas measured (falling in love, romantic relationships, kissing, petting, intercourse). However, a majority of the transsexual adolescents had fallen in love and approximately half had been involved in romantic relationships. One quarter had experienced petting while undressed, and 5% had experienced sexual intercourse. Fewer transsexual adolescents than the adolescents in the same-aged population (24% vs. 48%)

valued sex as important. In a descriptive study of clinically presenting adolescents with GD in the USA,<sup>45</sup> nearly half of the respondents (mean [SD] age 19.2 [2.9] years) reported being sexually active.

Some population studies provide information regarding transgender identity in adolescence and aspects of sexual development. Korchmaros et al<sup>84</sup> compared the romantic relationships of lesbian, gay, bisexual, transgender and questioning (LGBTQ) adolescents and those of adolescents with mainstream sexual and gender identities. Contrary to expectations, the LGBTQ adolescents were more experienced with romantic relationships and more active in initiating relationships both online and offline. Results were not reported separately for the transgender group. Robinson and Espelage<sup>87</sup> reported that LGBTQ adolescents were more likely to display risky sexual behaviors than same-aged non-LGBTQ youth. However, in more detailed analyses, the risk was associated with homosexual/bisexual orientation and not with transgender identity. Veale et al<sup>88</sup> set out to study pregnancy involvement among transgender youth and the health correlates of this involvement. In a large (n=923) sample of transgender-identifying youth, 540 responded to the pregnancy involvement item. Almost 5% of Canadian transgender adolescents had ever been pregnant or impregnated a partner; approximately the same proportion as their same-aged peers. Those with a history of pregnancy involvement were also more likely to have a history of sexually transmitted disease, but they did not differ from the rest of the transgender youth in terms of hormone use, living in the felt gender, self-reported mental health and level of social support.

Sexual harassment is a common problem among adolescent populations.<sup>89</sup> Transgender-identifying adolescents appear to be at the greatest risk of sexual harassment and to experience the greatest distress due to it.<sup>89</sup> Sexual harassment is suggested to function to maintain heteronormativity, which transgender adolescents likely challenge. Their perception of sexual harassment as more distressing compared with other adolescents could be due to harsher harassment, increased vulnerability due to uncertainty about self, or fear.<sup>89</sup>

Similarly, in a large school-based survey study on teen dating violence,<sup>90</sup> the few transgender-identifying youth in the sample reported the highest victimization rates for physical dating violence, psychological dating abuse, cyber dating abuse and sexual coercion. Differences from cisgender adolescents varied from 2- to 7-fold for the different forms of violence. However, the transgender-identifying youth also reported the highest rates of perpetrating dating violence. Minority stress theory<sup>91</sup> posits that the chronic stressors that

minorities experience (e.g., gender-based discrimination) shape their coping mechanisms (such as substance use, aggression) and lead to adverse psychosocial and health outcomes. The particular vulnerability to perpetrating dating violence observed among transgender adolescents by Dank et al<sup>90</sup> could be understood through minority stress theory, but more research is needed.

Transgender adolescents and young adults, particularly trans females, are at a disproportionately high risk of contracting human immunodeficiency virus and other sexually transmitted diseases.<sup>79,92</sup> The risk of unprotected sex in this population has been associated with sex work and drug use, which are further associated with rejection, stigma and discrimination.<sup>79,92</sup> Of the studies of referred samples, only one addressed sex work.<sup>45</sup> In that sample, 6% of the referred adolescents reported engaging in the trading of sex.

Sexual education is an important way to promote positive and responsible sexual behaviors in youth. The planned curricula and practical applications likely vary widely across countries and schools. Sexual- and gender-minority youth were found to desire minority-inclusive sexual education in a study by Gowen and Wingez-Yanez.<sup>93</sup> The sexual- and gender-minority youth felt that the sexual education that was offered isolated them by silencing them, adopting a hetero-centric perspective and pathologizing minorities. Reflecting on the available sexual education in light of these findings is appropriate for all educators.

## Preparing for occupation: academic performance and socioeconomic status

To the best of our knowledge, research has not specifically focused on academic performance and the progression to work life among adolescents with GD, but given the burden of psychiatric comorbidities among gender-referred youth, special needs regarding education are likely to exist.

Aspects of social relationships are relevant to well-being in school, school performance and pathways to occupation. Transgender youth have been reported to experience bullying and discrimination in schools, not only by peers but even by teachers; consequently, they perceive schools as unsafe places, which again increases the risk of non-attendance and poorer results.<sup>75,94</sup> Gender- and sexuality-related victimization may impair academic performance through, for example, decreased motivation, concentration and self-efficacy and the resulting school avoidance and harmful coping strategies.<sup>94,95</sup> Nevertheless, being “out” at school improves self-esteem among gender- and sexual-minority youth and increases their

well-being, which can have a positive impact on academic performance.<sup>94</sup>

School dropout is strongly linked to social exclusion. School dropout was associated with high masculinity in girls and low masculinity combined with high femininity in boys in a study of late-adolescent school dropouts and attenders in the Netherlands.<sup>96</sup> The authors suggested that such deviation from gender norms increases the risk of unpopularity among peers, which again predisposes individuals toward school dropout. However, school dropout was also associated with very masculine attitudes and self-evaluations among boys. The role of gendered behaviors, attitudes and experiences in school adjustment and academic performance deserves further research.

In Clark et al's<sup>51</sup> school-based survey, adolescents reporting non-cisgender identity came disproportionately often from families with high socioeconomic deprivation and less often felt that their family got along. Any explanation for this remains unknown; however, young people are likely to stay in the same socioeconomic position as their parents.<sup>97</sup> Jacob and Cox<sup>98</sup> also pinpointed transgender people's greater risk of having a disadvantaged socioeconomic status (in the USA), associating this with increased unemployment, and employment in low-paid jobs, because of stigmatization.

## Why the increase in referrals?

Zucker et al<sup>99</sup> observed an increase in the number of adolescents presenting at gender identity services in the early 2000s. Since then, several gender identity services for minors from across Western countries have reported increases.<sup>8,9,42,49</sup> Simultaneously, the earlier overrepresentation of natal boys has equaled or turned to overrepresentation of natal girls.<sup>9</sup> Natal girls now comprise from half<sup>9</sup> to ~90%<sup>39</sup> of clinical adolescent samples. The reasons for these changes are not known. The increase in referrals could be attributable to enhanced provision of services, or the threshold for seeking help may now be lower due to increased knowledge and improved societal acceptance. Aitken et al,<sup>9</sup> however, did not find evidence supporting a lowered threshold to gender identity services. Sociocultural features related to what kind of identities are available for whom, and sex-related differences of pressure to conform may play a role.

## Comments

Research regarding the clinical treatment of adolescents with GD has mainly focused on childhood-onset GD that intensifies during puberty, and the Dutch treatment protocol is also tailored for this group. There is little empirical knowledge

regarding young people who experience their first signs of GD in adolescence, well after the onset of puberty, especially regarding biological girls.<sup>50,100</sup> Among a treatment-seeking sample in the UK, 18% experienced their first feelings of GD in adolescence<sup>46</sup> compared with approximately two-thirds of the Finnish sample,<sup>39</sup> and for the majority of adolescent-onset cases, GD presented in the context of severe mental disorders and general identity confusion. In such situations, appropriate treatment for psychiatric comorbidities may be warranted before conclusions regarding gender identity can be drawn. Gender-referred adolescents actually display psychopathology to the same extent as mental health-referred youth.<sup>48,50</sup> In a nationwide long-term follow-up study of adult cases, psychiatric morbidity, suicide attempts and suicide mortality persisted as elevated after juridical and medical SR.<sup>101</sup>

Emerging discussions raise concern for post-pubertally abruptly emerging cross-gender identification (“rapid onset”), particularly among biological girls, suggesting a role for intensive media influences and generous group validation as shaping the understanding of, and giving new meanings to, the body discomfort common among female adolescents at large.<sup>100</sup> The persistence of increasing adolescent-onset transgender identification is not known.<sup>5,100</sup>

More empirical research is needed regarding virtually all aspects of GD in adolescence to create treatment approaches that optimize these young people’s future psychosocial health and well-being. It seems unlikely that all the psychopathology observed in the referred samples is secondary to gender identity issues and would resolve with hormonal and later surgical treatments. There is still no clear consensus regarding hormonal treatment for adolescents because long-term data are unavailable;<sup>36</sup> actually, only one long-term follow up has been carried out, with a highly selected intervention group and an at baseline non-comparable comparison group.<sup>102</sup>

An affirmative approach<sup>103</sup> is increasingly implemented in the health care of gender nonconforming children. This includes, based on a comprehensive psychological and psychosocial assessment, work with the children and their families and schools to support the gender-nonconforming minors to express themselves in a way that feels most comfortable for them. With the starting point that gender presentations are fluid and changing over time, gender variant children need to be allowed to freely explore a range of gender identities and expressions. A debate concerns whether or not a prepubertal child should be allowed to completely transition to live in other than birth gender. Concerns include that childhood transition may be forcing adolescents to proceed to biomedical interventions, as stepping back may be psychologically

troublesome, even though identity development has taken a new direction.<sup>28,104</sup>

The etiology of gender incongruence remains unknown. Gender identity differentiation does not occur in a psychosocial vacuum; instead, research in the field suggests that the developmental course is influenced by numerous psychosocial factors, likely in continuous interaction with biological factors.<sup>23,105</sup> Gray et al<sup>15</sup> noted that the general narrative in the research literature concerning gender variation among children focuses on gender “atypical” behavior and deviation from “normative patterns”, thus viewing gender in a binary way instead of as a wider spectrum of (healthy) identities, personalities and behaviors among children. This is surely relevant for adolescents as well. These authors also requested a shift in research paradigms away from the study of outcomes of sexuality and gender identity and the child/adolescent in isolation toward outcomes of adjustment and the child/adolescent in contexts that affect adjustment. Along with further discussions of the best treatment interventions, it is relevant to attempt to contribute to societal attitudes that enable children and adolescents with gender variance to express themselves and successfully complete the developmental tasks common to all, independent of gender.

## Disclosure

The authors report no conflicts of interest in this work.

## References

1. American Psychiatric Association. *Diagnostic and Statistical Manual of Mental Disorders*. 5th ed. Washington, DC: American Psychiatric Press; 2013.
2. World Health Organization. *The ICD-10 Classification of Mental and Behavioural Disorders: Clinical Descriptions and Diagnostic Guidelines*. Geneva: World Health Organization; 1992.
3. Drescher J, Cohen-Kettenis PT, Reed GM. Gender incongruence of childhood in the ICD-11: controversies, proposal, and rationale. *Lancet Psychiatry*. 2016;3(3):297–304.
4. Arcelus J, Bouman WP, Van Den Noortgate W, Claes L, Witcomb G, Fernandez-Aranda F. Systematic review and meta-analysis of prevalence studies in transsexualism. *Eur Psychiatry*. 2015;30(6):807–815.
5. Zucker KJ. Epidemiology of gender dysphoria and transgender identity. *Sex Health*. 2017;14(5):404–411.
6. Kuyper L, Wijnen C. Gender identities and gender dysphoria in the Netherlands. *Arch Sex Behav*. 2014;43(2):377–385.
7. Crissman HP, Berger MB, Graham LF, Dalton VK. Transgender demographics: a household probability sample of US adults, 2014. *Am J Public Health*. 2017;107(2):213–215.
8. Wood H, Sasaki S, Bradley SJ, et al. Patterns of referral to a gender identity service for children and adolescents (1976–2011): age, sex ratio, and sexual orientation. *J Sex Marital Ther*. 2013;39(1):1–6.
9. Aitken M, Steensma TD, Blanchard R, et al. Evidence for an altered sex ratio in clinic-referred adolescents with gender dysphoria. *J Sex Med*. 2015;12(3):756–763.
10. Connolly MD, Zervos MJ, Barone CJ, Johnson CC, Joseph CL. The mental health of transgender youth: advances in understanding. *J Adolesc Health*. 2016;59(5):489–495.

11. Sumia M, Lindberg N, Tyolajarvi M, Kaltiala-Heino R. Current and recalled childhood gender identity in community youth in comparison to referred adolescents seeking sex reassignment. *J Adolesc*. 2017;56:34–39.
12. Kroger J. *Identity Development: Adolescence Through Adulthood*. 2nd ed. Thousand Oaks, CA: Sage Publications; 2007.
13. Kroger J, Martinussen M, Marcia JE. Identity status change during adolescence and young adulthood: a meta-analysis. *J Adolesc*. 2010;33(5):683–698.
14. Martin CL, Ruble DN. Patterns of gender development. *Annu Rev Psychol*. 2010;61:353–381.
15. Gray SAO, Carter AS, Levitt H. A critical review of assumptions about gender variant children in psychological research. *J Gay Lesbian Ment Health*. 2012;16(1):4–30.
16. Wharton A. *The Sociology of Gender—An Introduction to Theory and Research*. 2nd ed. Chichester: Wiley-Blackwell; 2012.
17. Alanko K, Santtala P, Harlaar N, et al. Common genetic effects of gender atypical behavior in childhood and sexual orientation in adulthood: a study of Finnish twins. *Arch Sex Behav*. 2010;39(1):81–92.
18. McCarthy MM, Auger AP, Bale TL, et al. The epigenetics of sex differences in the brain. *J Neurosci*. 2009;29(41):12815–12823.
19. Meyer-Bahlburg HF. Introduction: gender dysphoria and gender change in persons with intersexuality. *Arch Sex Behav*. 2005;34(4):371–373.
20. Knickmeyer R, Wang J, Zhu H, et al. Impact of sex and gonadal steroids on neonatal brain structure. *Cereb Cortex*. 2014;24:2721–2731.
21. Marrocco J, McEwen BS. Sex in the brain: hormones and sex differences. *Dialogues Clin Neurosci*. 2016;18(4):373–338.
22. Ratnu V, Emami MR, Bredy TV. Genetic and epigenetic factors underlying sex differences in the regulation of gene expression in the brain. *J Neurosci Res*. 2017, 95:301–310.
23. Fausto-Sterling A, Crews D, Sung J, Garcia-Coll C, Seifer R. Multimodal sex-related differences in infant and in infant-directed maternal behaviors during months three through twelve of development. *Dev Psychol*. 2015;51(10):1351–1366.
24. Drescher J. Controversies in gender diagnoses. *LGBT Health*. 2013;1(1):10–14.
25. Steensma TD, McGuire JK, Kreukels BP, Beekman AJ, Cohen-Kettenis PT. Factors associated with desistance and persistence of childhood gender dysphoria: a quantitative follow-up study. *J Am Acad Child Adolesc Psychiatry*. 2013;52(6):582–590.
26. Steensma TD, Biemond R, de Boer F, Cohen-Kettenis PT. Desisting and persisting gender dysphoria after childhood: a qualitative follow-up study. *Clin Child Psychol Psychiatry*. 2011;16(4):499–516.
27. Steensma TD, Cohen-Kettenis PT. More than two developmental pathways in children with gender dysphoria? *J Am Acad Child Adolesc Psychiatry*. 2015;54(2):147–148.
28. Ristori J, Steensma TD. Gender dysphoria in childhood. *Int Rev Psychiatry*. 2016;28(1):13–20.
29. Singh D. *A Follow-up Study of Boys with Gender Identity Disorder* [Academic dissertation]. University of Toronto; 2012. Available from: [https://tspace.library.utoronto.ca/bitstream/1807/34926/1/Singh\\_Devita\\_201211\\_PhD\\_Thesis.pdf](https://tspace.library.utoronto.ca/bitstream/1807/34926/1/Singh_Devita_201211_PhD_Thesis.pdf). Accessed January 11, 2018.
30. Hembree WC, Cohen-Kettenis PT, Gooren L, et al. Endocrine treatment of gender-dysphoric/gender-incongruent persons: an endocrine society clinical practice guideline. *J Clin Endocrinol Metab*. 2017;102(11):3869–3903.
31. Coleman E, Bockting W, Botzer M, et al. Standards of care for the health of transsexual, transgender, and gender-nonconforming people, version 7. *Int J Transgend*. 2011;13(4):165–232.
32. de Vries AL, Cohen-Kettenis PT. Clinical management of gender dysphoria in children and adolescents: the Dutch approach. *J Homosex*. 2012;59(3):301–320.
33. Korte A, Lehmkuhl U, Goecker D, Beier KM, Krude H, Gruters-Kieslich A. Gender identity disorders in childhood and adolescence: currently debated concepts and treatment strategies. *Dtsch Arztebl Int*. 2008;105(48):834–841.
34. Viner R, Brain C, Carmichael P, Di Ceglie D. Sex on the brain: dilemmas in the endocrine management of children and adolescents with gender identity disorder. *Arch Dis Child*. 2005;90(2):A78.
35. Costa R, Carmichael P, Colizzi M. To treat or not to treat: puberty suppression in childhood-onset gender dysphoria. *Nat Rev Urol*. 2016;13(8):456–462.
36. Vroenenraets LJ, Fredriks AM, Hannema SE, Cohen-Kettenis PT, de Vries MC. Early Medical treatment of children and adolescents with gender dysphoria: an empirical ethical study. *J Adolesc Health*. 2015;57(4):367–373.
37. Shumer DE, Spack NP. Paediatrics: transgender medicine—long-term outcomes from ‘the Dutch model’. *Nat Rev Urol*. 2015;12(1):12–13.
38. Spack NP, Edwards-Leeper L, Feldman HA, et al. Children and adolescents with gender identity disorder referred to a pediatric medical center. *Pediatrics*. 2012;129(3):418–425.
39. Kaltiala-Heino R, Sumia M, Tyolajarvi M, Lindberg N. Two years of gender identity service for minors: overrepresentation of natal girls with severe problems in adolescent development. *Child Adolesc Psychiatry Ment Health*. 2015;9:9.
40. Zucker KJ, Bradley SJ, Owen-Andersen A, Singh D. Puberty-blocking hormonal therapy for adolescents with gender identity disorder: a descriptive clinical study. *J Gay Lesbian Ment Health*. 2011;15:58–82.
41. de Vries AL, Doreleijers TA, Steensma TD, Cohen-Kettenis PT. Psychiatric comorbidity in gender dysphoric adolescents. *J Child Psychol Psychiatry*. 2011;52(11):1195–1202.
42. Chen M, Fuqua J, Eugster EA. Characteristics of referrals for gender dysphoria over a 13-year period. *J Adolesc Health*. 2016;58(3):369–371.
43. Meyenburg B. Gender dysphoria in adolescents: difficulties in treatment. *Prax Kinderpsychol Kinderpsychiatr*. 2014;63(6):510–522.
44. Khatchadourian K, Amed S, Metzger DL. Clinical management of youth with gender dysphoria in Vancouver. *J Pediatr*. 2014;164(4):906–911.
45. Olson J, Schrage SM, Belzer M, Simons LK, Clark LF. Baseline physiologic and psychosocial characteristics of transgender youth seeking care for gender dysphoria. *J Adolesc Health*. 2015;57(4):374–380.
46. Holt V, Skagerberg E, Dunsford M. Young people with features of gender dysphoria: demographics and associated difficulties. *Clin Child Psychol Psychiatry*. 2016;21(1):108–118.
47. Steensma TD, Zucker KJ, Kreukels BP, et al. Behavioral and emotional problems on the Teacher’s Report Form: a cross-national, cross-clinic comparative analysis of gender dysphoric children and adolescents. *J Abnorm Child Psychol*. 2014;42(4):635–647.
48. de Vries AL, Steensma TD, Cohen-Kettenis PT, VanderLaan DP, Zucker KJ. Poor peer relations predict parent- and self-reported behavioral and emotional problems of adolescents with gender dysphoria: a cross-national, cross-clinic comparative analysis. *Eur Child Adolesc Psychiatry*. 2016;25(6):579–588.
49. Rodríguez MF, Mora PG, Sánchez EM, Gidseen G. Características de los menores de edad con disforia de género que acuden a la unidad de tratamiento de identidad de género. *Rev Esp Salud Publica*. 2017;91(1):e1–e9.
50. Zucker KJ, Bradley SJ, Owen-Anderson A, et al. Demographics, behavior problems, and psychosexual characteristics of adolescents with gender identity disorder or transvestic fetishism. *J Sex Marital Ther*. 2012;38:151–189.
51. Clark TC, Lucassen MF, Bullen P, et al. The health and well-being of transgender high school students: results from the New Zealand adolescent health survey (Youth’12). *J Adolesc Health*. 2014;55(1):93–99.
52. Diemer EW, Grant JD, Munn-Chernoff MA, Patterson DA, Duncan AE. Gender identity, sexual orientation, and eating-related pathology in a national sample of college students. *J Adolesc Health*. 2015;57(2):144–149.
53. Van Der Miesen AI, Hurley H, De Vries AL. Gender dysphoria and autism spectrum disorder: a narrative review. *Int Rev Psychiatry*. 2016;28(1):70–80.

54. Lai M, Lombardo MV, Baron-Cohen S. Autism. *Lancet*. 2014; 383(9920):896–910.
55. Strang JF, Kenworthy L, Dominska A, et al. Increased gender variance in autism spectrum disorders and attention deficit hyperactivity disorder. *Arch Sex Behav*. 2014;43(8):1525–1533.
56. van der Miesen AIR, deVries ALC, Steensma TD, Hartman CA. Autistic symptoms in children and adolescents with gender dysphoria. *J Autism Dev Disord*. Epub 2017 Nov 30.
57. Havighurst RJ. *Developmental Tasks and Education*. Chicago: University of Chicago Press; 1948.
58. Scheiffge-Krenke I, Gelhaar T. Does successful attainment of developmental tasks lead to happiness and success in later developmental tasks? A test of Havighurst's (1948) theses. *J Adolesc*. 2008;31: 33–52.
59. Harvey RG, Stone Fish L. Queer youth in family therapy. *Fam Process*. 2015;54(3):396–417.
60. Gray SA, Sweeney KK, Randazzo R, Levitt HM. "Am I doing the right thing?" Pathways to parenting a gender variant child. *Fam Process*. 2016;55(1):123–138.
61. Mayer KH, Garofalo R, Makadon HJ. Promoting the successful development of sexual and gender minority youths. *Am J Public Health*. 2014;104(6):976–981.
62. Gonzalez KA, Rostosky SS, Odom RD, Riggle ED. The positive aspects of being the parent of an LGBTQ child. *Fam Process*. 2013;52(2):325–337.
63. Travers R, Bauer G, Pyne J, et al. Impacts of strong parental support for trans youth: a report prepared for children's aid society of Toronto and delisle youth services; 2012. Available from: <http://transpulseproject.ca>. Accessed October 4, 2017.
64. Wilson EC, Iverson E, Garofalo R, Belzer M. Parental support and condom use among transgender female youth. *J Assoc Nurses AIDS Care*. 2012;23(4):306–317.
65. Le V, Arayasirikul S, Chen YH, Jin H, Wilson EC. Types of social support and parental acceptance among transfemale youth and their impact on mental health, sexual debut, history of sex work and condomless anal intercourse. *J Int AIDS Soc*. 2016;19(2):20781.
66. Simons L, Schragger SM, Clark LF, Belzer M, Olson J. Parental support and mental health among transgender adolescents. *J Adolesc Health*. 2013;53(6):791–793.
67. Sumia M, Lindberg N, Tyläljärvi M, Kaltiala-Heino R. Early pubertal timing is common among adolescent girl-to-boy sex reassignment applicants. *Eur J Contracept Reprod Health Care*. 2016;21(6):483–485.
68. Hall-Lande JA, Eisenberg ME, Christenson SL, Neumark-Sztainer D. Social isolation, psychological health, and protective factors in adolescence. *Adolescence*. 2007;42(166):265–286.
69. Laursen B, Hartl AC. Understanding loneliness during adolescence: developmental changes that increase the risk of perceived social isolation. *J Adolesc*. 2013;36(6):1261–1268.
70. Kornienko O, Santos CE, Martin CL, Granger KL. Peer influence on gender identity development in adolescence. *Dev Psychol*. 2016;52(10):1578–1592.
71. Shevlin M, Murphy S, Mallett J, Stringer M, Murphy J. Adolescent loneliness and psychiatric morbidity in Northern Ireland. *Br J Clin Psychol*. 2013;52(2):230–234.
72. Harris RA, Qualter P, Robinson SJ. Loneliness trajectories from middle childhood to pre-adolescence: impact on perceived health and sleep disturbance. *J Adolesc*. 2013;36(6):1295–1304.
73. Kaltiala-Heino R, Frojd S. Correlation between bullying and clinical depression in adolescent patients. *Adolesc Health Med Ther*. 2011;2:37–44.
74. Shiffman M, VanderLaan DP, Wood H, Lumley MM, Lollis SP, Zucker KJ. Behavioral and emotional problems as a function of peer relationships in adolescents with gender dysphoria: a comparison with clinical and nonclinical controls. *Psychol Sex Orient Gender Divers*. 2016;3(1):27–36.
75. McGuire JK, Anderson CR, Toomey RB, Russell ST. School climate for transgender youth: a mixed method investigation of student experiences and school responses. *J Youth Adolesc*. 2010;39(10):1175–1188.
76. Bos H, Sandfort T. Gender nonconformity, sexual orientation, and Dutch adolescents' relationship with peers. *Arch Sex Behav*. 2015;44(5):1269–1279.
77. Toomey RB, Ryan C, Diaz RM, Card NA, Russell ST. Gender-nonconforming lesbian, gay, bisexual, and transgender youth: school victimization and young adult psychosocial adjustment. *Dev Psychol*. 2010;46(6):1580–1589.
78. Baams L, Beek T, Hille H, Zevenbergen FC, Bos HM. Gender non-conformity, perceived stigmatization, and psychological well-being in Dutch sexual minority youth and young adults: a mediation analysis. *Arch Sex Behav*. 2013;42(5):765–773.
79. Reisner SL, Veters R, Leclerc M, et al. Mental health of transgender youth in care at an adolescent urban community health center: a matched retrospective cohort study. *J Adolesc Health*. 2015;56(3):274–279.
80. Lawrence AA. Sexual orientation versus age of onset as bases for typologies (subtypes) for gender identity disorder in adolescents and adults. *Arch Sex Behav*. 2010;39(2):514–545.
81. Romeo KE, Kelley MA. Incorporating human sexuality content into a positive youth development framework: implications for community prevention. *Child Youth Serv Rev*. 2009;31(9):1001–1009.
82. Madkour AS, Farhat T, Halpern CT, Godeau E, Gabhainn SN. Early adolescent sexual initiation as a problem behavior: a comparative study of five nations. *J Adolesc Health*. 2010;47(4):389–398.
83. Savioja H, Helminen M, Frojd S, Marttunen M, Kaltiala-Heino R. Delinquency and sexual experiences across adolescence: does depression play a role? *Eur J Contracept Reprod Health Care*. 2017;22(4): 298–304.
84. Korchmaros JD, Ybarra ML, Mitchell KJ. Adolescent online romantic relationship initiation: differences by sexual and gender identification. *J Adolesc*. 2015;40:54–64.
85. Savioja H, Helminen M, Fröjd S, Marttunen M, Kaltiala-Heino R. Sexual experience and self-reported depression across adolescent years. *Health Psychol Behav Med*. 2015;3(1):337–347.
86. Bungener SL, Steensma TD, Cohen-Kettenis PT, de Vries ALC. Sexual and romantic experiences of transgender youth before gender-affirmative treatment. *Pediatrics*. 2017;139(3):e20162283.
87. Robinson JP, Espelage DL. Peer victimization and sexual risk differences between lesbian, gay, bisexual, transgender, or questioning and nontransgender heterosexual youths in grades 7–12. *Am J Public Health*. 2013;103(10):1810–1819.
88. Veale J, Watson RJ, Adjei J, Saewyc E. Prevalence of pregnancy involvement among Canadian transgender youth and its relation to mental health, sexual health, and gender identity. *Int J Transgend*. 2016;17(3–4):107–113.
89. Mitchell KJ, Ybarra ML, Korchmaros JD. Sexual harassment among adolescents of different sexual orientations and gender identities. *Child Abuse Negl*. 2014;38(2):280–295.
90. Dank M, Lachman P, Zweig JM, Yahner J. Dating violence experiences of lesbian, gay, bisexual, and transgender youth. *J Youth Adolesc*. 2014;43(5):846–857.
91. Meyer IH. Prejudice, social stress, and mental health in lesbian, gay, and bisexual populations: conceptual issues and research evidence. *Psychol Bull*. 2003;129(5):674–697.
92. Poteat T, Scheim A, Xavier J, Reisner S, Baral S. Global epidemiology of HIV infection and related syndemics affecting transgender people. *J Acquir Immune Defic Syndr*. 2016;72(3):S210–S219.
93. Gowen LK, Winges-Yanez N. Lesbian, gay, bisexual, transgender, queer, and questioning youths' perspectives of inclusive school-based sexuality education. *J Sex Res*. 2014;51(7):788–800.
94. Kosciw JG, Palmer NA, Kull RM. Reflecting resiliency: openness about sexual orientation and/or gender identity and its relationship to well-being and educational outcomes for LGBT students. *Am J Community Psychol*. 2015;55(1–2):167–178.

95. Poteat VP, Scheer JR, Mereish EH. Factors affecting academic achievement among sexual minority and gender-variant youth. *Adv Child Dev Behav.* 2014;47:261–300.
96. Theunissen MJ, de Man I, Verdonk P, Bosma H, Feron F. Are Barbie and Ken too cool for school? A case-control study on the relation between gender and dropout. *Eur J Public Health.* 2015;25(1):57–62.
97. Carvalho L. Childhood circumstances and the intergenerational transmission of socioeconomic status. *Demography.* 2012;49(3):913–938.
98. Jacob M, Cox SR. Examining transgender health through the international classification of functioning, disability, and health's (ICF) contextual factors. *Qual Life Res.* 2017;26(12):3177–3185.
99. Zucker KJ, Bradley SJ, Owen-Anderson A, Kibblewhite SJ, Cantor JM. Is gender identity disorder in adolescents coming out of the closet? *J Sex Marital Ther.* 2008;34(4):287–290.
100. Marchiano L. Outbreak: on transgender teens and psychic epidemics. *Psychol Perspect.* 2017;60:345–366.
101. Dhejne C, Lichtenstein P, Boman M, Johansson ALV, Långström N, Landén M (2011). Long-term follow-up of transsexual persons undergoing sex reassignment surgery: cohort study in Sweden. *PLoS One.* 6(2):e16885.
102. de Vries AL, Steensma T, Doreleijers T, Cohen-Kettenis P. Puberty suppression in adolescents with gender identity disorder: a prospective follow-up study. *J Sex Med.* 2011;8:2276–2283.
103. Hidalgo MA, Ehrensaft D, Tishelman AC, et al. The gender affirmative model: what we know and what we aim to learn. *Hum Dev.* 2013;56(5):285–290.
104. Steensma TD, Cohen-Kettenis PT. Gender transitioning before puberty? *Arch Sex Behav.* 2011;40:649–650.
105. Zucker KJ. Persistence and desistence in children and adolescents with gender variance: a comparative-developmental perspective. *J Am Acad Child Adolesc Psychiatry.* 2016;55(10):S80.

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# Sex- and Gender-specific Observations and Implications for COVID-19

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*Disclaimer: Due to the rapidly evolving nature of this outbreak, and in the interests of rapid dissemination of reliable, actionable information, this paper went through expedited peer review. Additionally, information should be considered current only at the time of publication and may evolve as the science develops. On February 11, 2020, the World Health Organization renamed the virus COVID-19.*

This is a critical time for medicine. As we observe the exponential rise in the number of individuals in the United States (US) who are infected with COVID-19, we try to prepare. Those in the front lines are trying to protect themselves and their patients with the daily ration of personal protective equipment and ventilation assistive equipment. Many individuals are racing against time to develop the needed novel treatments and vaccines. Public health officials work with what little information is known in order to make effective recommendations for prevention. However, at this pivotal time in history where every detail obtained by US health officials could be lifesaving, we are leaving out vital information.

Descriptive and observational data from Wuhan, China, note that the majority (51%-66.7%) of affected patients have been male. In addition, male sex was an independent risk factor associated with refractory disease and death (2.8% death rate for men vs 1.7% for female).<sup>1,2</sup> Currently, men represent 58% of COVID-19 infected patients in Italy and 70% of COVID-related deaths.<sup>3</sup> As coronavirus cases and deaths in the US continue to soar, sex-specific, comprehensive data with regard to US patients is not yet available.

Sex- and gender-based medicine (SGBM) incorporates how biological *sex* and the sociocultural aspects of *gender* affect health and illness. It acknowledges the interrelationship between sex and gender on health outcomes and promotes consideration of this variable in both research and clinical practice. SGBM has demonstrated significant evidence-based impact on cardiovascular disease, stroke,

sports medicine, and pain management, just to name a few

Sex and gender differences have been observed in infectious diseases previously. On a broad and critical scale, Nasir et al demonstrated that males with all-cause infectious sepsis have a 70% greater mortality than their female counterparts. Likewise, respiratory infection-specific epidemiological data from recent SARS (2003) and MERS (2012) outbreaks demonstrated a significantly higher case fatality rate in males as compared to females, 21.9% vs 13.2%.<sup>4,5</sup>

## Sex-specific Factors

Is the biological male more susceptible to an increased severity of infection? Or does the biological woman have natural protection against these viruses? In a 2017 *BMJ* article, Dr. Kyle Sue demonstrated the effect of sex hormones, estrogen and testosterone, on immune system response and engagement, resulting in a less robust immunologic response in males and subsequent increased morbidity and mortality from viral respiratory illnesses.<sup>6</sup> In addition, the X chromosome carries the largest number of immune-related genes in the human genome, perhaps also contributing to female's superior immune response (as well as a female preponderance in autoimmune diseases).<sup>7</sup>

Angiotensin-converting enzyme 2 (ACE2) and its role in viral transmission and associated morbidity has also been a topic of recent COVID-19 associated discussion. ACE2 receptors on pulmonary endothelium serve as a main entry point for coronavirus. Several previous animal models have demonstrated increased ACE2 activity in the male or ovariectomized model, suggesting a sex hormone influence.<sup>8</sup> The gene for the ACE2 receptor is also, interestingly, on the X chromosome.<sup>9</sup>

## Gender-specific Factors

Behavioral and cultural variables have also influenced current COVID-19 epidemiology. Smoking in particular has

been implicated as a significant contributor to disease severity, and gender-specific patterns are quite apparent here. The smoking rate in China is much higher in men than in women (288 million men vs 12.6 million women; 2018 data).<sup>10</sup> Likewise, in Italy, men are more likely to smoke than women at any age (1.12x to 1.7x, depending on age cohort; 2018 data).<sup>11</sup> Similar gender-specific trends are also present in the US, where 17.6% of smokers are men as compared to 13.6% of women.<sup>12</sup>

In addition, as the traditional caregivers and coordinators of care for their loved ones, women, particularly working mothers, tend to spend more time than men focused on medical issues related to both their own healthcare and that of their families.<sup>13</sup> In general, men are more likely to engage in health-related risks which, even prior to the COVID-19 pandemic, has been shown to result in higher rates of injury and disease.<sup>14</sup> Suen et al demonstrated in 2019 that being a middle-aged female was a protective factor with regard to hand hygiene knowledge and practice.<sup>15</sup>

### Implications for COVID-19 Management

As clinical researchers and pharmaceutical companies race to find an effective treatment strategy or vaccine for COVID-19, no sex- or gender-specific recommendations have been released with regard to the care and management of individuals affected by the novel coronavirus. Appreciating the weight of known sex- and gender-specific epidemiologic observations thus far, however, will be an important highlight of the information gathered to date. This, combined with what is already known about sex- and gender-based pulmonary and infectious disease pathology, may lead to important treatment breakthroughs that consider the sex and/or gender of patient in the comprehensive management plan.

In addition, the current pandemic weighs heavily on emotional wellness along with physical health. COVID-19 has also released a contagion of fear, anxiety, and stigma that will have implications for downstream mental health effects including post-traumatic stress disorder (PTSD). In general, the prevalence of PTSD has been shown to be substantially higher in women.<sup>16</sup> This has been re-substantiated in the setting of the COVID-19 outbreak in Wuhan, China, where women scored significantly higher on the PCL-5 (DSM-5 self-report measure for PTSD) than men, including higher rates of re-experiencing and negative alterations in cognition or mood.<sup>17</sup> Early recognition and effective treatment can ameliorate the burden of PTSD on both the individual and society, particularly for women who have been shown to have a modest advantage with regard to treatment response.<sup>18</sup>

### Future Considerations

Since 2016, the NIH has required inclusion of sex as a biological variable (SABV) in the study design for funded

#### *Population Health Research Capsule*

What do we already know about this issue?

*COVID-19 represents an unparalleled public health crisis. Like many other infectious diseases, sex and gender differences in health outcomes have already been globally observed.*

What was the research question?

*We sought to summarize and explain known COVID-19-related sex and gender differences.*

What was the major finding of the study?

*Sex and gender differences are having significant impacts on current COVID-19 health outcomes.*

How does this improve population health?

*This perspective brings attention to the importance of sex and gender; specifically as they impact current clinical management and research during the COVID-19 pandemic.*

research.<sup>19</sup> Recognizing the weight these variables play in disease outcome should result in universal adoption of SABV as scientists develop and engage in COVID-19 research. While men appear to be disproportionately affected and at risk for COVID-19 infection and associated morbidity, researchers should avoid the slippery slope of the traditional male-dominant test and analysis approach.

When considering pharmaceutical therapy advances, several previous studies have established that women are much more likely to experience adverse drug reactions (ADR) than men.<sup>20</sup> In fact, historically the majority of drugs recalled from the market were done so due to serious ADRs reported by women, quite often because they were never tested on women during clinical trials. Several sex-specific pharmacokinetic and pharmacodynamic differences have been well documented.<sup>21</sup>

Yes, time is of the essence right now; however, COVID-19 does not get a “pass” in considering sex and gender when gathering data or testing treatments. Sex and gender have already proven to be crucial variables in the short history of COVID-19; they will continue to be factors of marked importance. Making healthcare providers and researchers aware of their impact in real time will be crucial to the integration of susceptibility profiles and improving outcomes during an active public health crisis.

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## REFERENCES

1. Mo P, Xing Y, Xiao Y, et al. Clinical characteristics of refractory COVID-19 pneumonia in Wuhan, China. *Clin Infect Dis*. 2020. In Press.
2. Chen N, Zhou M, Dong X, et al. Epidemiological and clinical characteristics of 99 cases of 2019 novel coronavirus pneumonia in Wuhan, China: a descriptive study. *Lancet*. 2020;395(10223):507–13.
3. Remuzzi A, Remuzzi G. COVID-19 and Italy: what next? *Lancet*. 2020;S0140-6736(20)30627-9.
4. Channappanavar R, Fett C, Mack M, et al. Sex-based differences in susceptibility to severe acute respiratory syndrome coronavirus infection. *J Immunol*. 2017;198(10):4046–53.
5. Matsuyama R, Nishiura H, Kutsuna S, et al. Clinical determinants of the severity of Middle East respiratory syndrome (MERS): a systematic review and meta-analysis. *BMC Public Health*. 2016;16(1):1203.
6. Sue K. The science behind “man flu.” *BMJ*. 2017;359:j5560.
7. Schurz H, Salie M, Tromp G, et al. The X chromosome and sex-specific effects in infectious disease susceptibility. *Hum Genomics*. 2019;13(1):2.
8. Liu J, Ji H, Zheng W, et al. Sex differences in renal angiotensin converting enzyme 2 (ACE2) activity are 17 $\beta$ -oestradiol-dependent and sex chromosome-independent. *Biol Sex Differ*. 2010;1(1):6.
9. Patel SK, Velkoska E, Freeman M, et al. From gene to protein—experimental and clinical studies of ACE2 in blood pressure control and arterial hypertension. *Front Physiol*. 2014;5:227.
10. Cai H. Sex difference and smoking predisposition in patients with COVID-19. *Lancet Respir Med*. 2020. In Press.
11. Statista Research Department. “Number of Smokers by Age and Gender in Italy 2018.” *Statista*. Available at: [www.statista.com/statistics/501615/italy-smokers-by-age-and-gender/](http://www.statista.com/statistics/501615/italy-smokers-by-age-and-gender/). Accessed March 27, 2020.
12. National Institute on Drug Abuse. “Are There Gender Differences in Tobacco Smoking?” *NIDA*. Available at: [www.drugabuse.gov/publications/research-reports/tobacco-nicotine-e-cigarettes/are-there-gender-differences-in-tobacco-smoking](http://www.drugabuse.gov/publications/research-reports/tobacco-nicotine-e-cigarettes/are-there-gender-differences-in-tobacco-smoking). Accessed March 27, 2020.
13. “A Health Care Consumer Gender Gap.” *Managed Care Magazine*. Available at: [www.managedcaremag.com/archives/2016/8/health-care-consumer-gender-gap](http://www.managedcaremag.com/archives/2016/8/health-care-consumer-gender-gap). Accessed March 27, 2019.
14. Harris CR, Jenkins M. Gender differences in risk assessment: Why do women take fewer risks than men? *Judgm Decis Mak*. 2006;1(1):48-63.
15. Suen LKP, So ZYY, Yeung SKW, et al. Epidemiological investigation on hand hygiene knowledge and behaviour: a cross-sectional study on gender disparity. *BMC Public Health*. 2019;19(1):401.
16. K Farhood L, Fares S, Hamady C. PTSD and gender: Could gender differences in war trauma types, symptom clusters and risk factors predict gender differences in PTSD prevalence? *Arch Womens Ment Health*. 2018;21(6):725–33.
17. Liu N, Zhang F, Wei C. Prevalence and predictors of PTSS during COVID-19 Outbreak in China Hardest-hit Areas: Gender differences matter. *Psychiatry Res*. 2020. In Press.
18. Blain LM, Galovksi TE, Robinson T. Gender differences in recovery from posttraumatic stress disorder: a critical review. *Aggress Violent Behav*. 2010;15(6):463-74.
19. “NIH Policy on Sex as a Biological Variable.” *National Institutes of Health*, U.S. Department of Health and Human Services. Available at: [orwh.od.nih.gov/sex-gender/nih-policy-sex-biological-variable](http://orwh.od.nih.gov/sex-gender/nih-policy-sex-biological-variable). Accessed March 27<sup>th</sup>, 2020.
20. Tran C, Knowles SR, Liu BA, et al. Gender differences in adverse drug reactions. *J Clin Pharmacol*. 1998;38(11):1003–9.
21. Franconi F, Campesi I. Pharmacogenomics, pharmacokinetics and pharmacodynamics: interaction with biological differences between men and women. *Br J Pharmacol*. 2014;171(3):580–94.

# Considering how biological sex impacts immune responses and COVID-19 outcomes

Eileen P. Scully, Jenna Haverfield, Rebecca L. Ursin, Cara Tannenbaum and Sabra L. Klein

**Abstract** | A male bias in mortality has emerged in the COVID-19 pandemic, which is consistent with the pathogenesis of other viral infections. Biological sex differences may manifest themselves in susceptibility to infection, early pathogenesis, innate viral control, adaptive immune responses or the balance of inflammation and tissue repair in the resolution of infection. We discuss available sex-disaggregated epidemiological data from the COVID-19 pandemic, introduce sex-differential features of immunity and highlight potential sex differences underlying COVID-19 severity. We propose that sex differences in immunopathogenesis will inform mechanisms of COVID-19, identify points for therapeutic intervention and improve vaccine design and increase vaccine efficacy.

The COVID-19 pandemic, caused by the emergence of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), has resulted in millions of infections and hundreds of thousands of deaths worldwide. Human biological sex plays a fundamental role in heterogeneous COVID-19 outcomes. Sex, defined as male, female or intersex on the basis of sex chromosome complement, reproductive tissues (ovaries or testes) and sex steroid hormone (oestrogen, progesterone and testosterone) concentrations, is a multidimensional biological characteristic that shapes infectious disease pathogenesis. We discuss how sex differences in basic molecular and cellular mechanisms can be leveraged to define the immune response to infection with SARS-CoV-2.

## Sex differences in COVID-19 severity

The precise drivers of death, regardless of sex, in COVID-19 remain unknown. There appears to be a subset of patients in whom high levels of dysregulated inflammation lead to severe multisystem organ pathology<sup>1,2</sup>. A postviral inflammatory syndrome has also emerged in children with COVID-19 (REFS<sup>3,4</sup>). As a result, research on therapeutics has focused on both antiviral and immunomodulatory pathways<sup>2,5</sup>

with the goal of achieving an optimized balance in immune response induction and resolution. Unfortunately, most studies fail to consider the sex of the patients, which may mask therapeutic targets.

Evidence of sex differences in COVID-19 severity emerged in China, where hospital admissions and mortality were higher among males than females<sup>6–8</sup>. In South Korea, where community testing was widespread, females represented ~60% of those testing positive for SARS-CoV-2, suggesting that females acquire infection, despite having a lower case fatality rate (CFR)<sup>9,10</sup>. In the United States, where testing was prioritized for people with symptomatic disease, the diagnosis rates were similar in males and females, but males had 1.5 times higher mortality<sup>11</sup>.

A male bias in COVID-19 mortality is currently reported in 37 of the 38 countries that have provided sex-disaggregated data (FIG. 1a). Our analyses show that the average male CFR across 38 countries is 1.7 times higher than the average female CFR ( $P < 0.0001$ ) (male CFR 7.3 (95% CI 5.4–9.2); female CFR 4.4 (95% CI 3.4–5.5)), which is consistent with other reports<sup>12,13</sup>. There is increased risk of death for both sexes with advancing age, but at all ages above 30 years males have a significantly higher risk of

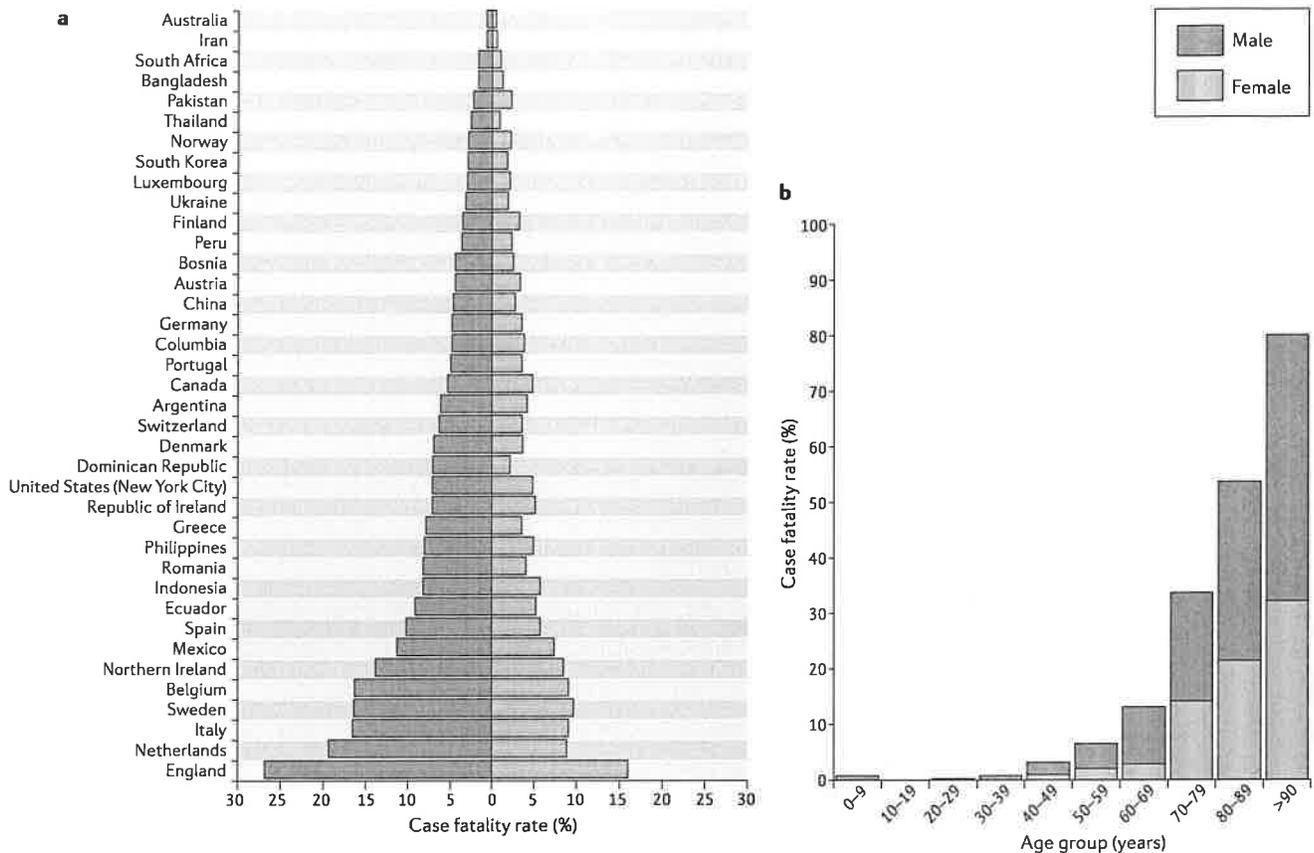
death than females ( $P < 0.05$ ) (FIG. 1b). A male predominance of deaths from COVID-19 is consistent with what was observed in the prior SARS<sup>14,15</sup> and Middle East respiratory syndrome (MERS)<sup>16</sup> epidemics (caused by SARS-CoV and MERS-CoV, respectively). Although gender-related social factors, including smoking, health care-seeking behaviours and some co-morbid conditions, may impact the outcomes of COVID-19 (REFS<sup>6,17</sup>) and contribute to male–female differences in disease severity, the cross-cultural emergence of increased risk of death for males points to biological risk determinants. In animal models of SARS-CoV infection, differences in mortality between male and female mice were observed and were attributed to steroid hormones<sup>18</sup>. Multiple dimensions of biological sex, including sex steroids, sex chromosomes and genomic and epigenetic differences between males and females, impact immune responses<sup>19–26</sup> and may affect responses to SARS-CoV-2 infection<sup>27</sup>.

## Ageing, sex and COVID-19

Although advancing age is associated with greater risk of death in both sexes, the male bias remains evident (FIG. 1b). An analysis of COVID-19 data from Italy, Spain, Germany, Switzerland, Belgium and Norway reveals that among all age groups older than 20 years, fatality rates are greater for males than females<sup>28</sup>. By contrast, male–female differences in the rate of confirmed SARS-CoV-2 infections are age dependent in all countries, being greater among females aged between 10 and 50 years and greater among males before the age of 10 years and after the age of 50 years<sup>28</sup>. The age-related male–female differences in confirmed cases of SARS-CoV-2 infections are consistent with reported confirmed cases of seasonal and pandemic influenza A virus infections in Australia and Japan<sup>29,30</sup>. We interpret these data to suggest that biological sex differences contribute to male-biased death, but gender-associated risk of exposure may affect rates of infection differently for males and females.

With a focus on biology, the impact of age on susceptibility to severe COVID-19 needs to be parsed, with both immunosenescence and dysregulation of innate immune responses as potential

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**Fig. 1 | Comparative analyses of COVID-19 case fatality rates by country, sex and age.** **a** | COVID-19 case fatality rates (CFRs) for males and females across 38 countries or regions reporting sex-disaggregated data on COVID-19 cases and deaths. CFR was calculated as the total number of deaths divided by the total number of cases for each sex multiplied by 100. The male CFR is higher than the female CFR in 37 of the 38 regions, with an average male CFR 1.7 times greater than the average female CFR ( $P < 0.0001$ , Wilcoxon signed rank test). **b** | Average COVID-19 CFRs for males and females stratified by age. The data represent 12 countries currently reporting sex- and age-disaggregated data on COVID-19 cases and deaths (Australia, Columbia, Denmark, Italy, Mexico, Norway, Pakistan, Philippines, Portugal, Spain, Switzerland and England). The COVID-19 CFR increases for both sexes with advancing age, but males have a significantly higher CFR than females at all ages from 30 years ( $P < 0.05$ , Wilcoxon signed rank test). The data were obtained from Global Health 50/50 and official government websites of each respective country on 7 May and 8 May 2020. For more information on the data source for a specific country, please contact the corresponding author.

mechanisms<sup>31,32</sup>. Biological sex differentially affects ageing of the immune system<sup>33</sup>, in part through changing concentrations of sex steroids<sup>34</sup>. In addition to reduced concentrations of sex steroids, an age-related mosaic loss of chromosome Y in leukocytes can alter transcriptional regulation of immunoregulatory genes<sup>35</sup>. Whether sex differences in the genomic signatures of aged immune cells translate to functional differences in the response to SARS-CoV-2 infection requires attention.

### Sex differences in immune responses

Biological sex affects innate and adaptive immune responses to self and foreign antigens, resulting in sex differences in autoimmunity as well as in responses to infections and vaccines<sup>36,37</sup>. Immune cell subsets have sex-specific patterns of gene expression, with most differentially

expressed genes found on autosomes, demonstrating sex-specific regulation of shared genetic material<sup>26</sup>. The sex chromosomes also directly contribute. Males are at higher risk of diseases caused by deleterious X-linked alleles. Incomplete inactivation of immunoregulatory genes on the X chromosome can also occur in females, which results in a gene dosage imbalance between sexes<sup>38,39</sup>. Incomplete X chromosome inactivation has been implicated in female-biased autoimmune diseases<sup>40</sup> and in vaccine efficacy<sup>41</sup>. The Y chromosome has immunoregulatory function, broadly impacting immune transcriptional profiles linked to autoimmune disease<sup>42</sup> and impacting outcomes of influenza virus and coxsackie virus infection in animals<sup>43,44</sup>. Sex-specific features of epigenomic organization also dictate differential availability of

transcriptional targets<sup>21,45</sup>. Superimposed on these genomic elements is the direct effect of sex steroid exposure. Oestrogens<sup>46,47</sup>, progesterone<sup>48-52</sup> and testosterone<sup>53</sup> have direct effects on immune cell function that are driven by the signalling of these hormones through their respective cellular receptors. The variation in sex steroid concentrations that occurs over the life course contributes to differences in immune profiles and disease susceptibility patterns at different ages<sup>20,52</sup>. Consistent with this variation, both sex and age contribute to unique transcriptional signatures of immune cells both at the baseline and after exposure to immunostimulants<sup>19,21,22</sup>. The summative effect is a sex-specific transcriptional regulatory network of genetic variants, epigenetic modifications, transcription factors and sex steroids that leads to a functional difference

in the immune response<sup>55,51</sup>. FIGURE 2 highlights intersections between SARS-CoV-2 infection and sources of sex bias in pathophysiology that warrant further investigation.

**Sex bias in SARS-CoV-2 infection**

**Virus entry receptors.** SARS-CoV-2 uses angiotensin-converting enzyme 2 (ACE2) as an entry receptor, with virus entry enhanced by cellular transmembrane serine protease 2 (TMPRSS2), which primes the spike protein of the virus<sup>55,56</sup>. ACE2 is an X chromosome-encoded gene that is downregulated by oestrogens<sup>57</sup> and exhibits tissue-specific expression patterns<sup>59</sup>. Differences in ACE2 expression may be driven by sex-differential expression of ACE2 variants<sup>58,60</sup>. ACE2 is associated with interferon gene expression<sup>61,62</sup>, which in turn shows sex-specific regulation. The cell-intrinsic regulation of ACE2 expression may change with age, in response to changing

levels of sex steroids, or following viral challenge. TMPRSS2 is regulated by androgen receptor signalling in prostate cells<sup>63</sup>. Initial investigations have not demonstrated a significant difference in TMPRSS2 mRNA expression in lung tissue from males and females, but it is unknown whether androgens may impact expression in the setting of infection with SARS-CoV-2 (REFS<sup>64,65</sup>) or whether the level of expression has an impact on SARS-CoV-2 burden. Further research is needed to determine whether sex-biased expression of ACE2, coupled with the regulation of TMPRSS2 by androgens, increases SARS-CoV-2 susceptibility of males compared with females.

**Interferons.** Innate sensing of viruses, production of interferons and activation of the inflammasome are the first line of defence against viruses<sup>66</sup>. In the case of SARS-CoV-2, where there is no pre-existing adaptive immune memory, the success of

this early antiviral response may be a determinant of disease outcome. Innate sensing of viral RNA by the pattern-recognition receptor Toll-like receptor 7 (TLR7) is sex biased, as TLR7 escapes X chromosome inactivation, resulting in greater expression in female immune cells; this has also been linked to sex differences in autoimmunity<sup>60,66</sup> and vaccine efficacy<sup>61</sup>. There is greater production of interferon- $\alpha$  (IFN $\alpha$ ) from plasmacytoid dendritic cells from adult females than from adult males<sup>67,68</sup>, an effect modulated by sex steroids<sup>69-71</sup>. In animal models of SARS-CoV infection, pretreatment with pegylated IFN $\alpha$  was associated with protection of lung tissue<sup>72</sup> but without consideration of biological sex. In SARS-CoV-2, emerging data suggest that there is aberrant activation of interferon responses but preserved chemokine signalling, which has been postulated to contribute to immunopathology<sup>73</sup>. Studies are needed

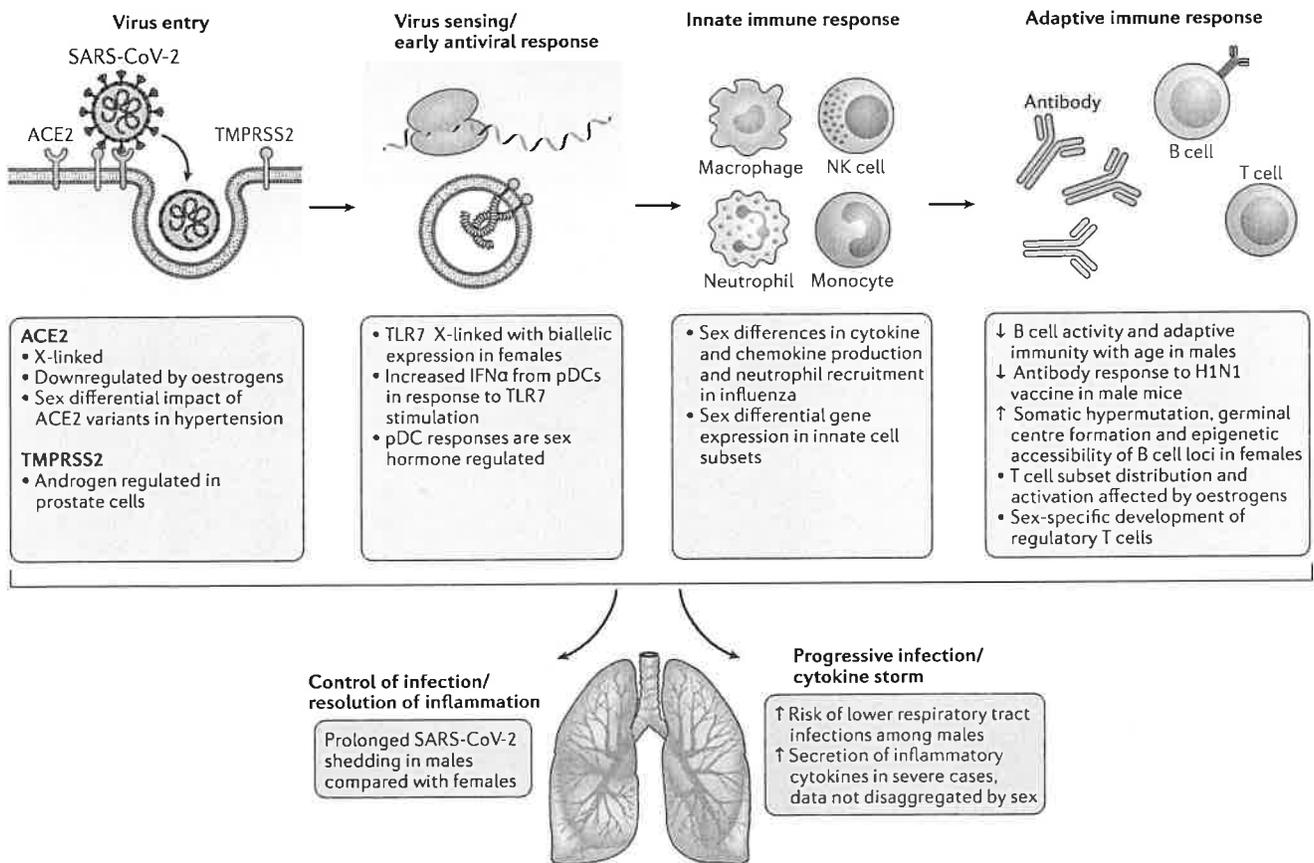


Fig. 2 | **Known sex differences that may impact immune responses to SARS-CoV-2 and COVID-19 progression.** An illustrative summary of the sequence of events in severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) infection and the associated immune responses. Broadly speaking (from left to right), there are the initial steps of virus entry, innate recognition of the virus with activation of antiviral programmes, the recruitment of innate immune cells and induction of an adaptive immune response. These major steps culminate either in successful control of infection and pathogen elimination or in a pathological inflammatory state. Sex differences that may be operative at multiple points along these pathways are indicated in the blue boxes. ACE2, angiotensin-converting enzyme 2; H1N1, H1N1 influenza virus; IFN $\alpha$ , interferon- $\alpha$ ; NK, natural killer; pDC, plasmacytoid dendritic cell; TLR7, Toll-like receptor 7; TMPRSS2, transmembrane protease serine 2.

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to determine whether differences in the magnitude or kinetics of the interferon response may contribute to a sex bias in the early control or severity of SARS-CoV-2 infection and may inform considerations of interferons as therapies for COVID-19 (REF.<sup>74</sup>). Early data suggest that male sex may be associated with a longer duration of viral detection, even within families<sup>75,76</sup>, raising the question of whether females have more efficient clearance of the virus. The rate of virus clearance will need to be assessed in evaluating the efficacy of innate and adaptive immune responses.

**Adaptive immunity.** Females generally mount greater antibody responses to viral infection and vaccination, albeit with higher levels of autoreactivity<sup>77</sup>. The mechanisms for sex differences in antibody production include oestrogenic enhancement of somatic hypermutation<sup>78</sup>, less stringent selection against autoreactive B cells<sup>77,79–82</sup> and sex differences in germinal centre formation<sup>83</sup> and in the epigenetic accessibility of B cell loci<sup>21</sup>. It is still unknown whether sex has an impact on antibody generation in SARS-CoV-2 infection. Early studies suggest that titres of antibodies to some viral epitopes are higher in patients with severe COVID-19 and that seroconversion may not be tightly linked to declining virus titres<sup>84,85</sup>. Ongoing studies evaluating the infusion of convalescent serum may provide answers as to the protective capacity of these antibodies<sup>86</sup>, but these studies are currently not considering biological sex. Generation of protective, neutralizing antibodies is a goal of vaccine development, with the cautionary note that in models of SARS-CoV vaccination some antibody responses induced potent inflammatory responses<sup>57</sup>. Persistence of antibodies, epitope targeting and non-neutralizing Fc-mediated antibody characteristics should be assessed with sex-stratified analyses. As vaccines are developed, the female bias towards both potent responses and adverse effects should be considered and sex-specific dosing should be tested, where appropriate<sup>87</sup>.

Sex impacts the development of regulatory T cells<sup>88–91</sup>, the distribution of lymphocyte subsets<sup>92</sup> and the overall quality of T cell responses<sup>93,94</sup>. In T cells, overexpression of X-encoded immune genes, including *CD40LG* and *CXCR3*, has been linked to incomplete X chromosome inactivation and T cell-specific epigenetic modifications of the X chromosome<sup>95,96</sup>. It is unknown whether T cell phenotypes contribute to COVID-19; data from the prior SARS outbreak did not link T cell

responses to outcomes in humans<sup>97</sup>, but mouse models suggest a role for CD4<sup>+</sup> T cells<sup>98</sup>. In patients with MERS, T cell responses were dysregulated<sup>99</sup>, but sex differences were not analysed. In the current pandemic, lymphopenia is associated with severe disease<sup>100,101</sup>, and early evidence suggests that the clinical markers of lymphocyte count may be lower in males and neutrophil–lymphocyte ratios may be higher<sup>17</sup>. Further work is needed to define the sex-differential role of T cells in acute infection, in acute lung injury phenotypes<sup>102</sup> and as potential vaccine targets.

**Severe infection and acute respiratory distress syndrome.** Severe cases of COVID-19 are typically marked by acute respiratory distress syndrome (ARDS), with respiratory failure requiring oxygen support and mechanical ventilation. The infection is primarily characterized by diffuse alveolar damage without a consistent pattern of cell infiltration<sup>75,103–105</sup>. The pathogenesis of ARDS involves the disruption of normal barrier function, inflammation and subsequent tissue repair. Whether there are sex-specific risks for ARDS and death from other causes, such as trauma, remains unknown<sup>106,107</sup>, although there is a suggestion of a higher risk of lower respiratory tract infections among males<sup>108</sup> and that steroid hormones modulate the immune response to respiratory viral pathogens<sup>109</sup>. In one cohort of patients with COVID-19, severe respiratory failure was associated with a pattern of inflammation, macrophage activation and depletion of lymphocytes that was distinct from bacterial infection<sup>110</sup>. There was a sex bias for severe COVID-19 not observed in the comparator group with bacterial infections<sup>110</sup>. Sex-differential production of IL-6 (REF.<sup>111</sup>), monocyte transcriptional patterns and inflammatory set point<sup>19,21,22</sup> could contribute to an enhanced risk of death in males and highlight the importance of sex-stratified analyses to guide deployment of safe and effective immunomodulatory therapeutics for males and females<sup>112</sup>.

## Conclusions

Emerging data demonstrating more favourable outcomes for community-dwelling adult females across age strata offer an immediate opportunity for comparative biology experiments to define features of COVID-19 pathogenesis and the associated immune response. The research pipeline should integrate sex as a biological variable in all stages, from fundamental research to preclinical

drug development, clinical trials and epidemiological analyses<sup>113</sup>. Considering the role of intersecting factors — including, but not limited to, gender, age, race and other identifying characteristics — is critical to understanding the biological and sociocultural factors contributing to heterogeneous COVID-19 outcomes. Sex is a driver of discovery and innovation<sup>114</sup>, and taking a sex-informed approach to COVID-19 research<sup>115</sup> and medicine<sup>116</sup> will uncover novel features of the host immune response to SARS-CoV-2 and ultimately result in more equitable health outcomes.

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1. Mehta, P. et al. COVID-19: consider cytokine storm syndromes and immunosuppression. *Lancet* **395**, 1033–1034 (2020).
2. Berlin, D. A., Gulick, R. M. & Martinez, F. J. Severe Covid-19. *N. Engl. J. Med.* <https://doi.org/10.1056/NEJMc2009575> (2020).
3. Verdoni, L. et al. An outbreak of severe Kawasaki-like disease at the Italian epicentre of the SARS-CoV-2 epidemic: an observational cohort study. *Lancet* **395**, 1771–1778 (2020).
4. Centers for Disease Control and Prevention. Multisystem inflammatory syndrome in children (MIS-C) associated with coronavirus disease 2019 (COVID-19). *CDC* <https://emergency.cdc.gov/han/2020/han00432.asp> (2020).
5. Vabret, N. et al. Immunology of COVID-19: current state of the science. *Immunity* <https://doi.org/10.1016/j.immuni.2020.05.002> (2020).
6. Chen, N. et al. Epidemiological and clinical characteristics of 99 cases of 2019 novel coronavirus pneumonia in Wuhan, China: a descriptive study. *Lancet* **395**, 507–513 (2020).
7. Guan, W. J. et al. Clinical characteristics of coronavirus disease 2019 in China. *N. Engl. J. Med.* **382**, 1708–1720 (2020).
8. Wu, Z. & McGoogan, J. M. Characteristics of and important lessons from the coronavirus disease 2019 (COVID-19) outbreak in China: summary of a report of 72314 cases from the Chinese center for disease control and prevention. *JAMA* <https://doi.org/10.1001/jama.2020.2648> (2020).
9. Dudley, J. P. & Lee, N. T. Disparities in age-specific morbidity and mortality from SARS-CoV-2 in China and the Republic of Korea. *Clin. Infect. Dis.* <https://doi.org/10.1093/cid/ciaa354> (2020).
10. Ministry of Health and Welfare of South Korea. Domestic occurrence status. *MOHW* [http://ncov.mohw.go.kr/bdBoardList\\_Real.do?brdId=1&brdGubun=11&ncvContSeq=&contSeq=&board\\_id=&gubun=](http://ncov.mohw.go.kr/bdBoardList_Real.do?brdId=1&brdGubun=11&ncvContSeq=&contSeq=&board_id=&gubun=) (2020).
11. NYC. COVID-19: data. *NYC.gov* <https://www1.nyc.gov/site/doh/covid/covid-19-data.page> (2020).

12. Jin, J.-M. et al. Gender differences in patients with COVID-19: focus on severity and mortality. *Front. Public Health* <https://doi.org/10.3389/fpubh.2020.00152> (2020).
13. Peckham, H. et al. Sex-bias in COVID-19: a meta-analysis and review of sex differences in disease and immunity. Preprint at *ResearchSquare* <https://doi.org/10.21203/rs.3.rs-23651/v2> (2020).
14. Karlberg, J., Chong, D. S. & Lai, W. Y. Do men have a higher case fatality rate of severe acute respiratory syndrome than women do? *Am. J. Epidemiol.* **159**, 229–231 (2004).
15. Leong, H. N. et al. SARS in Singapore—predictors of disease severity. *Ann. Acad. Med. Singap.* **35**, 326–331 (2006).
16. Alghamdi, I. G. et al. The pattern of Middle East respiratory syndrome coronavirus in Saudi Arabia: a descriptive epidemiological analysis of data from the Saudi Ministry of Health. *Int. J. Gen. Med.* **7**, 417–423 (2014).
17. Meng, Y. et al. Sex-specific clinical characteristics and prognosis of coronavirus disease-19 infection in Wuhan, China: a retrospective study of 168 severe patients. *PLoS Pathog.* **16**, e1008520 (2020).
18. Channappanavar, R. et al. Sex-based differences in susceptibility to severe acute respiratory syndrome coronavirus infection. *J. Immunol.* **198**, 4046–53 (2017).
19. Bongen, E. et al. Sex differences in the blood transcriptome identify robust changes in immune cell proportions with aging and influenza infection. *Cell Rep.* **29**, 1961–75 e4 (2019).
20. Ghosh, S. & Klein, R. S. Sex drives dimorphic immune responses to viral infections. *J. Immunol.* **198**, 1782–90 (2017).
21. Marquez, E. J. et al. Sexual-dimorphism in human immune system aging. *Nat. Commun.* **11**, 751 (2020).
22. Piasecka, B. et al. Distinctive roles of age, sex, and genetics in shaping transcriptional variation of human immune responses to microbial challenges. *Proc. Natl Acad. Sci. USA* **115**, E488–E497 (2018).
23. Schurz, H. et al. The X chromosome and sex-specific effects in infectious disease susceptibility. *Hum. Genomics* **13**, 2 (2019).
24. vom Steeg, L. G. & Klein, S. L. SexX matters in infectious disease pathogenesis. *PLoS Pathog.* **12**, e1005374 (2016).
25. Ober, C., Loisel, D. A. & Gilad, Y. Sex-specific genetic architecture of human disease. *Nat. Rev. Genet.* **9**, 911–922 (2008).
26. Schmiedel, B. J. et al. Impact of genetic polymorphisms on human immune cell gene expression. *Cell* **175**, 1701–15.e16 (2018).
27. Klein, S. L. et al. Biological sex impacts COVID-19 outcomes in the United States. *PLoS Pathog.* (in the press).
28. Marina S. & Piemonti L. Gender and age effects on the rates of infection and deaths in individuals with confirmed SARS-CoV-2 infection in six European countries. *Lancet* <https://doi.org/10.2139/ssrn.3576790> (2020).
29. Wong, K. C., Luscombe, G. M. & Hawke, C. Influenza infections in Australia 2009–2015: is there a combined effect of age and sex on susceptibility to virus subtypes? *BMC Infect. Dis.* **19**, 42 (2019).
30. Eshima, N. et al. Sex- and age-related differences in morbidity rates of 2009 pandemic influenza A H1N1 virus of swine origin in Japan. *PLoS ONE* **6**, e19409 (2011).
31. Koff, W. C. & Williams, M. A. Covid-19 and immunity in aging populations — a new research agenda. *N. Engl. J. Med.* <https://doi.org/10.1056/NEJMp2006761> (2020).
32. Nikolich-Zugich, J. et al. SARS-CoV-2 and COVID-19 in older adults: what we may expect regarding pathogenesis, immune responses, and outcomes. *GeroScience* **42**, 505–514 (2020).
33. Leng, S. X. & Margolick, J. B. Aging, sex, inflammation, frailty, and CMV and HIV infections. *Cell Immunol.* **348**, 104024 (2020).
34. Potluri, T. et al. Age-associated changes in the impact of sex steroids on influenza vaccine responses in males and females. *NPJ Vaccines* **4**, 29 (2019).
35. Dumanski, J. P. et al. Immune cells lacking Y chromosome have widespread dysregulation of autosomal genes. Preprint at *bioRxiv* <https://doi.org/10.1101/673459> (2020).
36. Klein, S. L. & Flanagan, K. L. Sex differences in immune responses. *Nat. Rev. Immunol.* **16**, 626–638 (2016).
37. Markle, J. G. & Fish, E. N. SexX matters in immunity. *Trends Immunol.* **35**, 97–104 (2014).
38. Carrel, L. & Brown, C. J. When the Lyonized chromosome roars: ongoing expression from an inactive X chromosome. *Philos. Trans. R. Soc. Lond. B Biol. Sci.* **372**, 20160355 (2017).
39. Tukkiainen, T. et al. Landscape of X chromosome inactivation across human tissues. *Nature* **550**, 244–248 (2017).
40. Souyris, M. et al. TLR7 escapes X chromosome inactivation in immune cells. *Sci. Immunol.* **3**, eaap8855 (2018).
41. Fink, A. L., Engle, K., Ursin, R. L., Tang, W. Y. & Klein, S. L. Biological sex affects vaccine efficacy and protection against influenza in mice. *Proc. Natl Acad. Sci. USA* **115**, 12477–82 (2018).
42. Case, L. K. et al. The Y chromosome as a regulatory element shaping immune cell transcriptomes and susceptibility to autoimmune disease. *Genome Res.* **23**, 1474–1485 (2013).
43. Kremontsov, D. N. et al. Genetic variation in chromosome Y regulates susceptibility to influenza A virus infection. *Proc. Natl Acad. Sci. USA* **114**, 3491–3496 (2017).
44. Robinson, D. P. et al. Sex chromosome complement contributes to sex differences in Coxsackievirus B3 but not influenza A virus pathogenesis. *Biol. Sex. Differ.* **2**, 8 (2011).
45. Golden, L. C. et al. Parent-of-origin differences in DNA methylation of X chromosome genes in T lymphocytes. *Proc. Natl Acad. Sci. USA* **116**, 26779–26787 (2019).
46. Straub, R. H. The complex role of estrogens in inflammation. *Endocr. Rev.* **28**, 521–574 (2007).
47. Peretz, J., Pekosz, A., Lane, A. P. & Klein, S. L. Estrogenic compounds reduce influenza A virus replication in primary human nasal epithelial cells derived from female, but not male, donors. *Am. J. Physiol. Lung Cell Mol. Physiol.* **310**, L415–L425 (2016).
48. Hall, O. J. & Klein, S. L. Progesterone-based compounds affect immune responses and susceptibility to infections at diverse mucosal sites. *Mucosal Immunol.* **10**, 1097–107 (2017).
49. Hall, O. J. et al. Progesterone-based therapy protects against influenza by promoting lung repair and recovery in females. *PLoS Pathog.* **12**, e1005840 (2016).
50. Hall, O. J. et al. Progesterone-based contraceptives reduce adaptive immune responses and protection against sequential influenza A virus infections. *J. Virol.* **91**, e02160-16 (2017).
51. Vom Steeg, L. G. & Klein, S. L. Sex steroids mediate bidirectional interactions between hosts and microbes. *Horm. Behav.* **88**, 45–51 (2017).
52. Vom Steeg, L. G. & Klein, S. L. Sex and sex steroids impact influenza pathogenesis across the life course. *Semin. Immunopathol.* **41**, 189–94 (2019).
53. Furman, D. et al. Systems analysis of sex differences reveals an immunosuppressive role for testosterone in the response to influenza vaccination. *Proc. Natl Acad. Sci. USA* **111**, 869–874 (2014).
54. Ellegren, H. & Parsch, J. The evolution of sex-biased genes and sex-biased gene expression. *Nat. Rev. Genet.* **8**, 689–698 (2007).
55. Hoffmann, M. et al. SARS-CoV-2 cell entry depends on ACE2 and TMPRSS2 and is blocked by a clinically proven protease inhibitor. *Cell* **181**, 271–280 (2020).
56. Matsuyama, S. et al. Enhanced isolation of SARS-CoV-2 by TMPRSS2-expressing cells. *Proc. Natl Acad. Sci. USA* **117**, 7001–7003 (2020).
57. Liu, J. et al. Sex differences in renal angiotensin converting enzyme 2 (ACE2) activity are 17beta-oestradiol-dependent and sex chromosome-independent. *Biol. Sex. Differ.* **1**, 6 (2010).
58. Zhang, Q. et al. Association of angiotensin-converting enzyme 2 gene polymorphism and enzymatic activity with essential hypertension in different gender: a case-control study. *Medicine* **97**, e12917 (2018).
59. Asselta, R., Paraboschi, E. M., Mantovani, A. & Duga, S. ACE2 and TMPRSS2 variants and expression as candidates to sex and country differences in COVID-19 severity in Italy. Preprint at *medRxiv* <https://doi.org/10.1101/2020.03.30.20047878> (2020).
60. Gibson, W. T., Evans, D. M., An, J. & Jones, S. J. ACE 2 coding variants: a potential x-linked risk factor for COVID-19 disease. Preprint at *bioRxiv* <https://doi.org/10.1101/2020.04.05.026633> (2020).
61. Ziegler, C. G. K. et al. SARS-CoV-2 receptor ACE2 is an interferon-stimulated gene in human airway epithelial cells and is enriched in specific cell subsets across tissues. *Cell* **181**, 1016–1035 (2020).
62. Sungnak, W. et al. SARS-CoV-2 entry factors are highly expressed in nasal epithelial cells together with innate immune genes. *Nat. Med.* **26**, 681–687 (2020).
63. Stopsack, K. H., Mucci, L. A., Antonarakis, E. S., Nelson, P. S. & Kantoff, P. W. TMPRSS2 and COVID-19: serendipity or opportunity for intervention? *Cancer Discov.* <https://doi.org/10.1158/2159-8290.CD-20-0451> (2020).
64. Baratchian, M. et al. No evidence that androgen regulation of pulmonary TMPRSS2 explains sex-discordant COVID-19 outcomes. Preprint at *bioRxiv* <https://doi.org/10.1101/2020.04.21.051201> (2020).
65. Iwasaki, A. A virological view of innate immune recognition. *Annu. Rev. Microbiol.* **66**, 177–196 (2012).
66. Souyris, M., Mejia, J. E., Chaumeil, J. & Guery, J. C. Female predisposition to TLR7-driven autoimmunity: gene dosage and the escape from X chromosome inactivation. *Semin. Immunopathol.* **41**, 153–64 (2019).
67. Berghofer, B. et al. TLR7 ligands induce higher IFN-alpha production in females. *J. Immunol.* **177**, 2088–2096 (2006).
68. Meier, A. et al. Sex differences in the Toll-like receptor-mediated response of plasmacytoid dendritic cells to HIV-1. *Nat. Med.* **15**, 955–959 (2009).
69. Griesbeck, M. et al. Sex differences in plasmacytoid dendritic cell levels of IRF5 drive higher IFN-alpha production in women. *J. Immunol.* **195**, 5327–5336 (2015).
70. Laffont, S. et al. X-Chromosome complement and estrogen receptor signaling independently contribute to the enhanced TLR7-mediated IFN-alpha production of plasmacytoid dendritic cells from women. *J. Immunol.* **193**, 5444–5452 (2014).
71. Seillet, C. et al. The TLR-mediated response of plasmacytoid dendritic cells is positively regulated by estradiol in vivo through cell-intrinsic estrogen receptor alpha signaling. *Blood* **119**, 454–464 (2012).
72. Haagmans, B. L. et al. Pegylated interferon-alpha protects type 1 pneumocytes against SARS coronavirus infection in macaques. *Nat. Med.* **10**, 290–293 (2004).
73. Blanco-Melo, D. N.-P. et al. Imbalanced host response to SARS-CoV-2 drives development of COVID-19. *Cell* <https://doi.org/10.1016/j.cell.2020.04.026> (2020).
74. Sallard, E., Lescure, F. X., Yazdanpanah, Y., Mentre, F. & Peiffer-Smadja, N. Type 1 interferons as a potential treatment against COVID-19. *Antivir. Res.* **178**, 104791 (2020).
75. Xu, K. et al. Factors associated with prolonged viral RNA shedding in patients with COVID-19. *Clin. Infect. Dis.* <https://doi.org/10.1093/cid/ciaa351> (2020).
76. Zheng, S. et al. Viral load dynamics and disease severity in patients infected with SARS-CoV-2 in Zhejiang province, China, January–March 2020: retrospective cohort study. *BMJ* **369**, m1443 (2020).
77. Sakiani, S., Olsen, N. J. & Kovacs, W. J. Gonadal steroids and humoral immunity. *Nat. Rev. Endocrinol.* **9**, 56–62 (2013).
78. Pauklin, S., Sernandez, I. V., Bachmann, G., Ramiro, A. R. & Petersen-Mahrt, S. K. Estrogen directly activates AID transcription and function. *J. Exp. Med.* **206**, 99–111 (2009).
79. Bynoe, M. S., Grimaldi, C. M. & Diamond, B. Estrogen up-regulates Bcl-2 and blocks tolerance induction of naive B cells. *Proc. Natl Acad. Sci. USA* **97**, 2703–2708 (2000).
80. Grimaldi, C. M., Cleary, J., Dagtas, A. S., Moussai, D. & Diamond, B. Estrogen alters thresholds for B cell apoptosis and activation. *J. Clin. Invest.* **109**, 1625–1633 (2002).
81. Grimaldi, C. M., Jeganathan, V. & Diamond, B. Hormonal regulation of B cell development: 17beta-estradiol impairs negative selection of high-affinity DNA-reactive B cells at more than one developmental checkpoint. *J. Immunol.* **176**, 2703–2710 (2006).
82. Hill, L., Jeganathan, V., Chinnasamy, P., Grimaldi, C. & Diamond, B. Differential roles of estrogen receptors alpha and beta in control of B-cell maturation and selection. *Mol. Med.* **17**, 211–220 (2011).
83. Zhao, R. et al. A GPR174-CCL21 module imparts sexual dimorphism to humoral immunity. *Nature* **577**, 416–20 (2020).
84. To, K. K. et al. Temporal profiles of viral load in posterior oropharyngeal saliva samples and serum antibody responses during infection by SARS-CoV-2: an observational cohort study. *Lancet Infect. Dis.* **20**, 565–74 (2020).

85. Wolfel, R. et al. Virological assessment of hospitalized patients with COVID-2019. *Nature* <https://doi.org/10.1038/s41586-020-2196-x> (2020).
86. Shen, C. et al. Treatment of 5 critically ill patients with COVID-19 with convalescent plasma. *JAMA* **323**, 1582–1589 (2020).
87. Flanagan, K. L., Fink, A. L., Plebanski, M. & Klein, S. L. Sex and gender differences in the outcomes of vaccination over the life course. *Annu. Rev. Cell Dev. Biol.* **33**, 577–99. (2017).
88. Dragin, N. et al. Estrogen-mediated downregulation of AIRE influences sexual dimorphism in autoimmune diseases. *J. Clin. Invest.* **126**, 1525–1537 (2016).
89. Polanczyk, M. J. et al. Cutting edge: estrogen drives expansion of the CD4+CD25+ regulatory T cell compartment. *J. Immunol.* **173**, 2227–2230 (2004).
90. Tai, P. et al. Induction of regulatory T cells by physiological level estrogen. *J. Cell Physiol.* **214**, 456–464 (2008).
91. Zhu, M. L. et al. Sex bias in CNS autoimmune disease mediated by androgen control of autoimmune regulator. *Nat. Commun.* **7**, 11350 (2016).
92. Page, S. T. et al. Effect of medical castration on CD4+CD25+ T cells, CD8+ T cell IFN-gamma expression, and NK cells: a physiological role for testosterone and/or its metabolites. *Am. J. Physiol. Endocrinol. Metab.* **290**, E856–E863 (2006).
93. Hewagama, A., Patel, D., Yariagadda, S., Strickland, F. M. & Richardson, B. C. Stronger inflammatory/cytotoxic T-cell response in women identified by microarray analysis. *Genes. Immun.* **10**, 509–516 (2009).
94. Yee Mon, K. J. et al. Differential sensitivity to IL-12 drives sex-specific differences in the CD8+ T cell response to infection. *Immunohorizons* **3**, 121–132 (2019).
95. Qu, K. et al. Individuality and variation of personal regulomes in primary human T cells. *Cell Syst.* **1**, 51–61 (2015).
96. Wang, J. et al. Unusual maintenance of X chromosome inactivation predisposes female lymphocytes for increased expression from the inactive X. *Proc. Natl Acad. Sci. USA* **113**, E2029–E2038 (2016).
97. Janice Oh, H. L., Ken-En Gan, S., Bertoletti, A. & Tan, Y. J. Understanding the T cell immune response in SARS coronavirus infection. *Emerg. Microbes Infect.* **1**, e23 (2012).
98. Chen, J. et al. Cellular immune responses to severe acute respiratory syndrome coronavirus (SARS-CoV) infection in senescent BALB/c mice: CD4+ T cells are important in control of SARS-CoV infection. *J. Virol.* **84**, 1289–1301 (2010).
99. Alosaimi, B. et al. MERS-CoV infection is associated with downregulation of genes encoding Th1 and Th2 cytokines/chemokines and elevated inflammatory innate immune response in the lower respiratory tract. *Cytokine* **126**, 154895 (2020).
100. Ruan, Q., Yang, K., Wang, W., Jiang, L. & Song, J. Clinical predictors of mortality due to COVID-19 based on an analysis of data of 150 patients from Wuhan, China. *Intensive Care Med.* **46**, 846–848 (2020).
101. Yang, X. et al. Clinical course and outcomes of critically ill patients with SARS-CoV-2 pneumonia in Wuhan, China: a single-centered, retrospective, observational study. *Lancet Respir. Med.* **8**, 475–481. (2020).
102. Halter, S. et al. T regulatory cells activation and distribution are modified in critically ill patients with acute respiratory distress syndrome: a prospective single-centre observational study. *Anaesth. Crit. Care Pain. Med.* **39**, 35–44 (2020).
103. Barton, L. M., Duval, E. J., Stroberg, E., Ghosh, S. & Mukhopadhyay, S. COVID-19 autopsies, Oklahoma, USA. *Am. J. Clin. Pathol.* **153**, 725–33. (2020).
104. Tian, S. et al. Pulmonary pathology of early-phase 2019 novel coronavirus (COVID-19) pneumonia in two patients with lung cancer. *J. Thorac. Oncol.* **15**, 700–704 (2020).
105. Tian, S. et al. Pathological study of the 2019 novel coronavirus disease (COVID-19) through postmortem care biopsies. *Mod. Pathol.* <https://doi.org/10.1038/s41379-020-0536-x> (2020).
106. Heffernan, D. S. et al. Gender and acute respiratory distress syndrome in critically injured adults: a prospective study. *J. Trauma* **71**, 878–883 (2011).
107. Liu, T. et al. The influence of sex on outcomes in trauma patients: a meta-analysis. *Am. J. Surg.* **210**, 911–921 (2015).
108. Falagas, M. E., Mourtzoukou, E. G. & Vardakas, K. Z. Sex differences in the incidence and severity of respiratory tract infections. *Respir. Med.* **101**, 1845–1863 (2007).
109. Robinson, D. P., Hall, O. J., Nilles, T. L., Bream, J. H. & Klein, S. L. 17beta-estradiol protects females against influenza by recruiting neutrophils and increasing virus-specific CD8 T cell responses in the lungs. *J. Virol.* **88**, 4711–4720 (2014).
110. Giamarellos-Bourboulis, E. J. et al. Complex immune dysregulation in COVID-19 patients with severe respiratory failure. *Cell Host Microbe* <https://doi.org/10.1016/j.chom.2020.04.009> (2020).
111. Naugler, W. E. et al. Gender disparity in liver cancer due to sex differences in MyD88-dependent IL-6 production. *Science* **317**, 121–124 (2007).
112. Bischof, E., Wolfe, J. & Klein, S. L. Clinical trials for COVID-19 should include sex as a variable. *J. Clin. Invest.* <https://doi.org/10.1172/JCI139306> (2020).
113. Canadian Institutes of Health Research. Why sex and gender need to be considered in COVID-19 research. *CIHR* <https://cihr-irsc.gc.ca/e/51939.html> (2020).
114. Klein, S. L. et al. Opinion: sex inclusion in basic research drives discovery. *Proc. Natl Acad. Sci. USA* **112**, 5257–5258 (2015).
115. Tannenbaum, C., Ellis, R. P., Eyssele, F., Zou, J. & Schiebinger, L. Sex and gender analysis improves science and engineering. *Nature* **575**, 137–46. (2019).
116. Bartz, D. et al. Clinical advances in sex- and gender-informed medicine to improve the health of all: a review. *JAMA Intern. Med.* **180**, 574–583 (2020).

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OPINION

# Biological sex impacts COVID-19 outcomes

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## Abstract

The current novel coronavirus disease 2019 (COVID-19) pandemic is revealing profound differences between men and women in disease outcomes worldwide. In the United States, there has been inconsistent reporting and analyses of male–female differences in COVID-19 cases, hospitalizations, and deaths. We seek to raise awareness about the male-biased severe outcomes from COVID-19, highlighting the mechanistic differences including in the expression and activity of angiotensin-converting enzyme 2 (ACE2) as well as in antiviral immunity. We also highlight how sex differences in comorbidities, which can be associated with both age and race, impact male-biased outcomes from COVID-19.

We are in the midst of a pandemic. Many of us predicted that the next “100 year pandemic” would be caused by an influenza A virus, like the H1N1 virus that caused the 1918 influenza pandemic. Instead, the current pandemic is caused by a novel  $\beta$ -coronavirus, the severe acute respiratory syndrome coronavirus 2 (SARS-CoV2). Currently, there are almost 2 million cases and over 100,000 deaths worldwide from the disease caused by this virus, called the novel coronavirus disease 2019 (COVID-19). Like the 1918 influenza pandemic [1], men are at greater risk of more severe COVID-19 outcomes than women, with both sex (i.e., biological differences) and gender (i.e., sociocultural and behavioral differences) playing fundamental roles.

The initial reports from China, followed by data from several countries in Europe, have highlighted that there are roughly similar numbers of confirmed SARS-CoV2 cases between men and women. The severity of COVID-19, as measured by hospitalization, admission to intensive care units, and rates of fatality, however, has consistently been 2-fold greater for men than women [2], with the Global Health 50/50 research initiative providing real-time sex-disaggregated data from most countries worldwide [3]. Unfortunately, despite the United States currently having the most COVID-19 cases in the world, considerably less attention has been paid to sex-disaggregation of data than in Europe and China.

We took this opportunity to evaluate the current situation in the US to both determine if similar patterns of male–female differences are observed and to document which states are or

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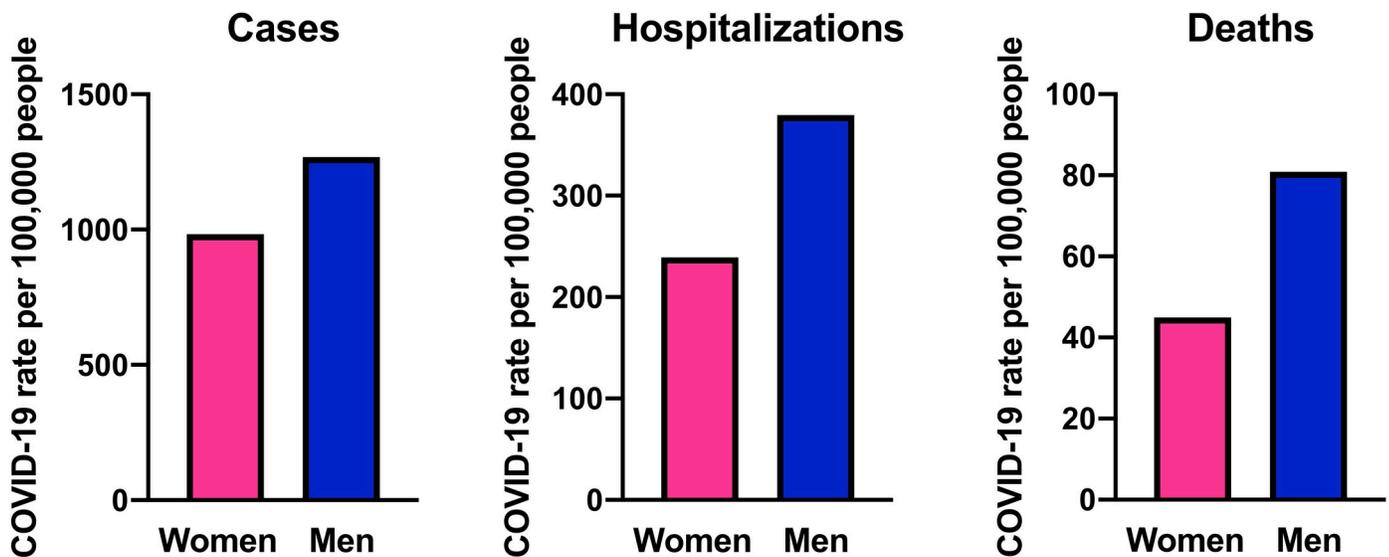
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are not disaggregating and analyzing data by sex. As of this writing, 26 states have more than 2,000 confirmed cases (<https://www.worldometers.info>) and from these states, only three (Louisiana, New Jersey, and Pennsylvania) have not sex-disaggregated cases of COVID-19. New York has the greatest number of COVID-19 cases in the US and is an epicenter of this pandemic. Of the data from the remaining 23 states, 7 states replicate the epidemiological pattern seen in New York City (NYC) (**Fig 1**) and elsewhere in the world [3], in which numbers of COVID-19 cases are similar between men and women. The other 16 states, however, suggest a female-bias exists in COVID-19 cases (i.e., 1 to 0.9/0.8 male to female ratio). This includes Washington state, which is another epicenter of the COVID-19 pandemic in the US. Of 167 COVID-19 cases from a Washington state long-term care facility, a majority of cases were women (68% of residents and 76% of healthcare workers) [4]. The total number of men and women among facility residents and healthcare workers was not provided and with women living longer than men and being more likely to work as healthcare providers [5], gender-associated factors may be involved [5].

Of the 26 states analyzed, only two counties within two different states (i.e., NYC and Bucks County, Pennsylvania) have reported rates of hospitalization broken down by sex, and both report greater rates of hospitalization from COVID-19 among men than women (**Fig 1**). Lastly, of the 26 states with more than 2,000 confirmed COVID-19 cases, only 13 (New York, Michigan, California, Illinois, Texas, Washington, Connecticut, Indiana, Colorado, Ohio, North Carolina, Wisconsin, and Alabama) have disaggregated fatalities from COVID-19 by sex and consistently show that fatality rates are 2-fold greater for men than women (**Fig 1**). Gender-associated factors have been reviewed elsewhere [2, 5]; thus, we seek to focus on biological mechanisms that could impact male–female differences in severe COVID-19 outcomes to call attention to sex-associated factors that could potentially provide novel insights into therapeutic interventions.

Angiotensin-converting enzyme 2 (ACE2) is a monocarboxypeptidase that counteracts the vasoconstrictor effects of angiotensin (Ang)-(1–8) by converting this octapeptide hormone to the vasodilator heptapeptide Ang-(1–7) [6]. In 2003, ACE2 was found to be the SARS-CoV (i.e., the virus that caused the 2002 to 2003 SARS outbreak) receptor [7, 8], with SARS-CoV2



**Fig 1. Sex-disaggregated numbers of COVID-19 cases, hospitalizations, and deaths per 100,000 people in NYC.** Data were accessed from <https://www1.nyc.gov/site/doh/covid/covid-19-data.page> on April 11, 2020. COVID-19, novel coronavirus disease 2019; NYC, New York City.

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binding ACE2 with even higher affinity [9]. ACE2 is expressed primarily in the kidney, heart, and testes, with the highest expression occurring in the kidney; it is, however, also expressed in the lungs at much lower levels [10]. The SARS-CoV2 virus S1 spike protein binds to the ACE2 receptor in alveolar epithelial cells of the lungs. ACE2 protein is expressed in a sex-specific manner in the mouse kidney; male mice have nearly 2-fold higher levels of renal ACE2 protein than female mice [11]. Furthermore, using the four core genotype mouse model in which gonadal sex (ovaries versus testes) is separated from the sex chromosome complement (XX versus XY) [12], we found renal ACE2 activity was greater in the male kidney regardless of the sex chromosome complement [11]. This sex difference in renal ACE2 activity was driven by estradiol reducing ACE2 activity regardless of the sex chromosome complement. These findings have implications for the observed sex differences in COVID-19 outcomes. It will be important to study the sex-specific regulation of ACE2 in the lung and other tissues involved in COVID-19 pathogenesis including the heart and brain [6] and also to investigate whether estrogens protects women from COVID-19 by reducing the expression levels of the receptor for the SARS-CoV2 virus.

The innate recognition and response to viruses as well as downstream adaptive immune responses during viral infections also differ between females and males [13]. We and others have illustrated that females generally mount greater inflammatory, antiviral, and humoral immune responses than males during viral infections [14], which contributes to better clearance of viruses, including SARS-CoV [15]. Enhanced immunity in females can, however, also result in greater immunopathology and tissue damage at later stages of viral disease, such as during influenza A virus infection [16]. To date, we have only identified two COVID-19 studies that have disaggregated and analyzed immunological outcome data by sex. In a published study of 168 patients with severe COVID-19 in Wuhan, China, it was reported that men were significantly more likely to remain hospitalized and die and less likely to be discharged from the hospital during the study period than women [17]. The male–female difference was most pronounced among individuals 60 years of age and older. In this study, the neutrophil to lymphocyte ratio and serum C-reactive protein concentrations were twice as high in male as in female COVID-19 patients, as well as in patients who died compared with patients who were discharged from the hospital (not disaggregated by sex) [17]. These data suggest that inflammatory immune responses and cell counts might be more elevated in men and associated with worse outcomes from COVID-19 than in women.

Mounting evidence suggests that humoral immune responses can be measured not only to confirm exposure to SARS-CoV2 but also to assess adaptive immune responses necessary for clearance of SARS-CoV2. As a result, convalescent plasma transfer studies are underway for compassionate care of severe COVID-19 patients [18], without, however, consideration of the sex of the donor. In a not yet peer-reviewed study of 331 patients with confirmed SARS-CoV2 infections in Wuhan, China, anti-SARS-CoV2 immunoglobulin G (IgG) responses were measured and compared among patients with either clinically diagnosed mild or severe disease. The sex distribution of recovering cases was 36% and 65% for men and women, respectively [19]. Among patients with mild COVID-19, anti-SARS-CoV2 IgG titers were similar between the sexes. In contrast, among patients with severe disease, women exhibited greater antibody responses than men, with production of antibodies at earlier phases of disease suggesting one possible immunological mechanism mediating better recovery from COVID-19 in women than men [19].

Development of mouse models for SARS-CoV2 will be instrumental for mechanistically assessing the causes of sex differences in the pathogenesis of COVID-19. In a mouse model of SARS-CoV infection, female mice had lower virus titers and less severe pulmonary damage from monocyte–macrophage infiltration and cytokine production, resulting in lower mortality

in female (20%) compared with male (80%) mice [15]; a sex distribution similar to that observed in human SARS [20]. Notably, the endogenous production of estradiol in female mice was important for this protection.

Comorbidities that are associated with more severe outcomes from COVID-19 in the US, include diabetes, obesity, hypertension, heart disease, chronic kidney disease, and chronic pulmonary disease. Notably, diabetes, obesity, and hypertension are the top three conditions associated with fatal COVID-19 cases in China and Italy [4, 21–23]. As of April 14, 2020, in New York, hypertension accounted for 56.8% and diabetes 42.4% of fatal cases [24]. In Louisiana, where New Orleans is the epicenter by death rate per capita, hypertension accounted for 59.8%, diabetes 38.1% and obesity 22.3% of fatal cases [25]. To date, no study has reported whether these comorbidities are influenced by sex or gender in COVID-19 patients. Biological (sex) as well as behavioral (gender) factors contribute to differences between men and women in these comorbidities [26]; the sex and gender-associated factors that underlie these comorbidities, however, have not been evaluated in context of COVID-19. There also has been no consideration about how sex intersects with age and race to further increase risk of severe COVID-19 outcomes in men, despite observations illustrating that older aged individuals [27] and African Americans [28] are also at risk for severe COVID-19 outcomes. For these reasons, we call on clinicians and epidemiologists to report data pertaining to comorbidities associated with COVID-19 disaggregated by sex, age, and race. We also emphasize the importance of considering the biological variable of sex when conducting basic science studies of COVID-19.

## References

1. Klein SL, Pekosz A, Passaretti C, Anker M, Olukoya P: Sex, gender and influenza. In. Geneva: World Health Organization; 2010: 1–58.
2. Gebhard C, Regitz-Zagrosek V, Neuhauser HK, Morgan R, Klein SL: Impact of sex and gender on COVID-19 outcomes in Europe. *Biology of Sex Differences* 11(1):29. <https://doi.org/10.1186/s13293-020-00304-9> PMID: 32450906
3. Sex, gender and COVID-19. [cited 2020 Apr 11]. <https://globalhealth5050.org/covid19/>
4. McMichael TM, Currie DW, Clark S, Pogojans S, Kay M, Schwartz NG, Lewis J, Baer A, Kawakami V, Lukoff MD et al: Epidemiology of Covid-19 in a Long-Term Care Facility in King County, Washington. *New England Journal of Medicine* March 27, 2020 <https://doi.org/10.1056/NEJMoa2005412> 2020
5. Wenham C, Smith J, Morgan R, Gender, Group C-W: COVID-19: the gendered impacts of the outbreak. *Lancet* 2020.
6. South AM, Diz D, Chappell MC: COVID-19, ACE2 and the Cardiovascular Consequences. *Am J Physiol Heart Circ Physiol* 2020.
7. Li W, Moore MJ, Vasilieva N, Sui J, Wong SK, Berne MA, Somasundaran M, Sullivan JL, Luzuriaga K, Greenough TC et al: Angiotensin-converting enzyme 2 is a functional receptor for the SARS coronavirus. *Nature* 2003, 426(6965):450–454. <https://doi.org/10.1038/nature02145> PMID: 14647384
8. Ge XY, Li JL, Yang XL, Chmura AA, Zhu G, Epstein JH, Mazet JK, Hu B, Zhang W, Peng C et al: Isolation and characterization of a bat SARS-like coronavirus that uses the ACE2 receptor. *Nature* 2013, 503(7477):535–538. <https://doi.org/10.1038/nature12711> PMID: 24172901
9. Chen Y, Guo Y, Pan Y, Zhao ZJ: Structure analysis of the receptor binding of 2019-nCoV. *Biochem Biophys Res Commun* 2020.
10. Donoghue M, Hsieh F, Baronas E, Godbout K, Gosselin M, Stagliano N, Donovan M, Woolf B, Robison K, Jeyaseelan R et al: A novel angiotensin-converting enzyme-related carboxypeptidase (ACE2) converts angiotensin I to angiotensin 1–9. *Circ Res* 2000, 87(5):E1–9. <https://doi.org/10.1161/01.res.87.5.e1> PMID: 10969042
11. Liu J, Ji H, Zheng W, Wu X, Zhu JJ, Arnold AP, Sandberg K: Sex differences in renal angiotensin converting enzyme 2 (ACE2) activity are 17beta-oestradiol-dependent and sex chromosome-independent. *Biology of sex differences* 2010, 1(1):6. <https://doi.org/10.1186/2042-6410-1-6> PMID: 21208466
12. Arnold AP, Cassis LA, Eghbali M, Reue K, Sandberg K: Sex Hormones and Sex Chromosomes Cause Sex Differences in the Development of Cardiovascular Diseases. *Arterioscler Thromb Vasc Biol* 2017, 37(5):746–756. <https://doi.org/10.1161/ATVBAHA.116.307301> PMID: 28279969

13. Klein SL, Flanagan KL: Sex differences in immune responses. *Nat Rev Immunol* 2016, 16(10):626–638. <https://doi.org/10.1038/nri.2016.90> PMID: 27546235
14. vom Steeg LG, Klein SL: Sex Matters in Infectious Disease Pathogenesis. *PLoS Pathog* 2016, 12(2): e1005374. <https://doi.org/10.1371/journal.ppat.1005374> PMID: 26891052
15. Channappanavar R, Fett C, Mack M, Ten Eyck PP, Meyerholz DK, Perlman S: Sex-Based Differences in Susceptibility to Severe Acute Respiratory Syndrome Coronavirus Infection. *J Immunol* 2017, 198(10):4046–4053. <https://doi.org/10.4049/jimmunol.1601896> PMID: 28373583
16. Vermillion MS, Ursin RL, Kuok DIT, Vom Steeg LG, Wohlgemuth N, Hall OJ, Fink AL, Sasse E, Nelson A, Ndeh R et al: Production of amphiregulin and recovery from influenza is greater in males than females. *Biol Sex Differ* 2018, 9(1):24. <https://doi.org/10.1186/s13293-018-0184-8> PMID: 30012205
17. Meng Y, Wu P, Lu W, Liu K, Ma K, Huang L, Cai J, Zhang H, Qin Y, Sun H et al: Sex-specific clinical characteristics and prognosis of coronavirus disease-19 infection in Wuhan, China: a retrospective study of 168 severe patients. *PLoS Pathog* 2020, <https://doi.org/10.1371/journal.ppat.1008520>
18. Bloch EM, Shoham S, Casadevall A, Sachais BS, Shaz B, Winters JL, van Buskirk C, Grossman BJ, Joyner M, Henderson JP et al: Deployment of convalescent plasma for the prevention and treatment of COVID-19. *J Clin Invest* 2020.
19. Zeng F, Dai C, Cai P, Wang J, Xu L, Li J, Hu G, Wang L: A comparison study of SARS-CoV-2 IgG antibody between male and female COVID-19 patients: a possible reason underlying different outcome between gender. *medRxiv preprint* 2020.
20. Karlberg J, Chong DS, Lai WY: Do men have a higher case fatality rate of severe acute respiratory syndrome than women do? *Am J Epidemiol* 2004, 159(3):229–231. <https://doi.org/10.1093/aje/kwh056> PMID: 14742282
21. Guan W-j, Ni Z-y, Hu Y, Liang W-h, Ou C-q, He J-x, Liu L, Shan H, Lei C-l, Hui DSC et al: Clinical Characteristics of Coronavirus Disease 2019 in China. *New England Journal of Medicine* February 28, 2020 <https://doi.org/10.1056/NEJMoa2002032> 2020
22. Bhatraju PK, Ghassemieh BJ, Nichols M, Kim R, Jerome KR, Nalla AK, Greninger AL, Pipavath S, Wurfel MM, Evans L et al: Covid-19 in Critically Ill Patients in the Seattle Region—Case Series. *New England Journal of Medicine* 2020.
23. Onder G, Rezza G, Brusaferro S: Case-Fatality Rate and Characteristics of Patients Dying in Relation to COVID-19 in Italy. *Jama* 2020.
24. New York State Department of Health COVID 19 Tracker. [cited 2020 Apr 11] <https://covid19tracker.health.ny.gov/views/NYS-COVID19-Tracker/NYSDOHCOVID-19Tracker-Fatalities?%3Aembed=yes&%3Atoolbar=no&%3Atabs=n>.
25. Louisiana Department of Health Coronavirus (COVID-19). [cited 2020 Apr 11]. <http://www.ldh.la.gov/coronavirus/>
26. Mauvais-Jarvis F, Bairey Merz CN, Barnes PJ, Brinton RD, Carrero J-J, DeMeo DL, de Vries GJ, Epperson CN, Govindan R, Klein SL et al: Sex and gender: Modifiers of health, disease and medicine. *Lancet* in press.
27. Chen N, Zhou M, Dong XP, Qu J, Gong F, Han Y, Qui Y, Wang J, Liu Y, Wei YQ et al: Epidemiological and clinical characteristics of 99 cases of 2019 novel coronavirus pneumonia in Wuhan, China: a descriptive study. *Lancet* 2020, 395(10223):507–513.
28. Yancy CW: COVID-19 and African Americans. *JAMA* 2020.

# The Inexorable Rise of Gender and the Decline of Sex: Social Change in Academic Titles, 1945–2001

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More than 30 million titles of “academic” articles, from the years 1945–2001, were surveyed for occurrences of the words *sex* and *gender*. At the beginning of this period, uses of *gender* were much rarer than uses of *sex*, and often used in the sense of a grammatical category. By the end of this period, uses of *gender* outnumbered uses of *sex* in the social sciences, arts, and humanities. Within the natural sciences, there was now more than 1 use of *gender* for every 2 uses of *sex*. The beginnings of this change in usage can be traced to Money’s introduction of the concept of “gender role” in 1955 (J. Money, 1955). However, the major expansion in the use of *gender* followed its adoption by feminists to distinguish the social and cultural aspects of differences between men and women (*gender*) from biological differences (*sex*). Since then, the use of *gender* has tended to expand to encompass the biological, and a *sex/gender* distinction is now only fitfully observed.

**KEY WORDS:** sex; gender; gender role; feminism.

## INTRODUCTION

In *The Mill on the Floss*, the novelist George Eliot (Mary Ann Evans) (1860) wrote “Public opinion, in these cases, is always of the feminine gender—not the world, but the world’s wife . . .” As this literary example shows, the use of *gender* as a synonym for *sex* has a long pedigree and is not a recent aberration as is sometimes claimed. The *Oxford English Dictionary* quotes uses of *gender* for *sex* from the fifteenth century, although in the first edition of the Dictionary in 1899 this usage was described as jocular. From the 1950s, however, a trickle of nonjocular uses of *gender* began to appear in the academic literature and, by the 1980s, this trickle had become a flood.

The most important factor was the adoption of *gender* in the 1970s by feminist scholars as a way of distinguishing “socially constructed” aspects of male–female differences (*gender*) from “biologically determined” aspects (*sex*). This distinction is now only fitfully respected, and *gender* is often used as a simple synonym of *sex*. The rise of *gender* has been accompanied by complaints

that the word should refer only to grammatical categories (Fletcher, 1991; Goodhart, 1992; Smyth, 1968) or to socially but not biologically determined differences (Fishman, Wick, & Koenig, 1999; Kim & Nafziger, 2000; Lewine, 1994; Pearson, 1996; Walker & Cook, 1998; Wilson, 2000).

In an attempt to document these changes in usage, I surveyed the titles of over 30 million academic articles, from the years 1945–2001, for occurrences of the words *sex* and *gender*. This work extends a similar analysis by Haig (2000) for the period 1988–1999. The quantitative analysis is followed by a discussion of the shifts in meaning of *gender* over this period.

## METHOD

The ISI Web of Science<sup>®</sup> is formed from the amalgamation of three databases: the Science Citation Index—Expanded (SCI) contains titles from 1945 until the present; the Social Sciences Citation Index (SSCI) contains titles from 1956 until the present; and the Arts & Humanities Citation Index (AHCI) contains titles from 1975 until the present. The contents of the three databases have considerable overlap. Thus, an article may be indexed in more

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than one database. The combined database contains over 30 million titles for the years 1945–2001.

The titles of all English-language articles in the Web of Science for the years 1945–2001 were searched for occurrences of “sex” and “gender.” Such searches retrieved titles that contain hyphenated constructs such as “sex-specific” or “gender-significant” but did not retrieve titles in which sex and gender appear as derived forms, such as “sexual” or “gendered.” In total, the searches found 59,262 sex-containing titles and 29,941 gender-containing titles (with some titles belonging to both categories).

The titles of articles in non-English languages often appear in the Web of Science as English translations. These articles were excluded from searches for “sex” and “gender,” but I was unable to exclude them from counts of the number of titles being searched. Therefore, whenever I calculated proportions of titles containing “sex” or “gender,” the numerator contained English language articles only, but the denominator contained articles in all languages. Most articles in the database are in English, but I do not have a measure of how the proportion of non-English articles has changed over time. This is a potentially confounding factor in the interpretation of Figs. 2–4. However, overall trends were little affected by including or excluding non-English language articles from searches.

Both sex and gender have uses for which the other would rarely, if ever, be substituted. Since the late 1970s, gender in its grammatical sense has contributed a small minority of all gender-containing titles. Of somewhat greater significance are biological uses of sex for which gender is not used (e.g., sex, in the sense of genetic recombination; sex chromosomes; sex hormones). But probably the most important uses of “sex” for which “gender” is not a synonym relate to copulation and other sexual activities (e.g., sex, in the sense of sexual intercourse; anal sex; safe sex; sex worker; sex slave). Such uses contribute a relatively small proportion of sex-containing titles in SCI, but a much greater proportion in SSCI and AHCI. (Analysis of a small sample suggests that about half of all sex-containing titles in SSCI and AHCI for the year 2001 belong in this category. I suspect that the advent of AIDS has increased the frequency of titles in this category, especially in SSCI, but I did not undertake a formal analysis.)

My analysis focused on usage in titles, but fashions in titles may not entirely reflect the content of articles. Articles may use gender in the text without it appearing in the title, or vice versa. In some cases, titles appeared to reflect editorial rather than authorial choices. For example, articles by Rothman and Liess (1976) and Harlap (1979) contained the first nongrammatical uses of gender in titles from the *New England Journal of Medicine* (with the en-

suaging correspondence, Harlap [1979] contributed six of 33 gender-containing titles in SCI for 1979). In both articles, however, gender appeared in the title but not in the text, where sex was used. Occasionally, tensions came to the surface. Ounsted and Taylor (1972) wrote in their edited volume, “As between the words ‘sex’ and ‘gender’ even, while preferring the scope of the latter term, we have accepted our authors’ preference for the former where they wish it” (p. vi). Despite this ecumenical principle, “gender” was used in the title of two chapters that used “sex” throughout the text, and the title of a third chapter contained “gender” in the Table of Contents but “sex” at the head of the chapter.

The journals indexed in the databases varied from year to year. Therefore, changes in the number of titles containing a particular word will depend only partly on changes in usage, but will also be influenced by what was and was not included in the database for a particular year. For example, several psychology journals that were covered by both SCI and SSCI in 1977 were no longer covered by SCI in 1978 (e.g., *Child Development*, *Journal of Personality and Social Psychology*). As a result, 28 SCI titles contained gender in 1977 but only nine contained gender in 1978. The latter figure would have been increased to 25 if titles, now included only in SSCI, had still been included in SCI. Thus, a corporate decision at the Institute for Scientific Information<sup>®</sup> accounted for most of the seemingly anomalous increase in the sex-to-gender ratio of SCI titles in 1978 (Fig. 1), although this factor does not explain the rebound to 33 SCI titles containing gender in 1979.

As another example of changes in coverage, the number of articles included in SSCI increased by 13% between 1994 and 1995. This increase appears to be due to the inclusion of additional journals not previously covered by SSCI. It is possible that the substantial increase in the proportion of titles containing gender that occurred in 1995 (Fig. 3), and the subsequent plateau in this measure, reflected a change in the composition of SSCI rather than any change of usage in the academic community; however, a 25% increase in the number of articles covered by SSCI between 1975 and 1976 does not appear to have affected the relative occurrence of sex and gender. An ideal analysis would separate effects of changes in usage from changes in coverage, but I doubt that such an analysis would change the gross trends detected by the present much simpler, and more easily replicable, analysis.

The databases did not contain book titles, except in book reviews, nor the texts of articles and books. Moreover, it is probable that use of gender in the titles of articles in indexed journals, at first, lagged behind conversational use. My quantitative analysis is restricted to indexed titles. The narrative that follows the quantitative analysis makes

use of other published sources that came to light in my readings.

**RESULTS**

Prior to the late 1960s, nongrammatical uses of gender were exceedingly rare. For the years 1945–1959, 1,685 (.14%) SCI titles out of 1,162,909 contained sex but only five (.0004%) contained gender. Of these, three used gender in a grammatical sense and two were sexological articles, both by Money (Money, 1955; Money, Hampson, & Hampson, 1957).

For the years 1960–1966, 2,094 (.17%) out of 1,253,631 titles in SCI contained sex and eight (.0006%) contained gender, of which three were grammatical uses and five were sexological (including three articles by Money and coauthors). For these same years, 819 (.24%) out of 353,069 titles in SSCI contained sex and 12 (.004%) contained gender (including four articles by Money and coauthors). Four gender-containing titles appeared in both SCI and SSCI.

Figure 1 presents changes in the ratio of sex-containing and gender-containing titles for the years 1966–2001

(from 1975 for AHCI). The ratio is expressed on a logarithmic scale because this is unbiased with respect to whether sex or gender appears in the denominator (i.e., 1:2 and 2:1 ratios are represented as equidistant from 1:1). There was substantial noise in the signal for the early years of this series because of the small number of gender-containing titles.

Some general observations can be made. The sex-to-gender ratio has always been lower in SSCI than in SCI, but this became more pronounced after 1973 when the SSCI initiated a sustained decline in the sex-to-gender ratio, which then leveled off in the 1990s (by which time gender-containing titles outnumbered sex-containing titles). A similar decline in the sex-to-gender ratio for SCI titles did not start until about 1980 and is still continuing. The ratio for AHCI followed closely that of SSCI, but with a slightly stronger preference for gender over sex. The first year for which gender-containing titles exceeded sex-containing titles was 1987 for AHCI and 1990 for SSCI. Sex-containing titles have always outnumbered gender-containing titles in SCI.

In 1993, the United States Food and Drug Administration (FDA) issued a Guideline requiring studies of “gender differences” in all new drug applications (Kessler,

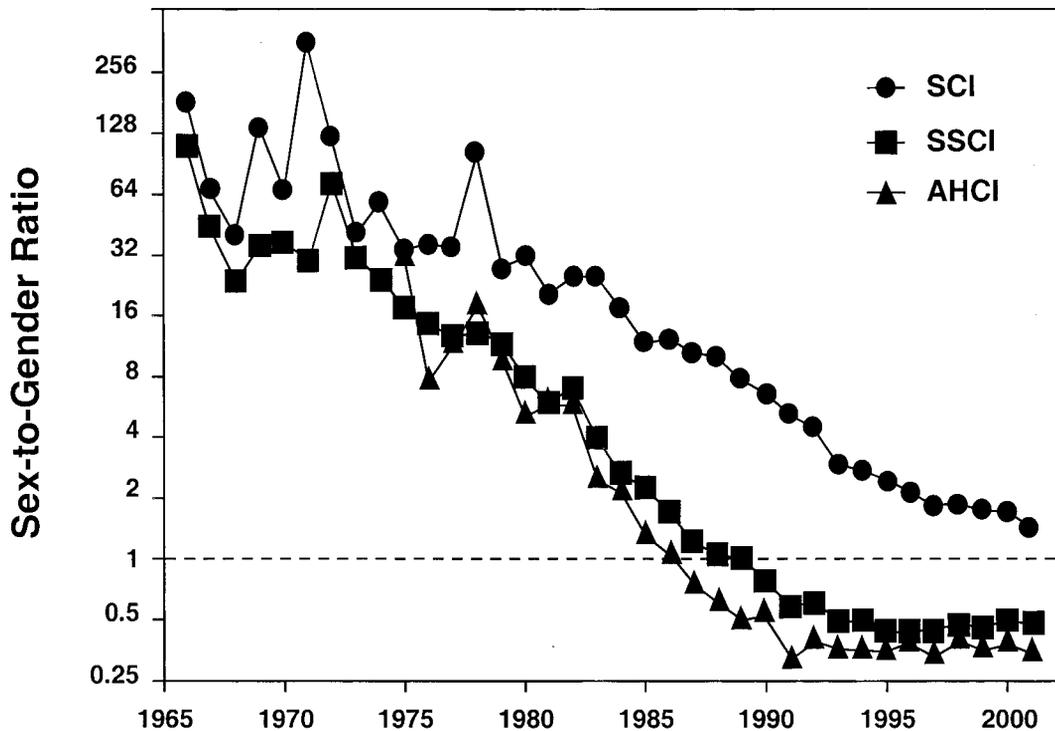


Fig. 1. The ratio of titles containing sex to titles containing gender for all articles in the Science Citation Index (SCI), Social Sciences Citation Index (SSCI), and Arts & Humanities Citation Index (AHCI).

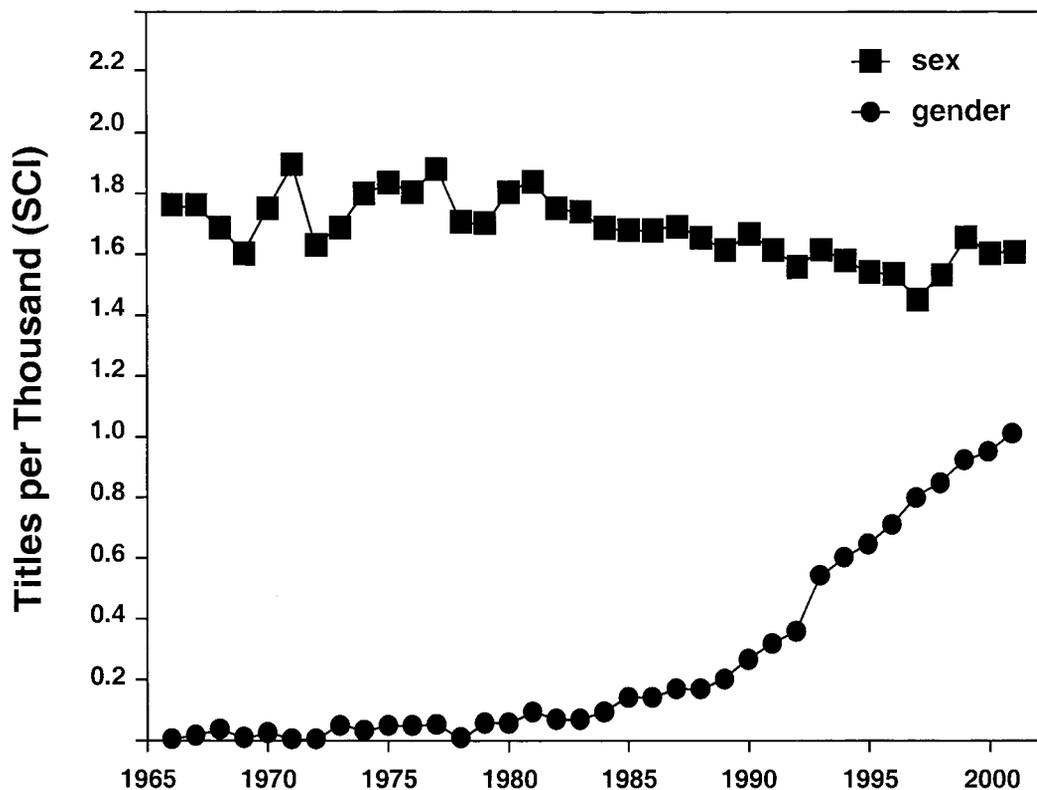


Fig. 2. Proportion of titles in the Science Citation Index containing the word *sex* and proportion containing the word *gender*.

1993). The decline in the sex-to-gender ratio in SCI began years before this Guideline and was not markedly affected by it, although there was a small acceleration in the decline in 1993. If the ratio of titles in SCI containing “sex differences” to titles containing “gender differences” is considered, this subsidiary ratio had been declining more rapidly than the overall sex-to-gender ratio since about 1985 (data not shown). Titles containing “gender differences” first outnumbered titles containing “sex differences” in 1994 (i.e., in the year following the Guideline) and have done so in every year since (except 1995).

Figures 2–4 present changes in the proportion of articles containing sex and gender (expressed as occurrences per thousand titles) for each of the three databases for the same years as covered in Fig. 1. Note that the vertical scales have been adjusted to reflect the fact that the proportion of titles containing sex and/or gender was far higher in SSCI than SCI, with AHCI intermediate. For SCI (Fig. 2), there was a small increase in the proportion of titles containing sex and/or gender over this period, from 1.8 per 1,000 in 1966 to 2.7 per 1,000 in 2001. From about 1980, gender began a steady increase in frequency, partly at the

expense of sex. The FDA Guideline on the evaluation of gender differences was possibly responsible for the extra large jump in the frequency of gender in 1993.

For SSCI (Fig. 3), there was a dramatic increase in the proportion of titles containing sex and/or gender from 3.4 per 1,000 in 1966 to 16.3 per 1,000 in 2001. Up until 1980, both gender and sex increased in tandem. During the 1980s, gender began a rapid rise in frequency at the expense of sex. From 1990, the frequency of sex has been roughly constant (as has the frequency of gender from 1995). Thus, there is a hint that the relative interest in sex-related subjects has reached a plateau in the social sciences.

The AHCI database contains data from 1975 until present. Figure 4 shows a dramatic increase over this period in the proportion of titles containing sex and/or gender, from .6 per 1,000 in 1975 to 7.1 per 1,000 in 2001, with a slight lag relative to the corresponding increase in SSCI (on the other hand, the fall in the sex-to-gender ratio in AHCI was slightly ahead of the decline in SSCI). The rapid rise in the frequency of gender began in about 1982, with a slower rise of sex from the late 1980s. Unlike SCI

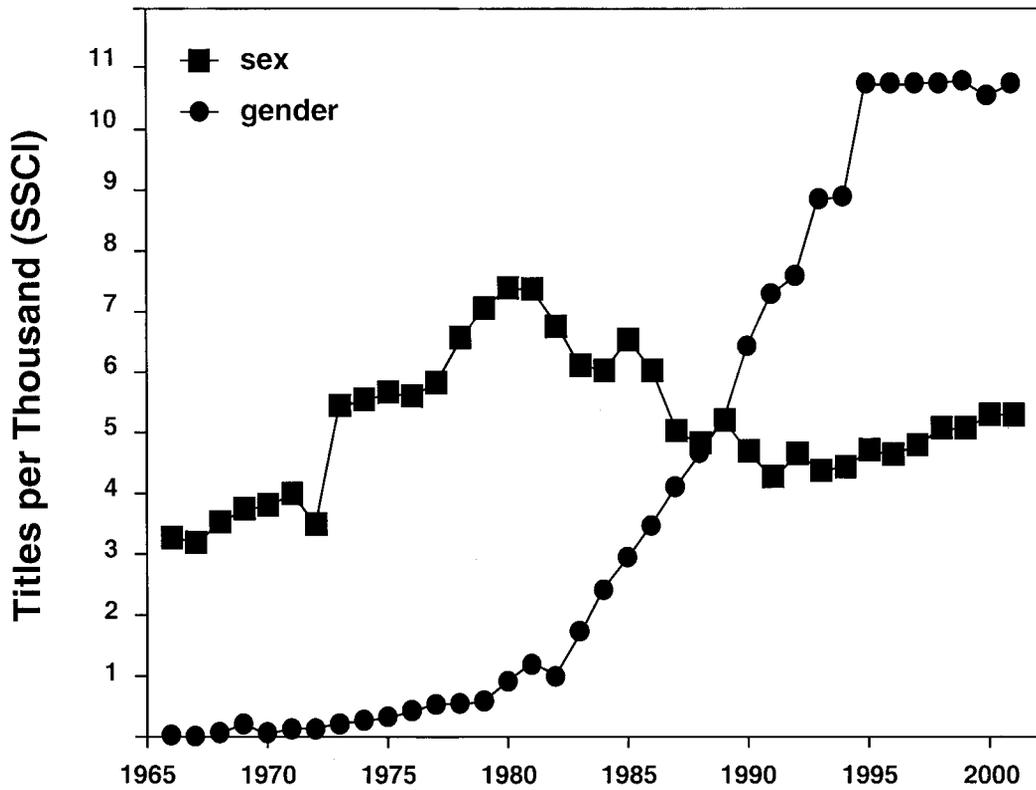


Fig. 3. Proportion of titles in the Social Sciences Citation Index containing the word *sex* and proportion containing the word *gender*.

and SSCI, the rapid rise of gender was not associated with a decline in the frequency of sex.

**DISCUSSION**

**Sexological Origins**

The first title in SCI to use gender in a nongrammatical sense was *Hermaphroditism, gender and precocity in hyperadrenocorticism: Psychologic findings* (Money, 1955). This article introduced the concept of a gender role: “The term *gender role* is used to signify all those things that a person says or does to disclose himself or herself as having the status of boy or man, girl or woman, respectively. It includes, but is not restricted to, sexuality in the sense of eroticism.” This was one of a series of papers by Money and his collaborators that appeared in the *Bulletin of the Johns Hopkins Hospital* during that year. Other papers in the series employed the concept of gender role (Money, Hampson, & Hampson, 1955a, 1955b), without gender appearing in their titles.

The juxtaposition of *role* and *status* in the above definition suggests that Money was influenced by Parson’s concept of *sex roles*. Money received his PhD in 1952 from the Department of Social Relations at Harvard University and listed Parsons among his teachers (Money, 1986, p. 5). For Parsons (1949), a status was “any patterned definition of who and what a person is” whereas a role was “the dynamic aspect of status, the behavior counterpart of the ideal or expected position defined by a status” (p. 43). Uses of *sex role* from the 1940s can be found in Parsons (1940, 1942), Cottrell (1942), and Mead (1949, p. 73). One of the many ironies to emerge from my analysis is that discussion of *sex roles* is now a staple of sociobiology (e.g., Vincent, 1994) without awareness of the term’s origin in sociology.

Money (1996) later wrote that he imported the term gender into sexological science “to make it possible to write about people who came into one’s office as either male or female, but of whom it could not be said that their sex role in the specific genital sense was either male or female insofar as they had a history of birth defect of the sex organs.” He then continued grandiloquently, “The

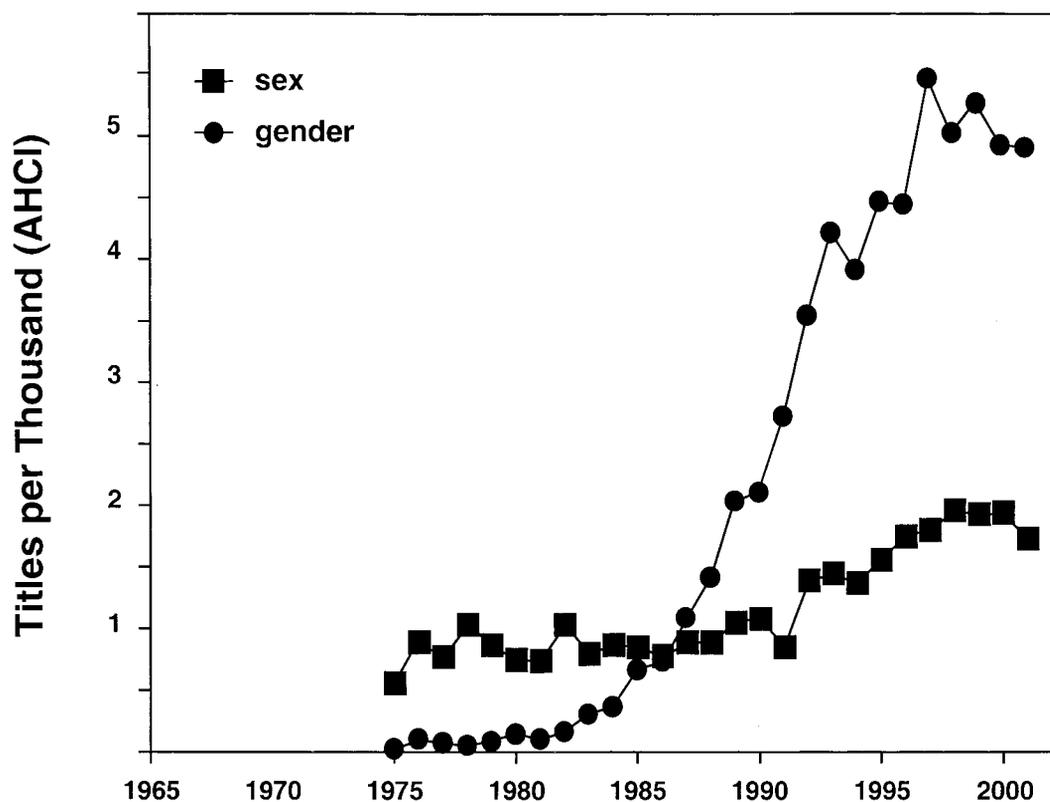


Fig. 4. Proportion of titles in the Arts & Humanities Citation Index containing the word *sex* and proportion containing the word *gender*.

majority of people who contributed to this new meaning of gender were hermaphrodites or intersexes. To them social science and social history overall owe a debt of gratitude. It is impossible to write about the political history of the second half of the twentieth century without reference to the concept of gender. This is particularly true with respect to the women's movement in politics" (p. xii). For similar reminiscences, and claims of priority, see Money (1973, 1985, 1995, p. 17ff.).

Excluding grammatical uses, most, if not all, of the gender-containing titles in SCI and SSCI from the 1960s appear to have derived the term from Money. These papers were mostly published in psychological journals and, at first, were concerned with individuals who did not conform to sexual stereotypes (hermaphrodites, transsexuals, transvestites, homosexuals, feminine boys, and masculine girls). However, in the late 1960s and early 1970s, gender began to appear in the titles of articles that addressed the behaviors and choices of individuals who conformed to gender stereotypes, with an emphasis on the extent to which the stereotypes were mutable or immutable, biological or social.

At this stage, it is worth discussing the causal connotations that had built up around *gender*. Money (1955) concluded that "Gender role and outlook as boy or man, girl or woman, was found to be in agreement with sex of rearing, except in three cases, and not to be automatically or instinctively determined by chromosomes, gonads or hormones." Similarly, Money et al. (1957) observed that "the sex of assignment and rearing is consistently and conspicuously a more reliable prognosticator of a hermaphrodite's gender role and orientation than is the chromosomal sex, the gonadal sex, the hormonal sex, the accessory internal reproductive morphology, or the ambiguous morphology of the external genitalia." They emphasized that "our findings indicate that neither a purely hereditary nor a purely environmental doctrine of the origins of gender role and orientation—of psychologic sex—is adequate."

Money and his co-workers offered two revealing analogies for the acquisition of a gender role: the first was the child's acquisition of a natural language (Money, 1955; Money et al., 1957); the second was the imprinting of a duckling on Konrad Lorenz when he imitated the

quacking of a mother duck (Money et al., 1957). In both these examples, the individual was seen as biologically primed to acquire a language or mother figure, but which language was acquired, or what individual was identified as mother, was determined by the environment. Consistent with these analogies, Money et al. (1957) believed that gender role was acquired very early in a child's development and once acquired was resistant to change: "Though the sex of rearing could transcend external genital morphology in psychologic importance, absence or correction of ambiguous genital appearance was psychologically beneficial. Reassignment of the sex of rearing after the early months of life was, without doubt, psychologically injurious." Although Money explicitly adopted an interactionist position as regards nature versus nurture, his work was implicitly read as lying at the nurture-end of the spectrum. Because a person's sex could differ from their gender role, gender became associated with a blurring of the male/female dichotomy, and the claim that upbringing trumped anatomy provided a powerful argument against the essential nature of sex differences.

An early person to employ the terminology of gender was the psychoanalyst Stoller (1964a, 1964b). For Stoller (1965), sex was biological but gender was social. The latter connoted "behavior learned from a tremendous pool of cues present in every culture and from a massive, intricate, though usually subtle, system of rewards and punishments in which every person lives from birth on" (p. 197). Although he did not deny some role for biology, Stoller (1968) wrote that "those aspects of sexuality that are called gender are primarily culturally determined" (p. xiii) and that "*gender* is a term that has psychological or cultural rather than biological connotations" (p. 9). Other psychoanalysts adopted a similar distinction between biological sex and social gender (e.g., Gershman, 1967; Ovesey & Person, 1973).

Stoller (1964b) and Greenson (1964) together introduced the term *gender identity* at the 23rd International Psycho-Analytical Congress in Stockholm (July–August 1963). The latter defined this to be "one's sense of being a member of a particular sex; it is expressed clinically in the awareness of being a man or male in distinction to being a woman or female." For Stoller (1968), "*gender identity* starts with the knowledge and awareness, whether conscious or unconscious, that one belongs to one sex and not the other . . . *gender role* is the overt behavior one displays in society, the role which he plays, especially with other people" (pp. 9–10). For Money and Ehrhardt (1972), "gender role is the public expression of gender identity, and gender identity is the private expression of gender role" (p. 4).

### Feminist Adoption

The origins of the use of gender among feminist scholars has been variously dated to the late 1960s (Nicholson, 1994) or the mid-1970s (Unger & Crawford, 1993). My own analysis suggests that its widespread adoption in feminist circles was delayed until the late 1970s or early 1980s. The first gender-containing title in the Web of Science that had an explicitly feminist context was *Some evolutionary aspects of human gender* (Tobach, 1971), in an issue of the *American Journal of Orthopsychiatry* devoted to *The Women's Movement: Social and Psychological Perspectives*. In this article, Tobach differentiated "biological sex" from "societally assigned gender" and warned against using "concepts from evolutionary biology to justify either retaining old traditions or changing them." Her article cited neither Money nor Stoller.

Other early feminist uses of gender occurred in books (not indexed in the Web of Science). Holter (1970) used sex and gender as interchangeable synonyms, seemingly for variety, whereas Millett (1970, p. 29) makes only passing reference to a sex/gender distinction, which she illustrates with a quote from Stoller (1968). Likewise, Bernard (1971, p. 16) derived her definitions of sex and gender directly from Stoller (1968). Oakley (1972) defined sex as biological and gender as psychological and cultural (pp. 16, 158). After a discussion of the work of Money and Stoller, she posed the rhetorical question "Does biology play any role at all in determining the development of gender identity in normal individuals?" and answered:

The consensus of opinion seems to be that its role is a minimal one, in that the biological predisposition to a male or female gender identity (if such a condition exists) may be decisively and ineradicably overridden by cultural learning. Those who have worked in the field of hermaphroditic disorders and problems of gender identity seem very impressed by the power of culture to ignore biology altogether. (p. 170).

Differences between successive editions of *Masculine/Feminine or Human?* (Chafetz, 1974, 1978) are particularly illuminating. In the first edition, Chafetz (1974) contrasted *innate* gender with *learned* sex roles. This edition contained no citations to Money. However, by the second edition (Chafetz, 1978), the terms had been swapped—*innate* sex was contrasted with *learned* gender roles—and references were added to Money and Ehrhardt (1972). Allowing for the time lags associated with publication, this suggests an absence of feminist consensus on the meaning of gender in the early 1970s with an emerging consensus by the late 1970s (see Gould & Kern-Daniels, 1977). This timing is supported by Unger (1979) who was able to write in the *American Psychologist*, "The term gender is

introduced for those characteristics and traits socioculturally considered appropriate to males and females” (my emphasis).

The only use of gender that I can find in *Women, Culture, and Society* (Rosaldo & Lamphere, 1974) is in the psychoanalytic chapter by Chodorow. Significantly, Ortner did not use gender in her influential chapter—*Is Female to Male as Nature is to Culture?* (an amended version of Ortner, 1972)—but 7 years later she was an editor of *Sexual Meanings: The Cultural Construction of Gender and Sexuality* (Ortner & Whitehead, 1981). In *Gender and Sex in Society*, Duberman (1975) defined sex as “an ascribed social status referring to the biological differences between people” whereas gender role referred to “the socially learned patterns of behavior that differentiate men from women in a given society” (p. 26). In *Toward an Anthropology of Women*, Rubin (1975) discussed the sex/gender system, which she defined as “the set of arrangements by which a society transforms biological sexuality into products of human activity, and in which these transformed sexual needs are satisfied” (p. 159).

Trends in feminist use of gender were assessed by scanning the contents of early issues of *Feminist Studies* (first issue in 1972) and *Signs: Journal of Women in Culture and Society* (first issue in 1975). The first gender-containing titles in *Feminist Studies* did not appear until Volume 5 (Davidoff, 1979) and Volume 6 (Vance, 1980). These authors derived their uses of gender from Oakley (1972) and Rubin (1975), respectively. Yudkin (1978) had earlier used gender in a philosophical discussion of transsexualism, but without the term appearing in the title. She constructed a trichotomy between biological *sex*, psychological *gender*, and social *sex role*. Her use of gender derived from Money and Stoller. The first issue of *Signs* defined the journal’s scope as including both sex and gender (Stimpson, Burstyn, Stanton, & Whisler, 1975), but use of gender was sparse in early issues (and predominantly by male authors). The first gender-containing title in *Signs* did not appear until the sixth volume (Baker, 1980), in a review of the biological literature on sex differences that contained numerous references to Money and coworkers. Gender-containing titles first exceeded sex-containing titles in Volume 11 of *Signs* (1986–1987).

Gender did not achieve uncontested acceptance by all feminists. In *Transsexual Empire*, Raymond (1979) treated *gender* as a technical or therapeutic term associated with the work of Money and Stoller. She found the term to have “certain problems for a feminist critic” as it gives “the impression that there is a fixed set of psychosocial conditions that determines gender identity and role.” Nevertheless, there were times that she found the word un-

avoidable despite her “dissatisfaction,” and in these places she “used it with reservation” (pp. 8–10).

From these small beginnings, use of gender became widely adopted by feminists during the 1980s. It is this adoption that I believe is principally responsible for the explosive growth in gender-containing titles that is observed in SSCI and AHCI during that decade (see Figs. 3 and 4). Feminists were able to embrace the concept of gender as their own contribution to discourse as the term’s earlier association with sexological science shifted into the background.

Feminist usage converged on a contrast between socially constructed gender and biologically determined sex. However, it proved difficult to maintain such a distinction. One problem with the simple dichotomization of biological sex and social gender was that no term remained to refer to situations in which causation was unknown, disputed, or involved an interaction between biology and culture. Thus, the choice of term for this middle ground became a simple matter of preference, blurring the conceptual distinction between terms. Moreover, among feminists, the domain of gender had a tendency to expand to subsume the category of sex, because the way that people talk about “male” and “female” bodies was also seen as socially constructed (discussed by Nicholson, 1994). Kessler and McKenna (1978) provided an uncompromising example of this position. They saw the element of social construction as primary in all aspects of maleness and femaleness: even to invoke two categories was a social construct. To emphasize their contention, they wrote of gender chromosomes and gender hormones. In a retrospective, McKenna and Kessler (2000) returned to this theme: “Retaining a separation between sex and gender, even if it is proposed that both are socially constructed, raises the question of why biology is so important that it merits a special category.”

Given the expansion in the domain of gender, and a certain indeterminacy in its meaning, it is hardly surprising that some authors who were unfamiliar with the subtleties of feminist debate interpreted gender as a simple synonym for sex and adopted it as such in their own writings. This is unambiguously demonstrated when gender is used in relation to the physiology of nonhuman animals, without any implication of a determining role of culture in the causation of observed differences. Such titles first appear in the 1970s (e.g., Hahn, Norton, & Fishman, 1977) and are now common in SCI.

The appearance of gender in a title from the natural sciences now communicates little if anything about causation or the ideology of the author. Among the reasons that working scientists have given me for choosing gender rather than sex in biological contexts are desires to

signal sympathy with feminist goals, to use a more academic term, or to avoid the connotation of copulation.

### Conclusion

This article addressed the history of terminology. During the first half of the twentieth century, gender appears to have been used predominantly in its grammatical sense, but its existing (albeit rare) use as a synonym of sex was readily available for anyone who wished to emphasize a dichotomy between different sources of sex-associated differences or to establish a separate domain for territory that had previously been considered part of the realm of sex. The expansion of the use of gender in the second half of the century appears to have derived from Money's concept of a gender role, introduced in the 1950s to refer to the self-identification of individuals whose genital sex was ambiguous. Significantly, in Money's usage, an individual's gender role could differ from various biological definitions of an individual's sex. From this beginning, there was a slow but gradual increase in the use of gender through the 1960s by writers, especially in the social sciences and among psychoanalysts, who wished to emphasize the environmental, social, or psychologic determinants of psychologic/behavioral differences between men and women. Some of these writers would have considered themselves feminists or at least sympathetic to the goals of the women's movement. Debates about nature versus nurture, the biological versus the social, and the autonomy of the social from the natural sciences, were of course much older than their association with a terminological sex vs. gender distinction.

Prior to the early 1980s, the rise in the use of gender in academic titles was not associated with an appreciable decline in the use of sex. The major increase in the use of gender, and the associated decline of sex, occurred in the 1980s after the adoption of *gender* as a technical term in feminist discourse. The available evidence strongly suggests that this usage was derived by descent with modification from Money. As the sex-to-gender ratio has declined, gender has come to be adopted as a simple synonym, perhaps a euphemism, for sex by many writers who are unfamiliar with the term's recent history.

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### REFERENCES

- Baker, S. W. (1980). Biological influences on human sex and gender. *Signs: Journal of Women in Culture and Society*, 6, 80–96.
- Bernard, J. (1971). *Women and the public interest*. Chicago: Aldine.
- Chafetz, J. S. (1974). *Masculine/feminine or human? An overview of the sociology of sex roles*. Itasca, IL: F. E. Peacock.
- Chafetz, J. S. (1978). *Masculine/feminine or human? An overview of the sociology of the gender roles* (2nd ed.). Itasca, IL: F. E. Peacock.
- Cottrell, L. S. (1942). The adjustment of the individual to his age and sex roles. *American Sociological Review*, 7, 617–620.
- Davidoff, L. (1979). Class and gender in Victorian England: The diaries of Arthur J. Munby and Hannah Culwick. *Feminist Studies*, 5(1), 87–141.
- Duberman, L. (1975). *Gender and sex in society*. New York: Praeger.
- Eliot, G. (1860). *The mill on the floss*. Edinburgh: Blackwood.
- Fishman, J. R., Wick, J. G., & Koenig, B. A. (1999). The use of "sex" and "gender" to define and characterize meaningful differences between men and women. In *Agenda for research on women's health for the 21st century*. (A report of the Task Force on the NIH Women's Health Research Agenda for the 21st century), (Vol. 2, pp. 15–19). Rockville, MD: Office of Research on Women's Health.
- Fletcher, D. E. (1991). The eternal battle of sex vs gender. *JAMA*, 266, 2833.
- Gershman, H. (1967). The evolution of gender identity. *Bulletin of the New York Academy of Medicine*, 43, 1000–1018.
- Goodhart, C. B. (1992). Sex and gender. *Nature*, 359, 182.
- Gould, M., & Kern-Daniels, R. (1977). Toward a sociological theory of gender and sex. *American Sociologist*, 12, 182–189.
- Greenson, R. R. (1964). On homosexuality and gender identity. *International Journal of Psychoanalysis*, 45, 217–219.
- Hahn, E. F., Norton, B. I., & Fishman, J. (1977). Influence of gender and castration on liver and CNS N-demethylation of morphine in rats. *Life Sciences*, 20, 95–99.
- Haig, D. (2000). Of sex and gender. *Nature Genetics*, 25, 373.
- Hartlap, S. (1979). Gender of infants conceived on different days of the menstrual cycle. *New England Journal of Medicine*, 300, 1445–1448.
- Holter, H. (1970). *Sex roles and social structure*. Oslo, Norway: Universitetsforlaget.
- Kessler, D. A. (1993). Guideline for the study and evaluation of gender differences in the clinical evaluation of drugs. *Federal Register*, 58(139), 39406–39416.
- Kessler, S. J., & McKenna, W. (1978). *Gender: An ethnomethodological approach*. New York: Wiley.
- Kim, J. S., & Nafziger, A. N. (2000). Is it sex or is it gender? *Clinical Pharmacology & Therapeutics*, 68, 1–3.
- Lewine, R. R. J. (1994). Sex: An imperfect marker of gender. *Schizophrenia Bulletin*, 20, 777–779.
- McKenna, W., & Kessler, S. J. (2000). Retrospective response. *Feminism and Psychology*, 10, 66–72.
- Mead, M. (1949). *Male and female: A study of the sexes in a changing world*. New York: William Morrow.
- Millett, K. (1970). *Sexual politics*. New York: Doubleday.
- Money, J. (1955). Hermaphroditism, gender and precocity in hyperadrenocorticism: Psychologic findings. *Bulletin of the Johns Hopkins Hospital*, 96, 253–264.
- Money, J. (1973). Gender role, gender identity, core gender identity: Usage and definition of terms. *Journal of the American Academy of Psychoanalysis*, 1, 397–402.
- Money, J. (1985). Gender history, theory and usage of the term in sexology and its relationship to nature nurture. *Journal of Sex and Marital Therapy*, 11, 71–79.
- Money, J. (1986). *Venuses/penuses*. Buffalo, NY: Prometheus.
- Money, J. (1995). *Gendermaps*. New York: Continuum.
- Money, J. (1996). Preface. In J. Money & A. A. Ehrhardt, *Man and woman, boy and girl. Gender identity from conception to maturity*. Northvale, NJ: Jason Aronson. (Original work published 1972)

- Money, J., & Ehrhardt, A. A. (1972). *Man and woman, boy and girl. The differentiation and dimorphism of gender identity from conception to maturity*. Baltimore: Johns Hopkins University Press.
- Money, J., Hampson, J. G., & Hampson, J. L. (1955a). Hermaphroditism: Recommendations concerning assignment of sex, change of sex, and psychologic management. *Bulletin of the Johns Hopkins Hospital*, 97, 284–300.
- Money, J., Hampson, J. G., & Hampson, J. L. (1955b). An examination of some basic sexual concepts: The evidence of human hermaphroditism. *Bulletin of the Johns Hopkins Hospital*, 97, 301–319.
- Money, J., Hampson, J. G., & Hampson, J. L. (1957). Imprinting and the establishment of gender role. *Archives of Neurology and Psychiatry*, 77, 333–336.
- Nicholson, L. (1994). Interpreting gender. *Signs: Journal of Women in Culture and Society*, 20, 79–105.
- Oakley, A. (1972). *Sex, gender, and society*. New York: Harper Colophon.
- Ortner, S. B. (1972). Is female to male as nature is to culture? *Feminist Studies*, 1(2), 5–31.
- Ortner, S. B., & Whitehead, H. (Eds.). (1981). *Sexual meanings: The cultural construction of gender and sexuality*. Cambridge, UK: Cambridge University Press.
- Ounsted, C., & Taylor, D. C. (Eds.). (1972). *Gender differences: Their ontogeny and significance*. Edinburgh: Churchill Livingstone.
- Ovesey, L., & Person, E. (1973). Gender identity and sexual psychopathology in men: A psychodynamic analysis of homosexuality, transsexualism, and tranvestitism. *Journal of the American Academy of Psychoanalysis*, 1, 53–72.
- Parsons, T. (1940). An analytical approach to the theory of social stratification. *American Journal of Sociology*, 45, 841–862.
- Parsons, T. (1942). Age and sex in the social structure of the United States. *American Sociological Review*, 7, 604–616.
- Parsons, T. (1949). *Essays in sociological theory pure and applied*. Glencoe, IL: Free Press.
- Pearson, G. A. (1996). Of sex and gender. *Science*, 274, 328–329.
- Raymond, J. G. (1979). *The transsexual empire: The making of the she-male*. Boston: Beacon.
- Rosaldo, M. Z., & Lamphere, L., (Eds.). (1974). *Women, culture, and society*. Stanford, CA: Stanford University Press.
- Rothman, K. J., & Liess, J. (1976). Gender of offspring after oral contraceptive use. *New England Journal of Medicine*, 295, 859–861.
- Rubin, G. (1975). The traffic in women: Notes on the “political economy” of sex. In R. R. Reiter (Ed.), *Toward an anthropology of women* (pp. 157–210). New York: Monthly Review Press.
- Smyth, D. H. (1968). Gender and sex. *British Medical Journal*, 2, 368.
- Stimpson, C. R., Burstyn, J. N., Stanton, D. C., & Whisler, S. M. (1975). Editorial. *Signs: Journal of Women in Culture and Society*, 1, v–viii.
- Stoller, R. J. (1964a). Gender-role change in intersexed patients. *JAMA*, 188, 684–685.
- Stoller, R. J. (1964b). A contribution to the study of gender identity. *International Journal of Psychoanalysis*, 45, 220–226.
- Stoller, R. J. (1965). Passing and the continuum of gender identity. In J. Marmor (Ed.), *Sexual inversion* (pp. 190–210). New York: Basic Books.
- Stoller, R. J. (1968). *Sex and gender: The development of masculinity and femininity*. London: Hogarth.
- Tobach, E. (1971). Some evolutionary aspects of human gender. *American Journal of Orthopsychiatry*, 41, 710–715.
- Unger, R. K. (1979). Toward a redefinition of sex and gender. *American Psychologist*, 34, 1085–1094.
- Unger, R. K., & Crawford, M. (1993). Sex and gender—The troubled relationship between terms and concepts. *Psychological Science*, 4, 122–124.
- Vance, C. S. (1980). Gender systems, ideology, and sex research: An anthropological analysis. *Feminist Studies*, 6, 129–143.
- Vincent, A. C. J. (1994). Seahorses exhibit conventional sex roles in mating competition, despite male pregnancy. *Behaviour*, 128, 135–151.
- Walker, P. L., & Cook, D. C. (1998). Gender and sex: Vive la difference. *American Journal of Physical Anthropology*, 106, 255–259.
- Wilson, D. R. (2000). Gender vs sex. *JAMA*, 284, 2997–2998.
- Yudkin, M. (1978). Transsexualism and women: A critical perspective. *Feminist Studies*, 4(3), 97–106.



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*AHR Forum*  
A History of “Gender”

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JOANNE MEYEROWITZ

SCHOLARLY ARTICLES TEND TO HAVE LIMITED SHELF LIVES, but twenty years on, Joan Scott’s “Gender: A Useful Category of Historical Analysis” has no discernible date of expiration. A cursory Google search leads to dozens of syllabi that feature it as required reading, and the figures from JSTOR attest to its durable popularity. Of all the *American Historical Review* articles on JSTOR, Scott’s has had by far the most traffic. Since JSTOR first began posting scholarly articles online in 1997, users have accessed “Gender” more than 38,000 times and printed more than 25,000 copies. For the past five years, it has consistently ranked in the top spot as the most frequently viewed and most frequently printed of JSTOR’s *AHR* articles.<sup>1</sup>

What elevates one article above the rest? What creates the reputation that makes an article required reading for more than twenty years? In part, it may be a matter of architecture. Scott built “Gender” with an artful use of argument. In one brief essay, she managed to summarize the advent of gender history, provide critiques of earlier theories of women’s subordination, introduce historians to deconstructionist methods, and lay out an agenda for future historical studies. But as we all know, academic reputation rests on more than compellingly structured argument, even when the argument is displayed well in a top-tier scholarly journal.<sup>2</sup> For historians, the surest way to explain a text is to place it in historical context. Thus, a short history of “Gender” the article might help us assess its rise to prominence and its influence within the field of U.S. history. And an even shorter history of “gender” the concept might suggest the article’s longer-lasting contribution to American social thought.

AS SCOTT NOTED, BY 1986, feminists had already adopted the term “gender” to refer to the social construction of sex differences, and theorists had already posed “gen-

For helpful comments on earlier drafts, many thanks to Margot Canaday, Regina Kunzel, Christina Simmons, and the editors of and anonymous reviewers for the *AHR*.

<sup>1</sup> Joan W. Scott, “Gender: A Useful Category of Historical Analysis,” *American Historical Review* 91, no. 5 (December 1986): 1053–1075. Thanks to Robert B. Townsend, Assistant Director for Research and Publications of the American Historical Association, for supplying these figures, which were compiled on December 27, 2007. The exact figures are 38,093 viewings and 25,180 printings. The closest competitors (based on total viewings plus total printings) were Robert Finlay, “The Refashioning of Martin Guerre,” *American Historical Review* 93, no. 3 (June 1988): 553–571, with 21,558 viewings and 11,183 printings, and Melvyn P. Leffler, “The Cold War: What Do ‘We Now Know?’” *American Historical Review* 104, no. 2 (April 1999): 501–571, with 22,075 viewings and 9,495 printings.

<sup>2</sup> For an attempt to theorize the sources of scholarly reputations, see, for example, Michèle Lamont, “How to Become a Dominant French Philosopher: The Case of Jacques Derrida,” *American Journal of Sociology* 93, no. 3 (1987): 584–622.

der" as an analytic category, akin to class and race. A few historians had begun to use the term "gender history" in addition to "women's history," and a handful had looked at men and masculinity as part of a gender history that did not focus solely on women. Scott intervened in this historiographic process at a critical moment. For some historians of women, the shift toward gender history was mostly unwelcome. To replace "women's history" with "gender history" and to include men and masculinity seemed to some at the time like a conservative retrenchment, a quest for respectability, or an abandonment of the study of marginalized and oppressed groups. Scott recognized the pitfalls and offered reassurance. She directly repudiated the use of "gender" as a de-politicized, social-scientized synonym for women or sex, and she promised to reinvigorate feminist history by expanding its realm of influence. In this way, she helped historians of women to approve (and other historians to discern) an emerging shift in historiography.

Scott outlined a problem faced by women's historians and proffered a solution. Two decades after the launching of the field, women's history was, she implied, stuck in a descriptive rut, relegated to the limited byways of social history inquiry. It had failed in its earlier claims to rewrite the master narrative of history, and it had not yet adequately explained the "persistent inequalities between women and men." Existing theories, Scott said, were ahistorical and reductionist. She offered a different approach for rethinking and rewriting history. Influenced by Derrida's deconstructionism and Foucault's formulation of dispersed power, she asked historians to analyze the language of gender, to observe how perceived sex differences had appeared historically as a natural and fundamental opposition. These perceived differences, she wrote, had often subordinated and constrained women, yes, but they had also provided a "primary way of signifying" other hierarchical relationships. This was the heart of her contribution: she invited us to look at how "the so-called natural relationship between male and female" structured, naturalized, and legitimated relationships of power, say, between ruler and ruled or between empire and colony. The history of gender could, it seems, inhabit more of the historical turf than could the history of women. It could even enter and remap the most resistant domains, such as the history of war, politics, and foreign relations.<sup>3</sup>

Although she promised to expand the realm of feminist influence, Scott could not deflect the critics from within her own fractious camp. Her embrace of poststructuralism and her consequent emphasis on the language of sex difference provoked a number of pointed rejoinders from prominent women's historians. Judith Bennett, for example, worried that "the Scottian study of gender ignore[d] women *qua* women," avoided reckoning with "material reality," and "intellectualize[d] and abstract[ed] the inequality of the sexes." Likewise, Linda Gordon suspected that a "focus on gender as difference in itself" as "a kind of paradigm for all other divides" had replaced "gender as a system of domination" and thereby substituted a pluralist vision of "multiple differences" for the study of "power differentials." Joan Hoff went further, even overboard. She accused poststructuralist gender historians, and Scott in particular, of nihilism, presentism, ahistoricism, obfuscation, elitism, obedience to patriarchy, ethnocentrism, irrelevance, and possibly racism. Poststructur-

<sup>3</sup> Scott, "Gender," 1066, 1067, 1073.

alism, she found, “erased woman as a category of analysis,” undermined the “traditional stage of historical fact-finding” for those groups of women whose history had not yet been written, and damaged political activism for women’s rights. She titled her essay “Gender as a Postmodern Category of Paralysis.”<sup>4</sup>

The critical commentary also came from historians who did not write women’s history, especially those who questioned the linguistic turn. Critiques of Scott’s work came from both the left and the right. Bryan Palmer, for example, decried her repudiation of historical materialism, and Gertrude Himmelfarb complained about the undermining of fact, reality, and objectivity.<sup>5</sup> In the United States, as others have suggested, “feminist historians” were “in the vanguard” of poststructuralist historical practice, especially in its manifestations outside of intellectual history, and Scott stood out at the front. In this sense, “Gender” came to represent something larger than itself. Scott served as the whipping girl not only for gender history but also for the challenges of poststructuralism, the revisionism of the latest new history, and the vogue—the “intellectual *haute couture*”—of imported French theory.<sup>6</sup> She may not have enjoyed the public flagellation, but it no doubt played a part in attracting readers to her essay.

DESPITE THE MISGIVINGS OF SOME HISTORIANS, gender soon took on a life of its own. Within the field of U.S. history, much of the new work on gender had little direct connection with Scott’s essay. Case studies of the intersections of race, class, and gender, for example, and accounts of how various groups of women and men participated differently in politics, labor, and consumption did not necessarily draw on Scott’s Derridean, Foucauldian model. Some new histories of gender in public cited Jürgen Habermas and Nancy Fraser more often than they cited Derrida and Scott.<sup>7</sup> But Scott’s article did have unquestionable influence, even among those authors who did not adopt the deconstructionist method wholesale. In the 1990s, it inspired a cohort of scholars who wrote gender history in a range of forms and fields. Within this cohort, a number of authors followed Scott’s proposal to foreground the dis-

<sup>4</sup> Judith M. Bennett, “Feminism and History,” *Gender and History* 1, no. 3 (1989): 258; Linda Gordon, review of Joan Wallach Scott, *Gender and the Politics of History*, *Signs* 15, no. 4 (1990): 858; Joan Hoff, “Gender as a Postmodern Category of Paralysis,” *Women’s History Review* 3, no. 2 (1994): 149, 162. For additional critical commentaries, see, for example, Sonya O. Rose et al., “Gender History/Women’s History: Is Feminist Scholarship Losing Its Critical Edge?” *Journal of Women’s History* 5, no. 1 (1990): 89–128. Some of these authors addressed Scott’s essays more generally, not just the article “Gender.”

<sup>5</sup> Bryan D. Palmer, *Descent into Discourse: The Reification of Language and the Writing of Social History* (Philadelphia, 1990), esp. chap. 5; Gertrude Himmelfarb, “Some Reflections on the New History,” *American Historical Review* 94, no. 3 (June 1989): 661–670.

<sup>6</sup> Joyce Appleby, Lynn Hunt, and Margaret Jacob, *Telling the Truth about History* (New York, 1994), 226; Sandra M. Gilbert and Susan Gubar, “Sexual Linguistics: Gender, Language, Sexuality,” *New Literary History* 16, no. 3 (1985): 521. On the “linguistic turn” in history, see, for example, John E. Toews, “Intellectual History after the Linguistic Turn: The Autonomy of Meaning and the Irreducibility of Experience,” *American Historical Review* 92, no. 4 (October 1987): 879–907; Kathleen Canning, “Feminist History after the Linguistic Turn: Historicizing Discourse and Experience,” *Signs* 19, no. 2 (1994): 368–404.

<sup>7</sup> See, for example, Mary Ryan, *Women in Public: Between Banners and Ballots, 1825–1880* (Baltimore, 1990), and Glenda Elizabeth Gilmore, *Gender and Jim Crow: Women and the Politics of White Supremacy in North Carolina, 1896–1920* (Chapel Hill, N.C., 1996).

cursive use of perceived sex differences and track how they constituted relationships of power. In U.S. history, the case studies of "women's worlds" and "female cultures" that had proliferated in the 1980s dwindled as accounts rose of the ways in which the language of gender had shored up hierarchies of race, class, region, politics, nation, and empire.

A quick (and, forgive me, incomplete) survey of just a few subfields of U.S. history establishes the point. In southern history, Jacquelyn Dowd Hall endorsed the gender project early on. "The South," she wrote in 1989, "provides a prime example of how gender signifies relations of power in hierarchical regimes." Other historians took up the task. Stephanie McCurry found that proslavery ministers and politicians repeatedly drew analogies between "the subordination of women" and "that of slaves," and thereby "endow[ed] slavery with the legitimacy of the family and especially marriage." They used the language of gender "to naturalize other social relations—class and race, for example." Laura Edwards reported similar analogies—between women and other "dependent" groups—in the Reconstruction-era writings of elite white southern men, who used the language of gender to legitimate their bid to monopolize political power. Historians also noted how the southern states themselves were coded as feminine within the United States. Nina Silber, for example, pointed to a post-bellum northern language of gender that portrayed the South as a "submissive" wife and helped to enable the "romance" of sectional reunion.<sup>8</sup>

In other areas, historians also attended to the ways that political theorists, government officials, and other writers used the language of sex difference to construct and sustain political and social hierarchies. In early American history, Mary Beth Norton described how seventeenth-century British male colonists established governments based on a gendered, hierarchical model of the family, and Kathleen Brown suggested that gender discourse shaped the emerging political order in Virginia from the first conflicts with the Indians through the course of Bacon's Rebellion. Jennifer Morgan illustrated how early European narratives of the New World "relied on gender," especially on accounts of monstrous Indian and African women, "to convey an emergent notion of racialized difference," and Toby Ditz delineated how eighteenth-century Philadelphia merchants stabilized their own fragile masculine status by feminizing and thereby stigmatizing their failed and dishonest colleagues as "weeping victims and harpies."<sup>9</sup> At the other end of the chronological span, historians of twentieth-century U.S. politics examined how male politicians

<sup>8</sup> Jacquelyn Dowd Hall, "Partial Truths," *Signs* 14, no. 4 (1989): 910; Stephanie McCurry, *Masters of Small Worlds: Yeoman Households, Gender Relations, and the Political Culture of the Antebellum South Carolina Low Country* (New York, 1995), 214, 224; Laura Edwards, *Gendered Strife and Confusion: The Political Culture of Reconstruction* (Urbana, Ill., 1997), esp. chap. 6; Nina Silber, *The Romance of Reunion: Northerners and the South, 1865–1900* (Chapel Hill, N.C., 1993), 10.

<sup>9</sup> Mary Beth Norton, *Founding Mothers and Fathers: Gendered Power and the Forming of American Society* (New York, 1996); Kathleen M. Brown, *Good Wives, Nasty Wenches, and Anxious Patriarchs: Gender, Race, and Power in Colonial Virginia* (Chapel Hill, N.C., 1996); Jennifer L. Morgan, "'Some Could Suckle over Their Shoulder': Male Travelers, Female Bodies, and the Gendering of Racial Ideology, 1500–1770," *William and Mary Quarterly* 54, no. 1 (1997): 168; Toby L. Ditz, "Shipwrecked; or, Masculinity Imperiled: Mercantile Representations of Failure and the Gendered Self in Eighteenth-Century Philadelphia," *Journal of American History* 81, no. 1 (1994): 54. On gender more generally in early American history, see Toby L. Ditz, "The New Men's History and the Peculiar Absence of Gendered Power: Some Remedies from Early American Gender History," *Gender and History* 16, no. 1 (2004): 1–35.

used the language of gender to create a hierarchy in which they stood above their male opponents. In the early twentieth century, they cast male reformers as feminine and therefore lacking, and in the late twentieth century, they attacked male liberals in somewhat similar form. Gail Bederman and Arnaldo Testi showed how Theodore Roosevelt shook off the gendered smear by combining his reform agenda with an imperialist, racist hypermasculinity, and Robert Dean and K. A. Cuordileone elucidated how John F. Kennedy attempted to repel the aspersion with an aggressive expression of liberalism.<sup>10</sup>

Perhaps most surprising, gender history also made significant forays into the history of foreign policy, the field of U.S. history that had seemed most immune to the women's history enterprise. Scott had specifically called for such an intervention; in 1990, Emily Rosenberg responded and made the case for the potential benefits of gender analysis. Gendered imagery, she said, pervaded accounts of international affairs, legitimating foreign relations of domination and dependence. Andrew Rotter pursued the lead and showed how mid-twentieth-century U.S. policymakers had imagined India as feminine and India's male leaders as passive, emotional, and lacking in virility. In this case, the "feminization" undermined the opportunity for alliance between the U.S. and India. In other cases, though, the "masculinization" of nations and their leaders damaged international relations, while "feminization" eased them. Frank Costigliola, for example, investigated the writings of Cold War architect George Kennan, who shifted from feminizing a beloved Russia in the 1930s to portraying Soviet leaders as "monstrously masculine" and rapacious in the post-World War II years. Petra Goedde traced the inverse shift with regard to Germany. During World War II, American soldiers vilified the Nazi leaders, whom they understood as brutally masculine, but after the war they "developed a feminized image" of Germans as a population in need of protection, and thus, Goedde claimed, "paved the way toward reconciliation."<sup>11</sup>

Historians also began to suggest that discourses of gender had promoted and sustained American military interventions. In *Fighting for American Manhood*, Kristin Hoganson explored "how gender politics provoked the Spanish-American and Philippine-American wars," as the subtitle of her book stated plainly. As they advocated war, jingoes and imperialists expressed heightened concern with masculinity and looked to the military to build and prove American manhood. They posed the

<sup>10</sup> Gail Bederman, *Manliness and Civilization: A Cultural History of Gender and Race in the United States, 1880-1917* (Chicago, 1995), chap. 5; Arnaldo Testi, "The Gender of Reform Politics: Theodore Roosevelt and the Culture of Masculinity," *Journal of American History* 81, no. 4 (1995): 1509-1533; Robert D. Dean, *Imperial Brotherhood: Gender and the Making of Cold War Foreign Policy* (Amherst, Mass., 2001); K. A. Cuordileone, *Manhood and American Political Culture in the Cold War* (New York, 2005).

<sup>11</sup> Emily S. Rosenberg, "Gender," *Journal of American History* 77, no. 1 (1990): 116-124; Andrew J. Rotter, "Gender Relations, Foreign Relations: The United States and South Asia, 1947-1964," *Journal of American History* 81, no. 2 (1994): 518-542; Frank Costigliola, "'Unceasing Pressure for Penetration': Gender, Pathology, and Emotion in George Kennan's Formation of the Cold War," *Journal of American History* 83, no. 2 (1997): 1333; Petra Goedde, "From Villains to Victims: Fraternalization and the Feminization of Germany, 1945-1947," *Diplomatic History* 23, no. 1 (1999): 2, 20. See also the essays on gender in the Winter 1994 issue of *Diplomatic History*, especially Geoffrey S. Smith, "Commentary: Security, Gender, and the Historical Process," *Diplomatic History* 18, no. 1 (1994): 79-90; Petra Goedde, *GIs and Germans: Culture, Gender, and Foreign Relations, 1945-1949* (New Haven, Conn., 2003). For a useful review essay, see Kristin Hoganson, "What's Gender Got to Do with It? Women and Foreign Relations History," *OAH Magazine of History* 19, no. 2 (2005): 14-18.

Spanish soon-to-be enemies as both distastefully feminine and repulsively masculine—"effeminate aristocrats" and "savage rapists"—and sometimes also feminized the Cubans and Filipinos as well as their own domestic opponents. Mary Renda outlined a somewhat different masculine discourse of "interventionist paternalism" that underwrote the American occupation of Haiti. The gendered language of fatherhood helped U.S. policymakers and marines to justify imperialist violence as a manly attempt to protect, educate, and discipline the allegedly childlike Haitians. And Robert Dean wrote of the threats to the "imperial masculinity" of the mid-twentieth-century U.S. foreign policy elite. Politicians and policymakers used the language of gender to defend their own manhood and diminish that of their rivals, and thereby engaged, Dean suggested, in a "politics of manhood" that "crucially shaped the tragedy of the Vietnam War." Hoganson, Renda, and Dean (and the other authors mentioned above) did not confine their analyses to the deconstruction of binary oppositions, but they provided evidence of how the language of gender constructed and legitimated American imperialism and its violent manifestations.<sup>12</sup>

Taken together, these various works point, as Scott predicted, to the multiplicity of meanings that gendered language conveyed. In different historical contexts, masculinity represented strength, protection, independence, camaraderie, discipline, rivalry, militarism, aggression, savagery, and brutality, and femininity represented weakness, fragility, helplessness, emotionality, passivity, domestication, nurturance, attractiveness, partnership, excess, and temptation. The so-called natural differences between the sexes had no fixed and unchangeable meaning, and in their variety they provided potential meaning for a range of other relationships. As other historians have protested, though, the ultimate impact of the language of gender remained hard to discern.<sup>13</sup> When (and how), as Scott asked, did the language of gender crucially structure experience and actually influence behavior and decision-making, and when did it simply add a convenient rhetorical flourish or embellish with a hollow cliché? When (and how), as Scott asked, did the language of gender constitute other relations of power, and when was it just a minor paragraph or a supplemental example within the narratives of social and political order? Even without all the answers, the growing number of studies of gender discourse pushed historians to recognize its pervasiveness, the diverse domains in which perceived sex differences appeared as model, analogy, and metaphor for hierarchical relationships, and the wide-ranging and changing meanings of masculinity and femininity in the modern era.

The studies also enhanced the reputation of Scott's essay and injected its message into traditional subfields of historical study. Almost all of the works cited above (and many other books and articles as well) mentioned "Gender," in the footnotes if not in the text. Some of them quoted it directly. It became a validating authority behind the monographic works that moved gender to the center of specialized subfields in which it had earlier stood at the margins.<sup>14</sup> By the end of the 1990s, through a process

<sup>12</sup> Kristin L. Hoganson, *Fighting for American Manhood: How Gender Politics Provoked the Spanish-American and Philippine-American Wars* (New Haven, Conn., 1998), 11; Mary A. Renda, *Taking Haiti: Military Occupation and the Culture of U.S. Imperialism, 1915–1940* (Chapel Hill, N.C., 2001); Dean, *Imperial Brotherhood*, 243.

<sup>13</sup> See, for example, Melvyn P. Leffler, "New Approaches, Old Interpretations, and Prospective Reconfigurations," *Diplomatic History* 19, no. 2 (1995): 195.

<sup>14</sup> Scott's article also had a significant impact on U.S. labor history. See especially Ava Baron, ed.,

of repetition, “Gender” had reshaped the commonplace wisdom of the discipline. As a measure of its success, Scott’s essay increasingly served as a voice from the recent past stating eloquently what everybody, it seems, already knew.

Meanwhile, Scott herself moved in new directions. In 1999, she questioned the ongoing vitality of the term “gender.” In the 1980s, she wrote, gender had “seemed a useful category of analysis precisely because it had an unfamiliar, destabilizing effect.” Now, however, it had “lost its ability to startle and provoke.” In everyday usage, gender had become “a synonym for women, for the differences between the sexes, for sex.” The word “gender” had crept into women’s history without necessarily transforming the field. It appeared often in “predictable studies of women, or . . . of differences in the status, experience, and possibilities open to women and men.” Many accounts failed to “examine how the meanings of ‘women’ and ‘men’” were “discursively established” or to address the “variations of subjectively experienced ‘womanhood.’” They thereby imposed a false solidity on the unstable and variable categories of “women” and “men.” Scott now avoided the word “gender” and wrote instead about “differences between the sexes and about sex as a historically variable concept.” She turned more concertedly to psychoanalysis, to the fantasies that enable identities, including the “phantasmatic projections that mobilize individual desires into collective identifications.” In her 2005 book, *Parité! Sexual Equality and the Crisis of French Universalism*, and her 2007 book, *The Politics of the Veil*, she entered into current debates in French politics. She focused less on the language of sex difference and more on the language of universalism in contemporary France. In these books, she did not renounce the study of “gender,” but she positioned French gender relations within a discursive analysis of “the abstract individualism” that animates French republican traditions.<sup>15</sup>

As one would expect, other historians also ventured into new territory. In U.S. women’s—and now gender—history, they brought in race, sexuality, and nationality as equally useful categories of historical analysis, and they borrowed from postcolonial, critical race, queer, and political theory. Other forms of perceived difference seem to have constituted gender as much as gender constituted them. In particular, the call to address race had at least as much impact on U.S. women’s history as the call to attend to gender. Historians of women and gender also turned to the policy history of welfare and wages, the legal history of marriage, and the social history of those who questioned and transgressed gender norms. Historians of women shifted away from the local community studies that had characterized social history and focused more on individual or collective biography, questions of law and citizenship, and transnational circulations of women and ideas about womanhood. They rewrote the history of women’s movements with a closer eye to differences among women and conflicts among competing schools of feminists. At the same time, historians of manhood produced a series of studies of shifting conceptions, multiple variants, and

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*Work Engendered: Toward a New History of American Labor* (Ithaca, N.Y., 1991). The article also had influence outside U.S. history, of course, but I will leave that to the other participants in this forum.

<sup>15</sup> Joan Wallach Scott, *Gender and the Politics of History*, rev. ed. (New York, 1999), xi–xii, 204; Scott, *Parité! Sexual Equality and the Crisis of French Universalism* (Chicago, 2005); Scott, *The Politics of the Veil* (Princeton, N.J., 2007), 154. See also Scott, “Fantasy Echo: History and the Construction of Identity,” *Critical Inquiry* 27, no. 2 (2001): 284–304.

repeated crises of masculinity. Gender history, then, continued (and continues) to thrive in several incarnations, and despite the fears of early (and later) critics, it coexists and overlaps with, instead of supplanting or displacing, the history of women.<sup>16</sup> Amid the profusion, Scott's article has taken on the emblematic role of a foundational text.

SCOTT'S ESSAY HAD ITS MOST OBVIOUS INFLUENCE in the fields of women's and gender history, but it also played a significant part in the broader shift from social to cultural history, from the study of the demography, experiences, and social movements of oppressed and stigmatized groups to the study of representations, language, perception, and discourse. In U.S. history, the rise of gender history was similar to and roughly simultaneous with changes in other identity-based fields of history, including African American, Latino/a, Asian American, immigrant, gay and lesbian, and working-class history. Gender history and the historical construction of masculinity had their counterparts in the history of race and the construction of whiteness, the history of ethnicity and the construction of national identity, the history of sexuality and the construction of heterosexuality, and the history of class and the construction of middle-classness. To a certain extent, the same left-leaning political energies that had informed much of the new social history informed the new cultural history as well. The irony is that social history, the alleged source of centrifugal fragmentation, had spun out into a cultural history that seems to have gravitated back—in the histories of masculinity, whiteness, national identity, heterosexuality, and middle-classness—to return, with a new and critical torque, to the pre-social-history center of historical inquiry.<sup>17</sup> "Gender," and Scott's other writings as well, provided a key piece of the theoretical grounding for this historiographic trend.

Like all historiographic moments, this one, too, will no doubt pass. And when it does, what will we remember? We might consider another context for understanding the significance of Scott's essay and its larger contribution beyond historiography. We have only begun to historicize "gender"—that is, to write the history of the concept of gender itself. Scott's essay belongs in that history; it represents a turning point when U.S. feminist scholars pulled "gender" away from its scientific and social scientific origins, reworked its meaning, and suggested its broader social, cultural, and historical impact.

Scott dated the term "gender," in its contemporary usage, to the 1970s feminist movement, but the word has a longer history, even as a reference to the non-biological components of sex. Before the 1950s, linguists used "gender," as Scott acknowledged, to refer to a form of grammatical classification. The concept of socially constructed sex differences did not yet have a word to connote it. Nonetheless, theories of the social construction of sex differences emerged in tandem with theories of the social construction of other forms of group difference. From the early twen-

<sup>16</sup> For more recent concerns that gender history will supplant women's history, see Alice Kessler-Harris, "Do We Still Need Women's History?" *Chronicle of Higher Education* 54, no. 15 (December 7, 2007): B6.

<sup>17</sup> For a recent account of this trend, see Daniel Wickberg, "Heterosexual White Male: Some Recent Inversions in American Cultural History," *Journal of American History* 92, no. 1 (2005): 136–157.

tieth century on, social scientists engaged in a profound questioning of biological determinism and the categories on which it relied, not only with regard to sex but also with regard to race, ethnicity, national character, sexuality, criminality, and mental illness. By the mid-twentieth century, anthropologists and sociologists wrote of “sex roles” to refer to the culturally determined expected behavior of women and men and “sexual status” to acknowledge that different cultures accorded different social rankings to women and men. Psychologists used the phrases “psychological sex” and “sex-role identification” to point to a person’s acquired sense of self as female or male.<sup>18</sup>

In the mid- to late 1950s, John Money, Joan Hampson, and John Hampson, all then at Johns Hopkins University, introduced the term “gender” into this scientific literature. In a series of articles on intersexuality, they argued for the environmental determinants of “gender,” “gender role,” and “gender role and orientation,” just as others had earlier argued for the environmental determinants of “sex roles” and “psychological sex.” Children learned “gender” in early childhood, they argued, in the same way they learned a language. Biological sex, however it was defined, did not determine one’s “gender role and orientation.”<sup>19</sup> Other scientists and social scientists picked up the new terminology. In 1962, psychoanalyst Robert Stoller and his colleagues at the University of California in Los Angeles opened the first Gender Identity Research Clinic (GIRC), and in 1968, Stoller published the book *Sex and Gender*, which seems to have been the first American book with the word “gender,” in its current non-linguistic form, in the title. For Stoller, gender referred to the particular balance of masculinity and femininity found in each person. It had “psychological or cultural rather than biological connotations.” Stoller was not a feminist. In fact, he worried about the erosion of gender roles and the developmental disturbance of “gender identity,” the new term he coined for “psychological sex.” He and his colleagues at the GIRC worked to instill masculinity in feminine boys and femininity in masculine girls. If gender was mostly socially constructed, then someone, they reasoned, had to repair it when it was improperly built. Stoller and his colleagues signed up for the job.<sup>20</sup>

Influenced by the women’s movement, American feminists appropriated the word “gender” in the 1970s and transformed its meaning. Like others before them, feminist social scientists used “gender” to reject the notion that the perceived sex

<sup>18</sup> On American social scientists and the social construction of sex differences, see, for example, Rosalind Rosenberg, *Beyond Separate Spheres: Intellectual Roots of Modern Feminism* (New Haven, Conn., 1982); Carl Degler, *In Search of Human Nature: The Decline and Revival of Darwinism in American Social Thought* (New York, 1991); Mari Jo Buhle, *Feminism and Its Discontents: A Century of Struggle with Psychoanalysis* (Cambridge, Mass., 1998).

<sup>19</sup> For uses of the new terms, see John Money, “Hermaphroditism, Gender, and Precocity in Hyperadrenocorticism: Psychologic Findings,” *Bulletin of the Johns Hopkins Hospital* 96 (1955): 253–264; John Money, Joan G. Hampson, and John L. Hampson, “Imprinting and the Establishment of Gender Role,” *American Medical Association Archives of Neurology and Psychiatry* 77 (1957): 333–336. Money later retreated from his early environmentalism; by the end of the 1960s, he speculated that early exposure to sex hormones and the neurophysiology of the brain (as well as environment) shaped gender identity. On Money, the Hampsons, and “gender,” see Bernice Hausman, *Changing Sex: Transsexualism, Technology, and the Idea of Gender* (Durham, N.C., 1995), chap. 3; Joanne Meyerowitz, *How Sex Changed: A History of Transsexuality in the United States* (Cambridge, Mass., 2002), chap. 3.

<sup>20</sup> Robert J. Stoller, *Sex and Gender: On the Development of Masculinity and Femininity* (New York, 1968), 9. On Stoller and the GIRC, see Meyerowitz, *How Sex Changed*, chap. 3; Phyllis Burke, *Gender Shock: Exploding the Myths of Male and Female* (New York, 1996).

differences in behavior, temperament, and intellect were simply natural or innate, but unlike their predecessors, they rejected functionalism and questioned whether gender and gender roles were necessary or good. If gender was artifice, then many 1970s feminists saw little reason to maintain it, especially when it played a part in subordinating women. But gender, in its multiple variations, was not so easily willed away. It was built into the structure and practice of families, education, labor markets, and government policies, and it had deep roots in the everyday behaviors and fantasies of individual women and men. Some academic feminists, especially in the humanities, turned away from the study of gender roles, gender systems, and gender segregation, and focused instead on the reconstruction and revaluation of femininities, women's writings, women's ethics, and women's worlds.<sup>21</sup>

Others searched for theoretical approaches that could explicate how perceptions of sex difference operated in language, psyche, and symbolic order. In the late 1970s and early 1980s, some American feminist literary critics turned to French poststructuralist theory. They drew on the works of Jacques Lacan, Roland Barthes, and Jacques Derrida, and they translated the writings of Hélène Cixous, Luce Irigaray, and Julia Kristeva. They expanded their purview from "the woman reader, women's culture, and the woman's text" to "the whole of literature and culture." Cixous wrote: "Every theory of culture, every theory of society, the whole conglomeration of symbolic systems . . . it is all ordered around hierarchical oppositions that come back to the man/woman opposition." By the early 1980s, male literary critics recognized the feminist affinity to poststructuralism. In 1983, in *Literary Theory*, Terry Eagleton suggested that "the movement from structuralism to post-structuralism was in part a response" to the demands of the women's movement. In this rendition, feminism stood front and center on the poststructuralist stage.<sup>22</sup>

In 1986, with the article "Gender," Joan Scott helped to bridge the gap between the feminist social scientists who critiqued "gender" and "gender roles" and the feminist literary critics who deconstructed textual representations of sex difference.<sup>23</sup> She wrote in a moment, as she noted, "of great epistemological turmoil," when social scientists were shifting "from scientific to literary paradigms," and when feminists were finding "scholarly and political allies" among poststructuralists. For Scott, gender was "a constitutive element of social relationships based on perceived differences between the sexes," and also "a primary way of signifying relationships of power." Scott's dual definition allowed her to bring together the social scientists who rejected biological determinism and questioned the allegedly natural differ-

<sup>21</sup> On 1970s feminists and "gender," see, for example, Suzanne J. Kessler and Wendy McKenna, *Gender: An Ethnomethodological Approach* (Chicago, 1978); see also Rosalind Rosenberg, "Gender," in Theodore M. Porter and Dorothy Ross, eds., *The Modern Social Sciences* (Cambridge, 2003), 678–692.

<sup>22</sup> Elaine Showalter, "Women's Time, Women's Space: Writing the History of Feminist Criticism," *Tulsa Studies in Women's Literature* 3, no. 1/2 (1984): 35; Hélène Cixous, "Castration or Decapitation?" *Signs* 7, no. 1 (1981): 44; Terry Eagleton, *Literary Theory: An Introduction* (Minneapolis, 1983), 149. For American feminist adaptations of French theory, see, for example, Elaine Marks and Isabelle de Courtivron, eds., *New French Feminisms: An Anthology* (Amherst, Mass., 1980); *Writing and Sexual Difference*, Special Issue, *Critical Inquiry* 8, no. 2 (1981); *Feminist Readings: French Texts/American Contexts*, Special Issue, *Yale French Studies* 62 (1981). For critical commentaries by historians, see Buhle, *Feminism and Its Discontents*, chap. 9; Claire Goldberg Moses, "Made in America: 'French Feminism' in Academia," *Feminist Studies* 24, no. 2 (1998): 241–274.

<sup>23</sup> Scott was soon joined in this endeavor by Judith Butler; see Butler, *Gender Trouble: Feminism and the Subversion of Identity* (New York, 1990).

ences on which it was based and the philosophers, psychoanalysts, and literary critics who suggested that the language of difference sustained Western social and political order. She was not alone in this kind of endeavor. A year earlier, for example, Henry Louis Gates, Jr. (and others) had posited race as a “trope of ultimate, irreducible difference” that naturalized distinctions between “cultures, linguistic groups, or adherents of specific belief systems.”<sup>24</sup> Within the United States, the scholarly study of difference and inequality, once firmly grounded in social science, had migrated to the humanities and taken root in the study of language. It soon spread beyond the analysis of literature and into the reading of multifarious texts, including the kinds of texts that historians typically use as evidence.

This abbreviated genealogy of gender might help to place Scott’s contribution in a broader context. For historians, Scott summarized explanations of gender inequality, captured an emerging historiographic trend, and imported theory to a discipline of committed empiricists. She promised both to expand the terrain of the new social and cultural history and to return to and revivify the traditional fields of historical study. In the 1980s and 1990s, her readers sustained her argument first by publicly debating its merits and then by applying its theory and its method of reading. Beyond the historical discipline, though, Scott’s essay entered into decades-long conversations on the social and symbolic constructions of sex difference. She helped to move the American concept of gender beyond its scientific and social scientific origins and nudged the American adaptations of poststructuralism beyond their recognized place in literary criticism. She suggested how the language of sex difference had historically provided a means to articulate relationships of power. In this way, she tied gender back to other forms of difference and pushed us to ponder the metanarratives that mutually constituted various social and political hierarchies. And ponder we should. This may, in the end, prove to be the enduring legacy of “Gender.”

<sup>24</sup> Scott, “Gender,” 1066, 1067; Henry Louis Gates, Jr., “Writing ‘Race’ and the Difference It Makes,” “Race,” *Writing, and Difference*, Special Issue, *Critical Inquiry* 12, no. 1 (1985): 5. Gates’s essay is the editor’s introduction to the issue; some of the other essays in the issue also address the language of race difference. See also Evelyn Brooks Higginbotham, “African American Women’s History and the Meta-language of Race,” *Signs* 17, no. 2 (1992): 251–274.

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## Essay

# Sex Determination: Why So Many Ways of Doing It?

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**Abstract:** Sexual reproduction is an ancient feature of life on earth, and the familiar X and Y chromosomes in humans and other model species have led to the impression that sex determination mechanisms are old and conserved. In fact, males and females are determined by diverse mechanisms that evolve rapidly in many taxa. Yet this diversity in primary sex-determining signals is coupled with conserved molecular pathways that trigger male or female development. Conflicting selection on different parts of the genome and on the two sexes may drive many of these transitions, but few systems with rapid turnover of sex determination mechanisms have been rigorously studied. Here we survey our current understanding of how and why sex determination evolves in animals and plants and identify important gaps in our knowledge that present exciting research opportunities to characterize the evolutionary forces and molecular pathways underlying the evolution of sex determination.

variance that is otherwise hidden [2]. While many unicellular organisms produce gametes of equal size (isogamy, see Box 1), sexual reproduction in most multicellular organisms has led to the evolution of female and male gametes differing in size (anisogamy), and often to the evolution of two separate sexes. Even though the outcome of sex determination—whether an individual produces relatively few large ova or many small sperm—is strongly conserved, a bewildering number of underlying mechanisms can trigger development as either a male or female [3,4].

In humans, sex is determined by sex chromosomes (XX females, XY males). The X and Y chromosomes harbor dramatically different numbers and sets of genes (about 1,000 genes on the X and only a few dozen genes on the Y), yet they originated from ordinary autosomes during the early evolution of mammals (Figure 1). Restriction of recombination followed by gene loss on the Y has resulted in the morphological differentiation of sex chromosomes (for a review of the molecular and evolutionary processes involved in Y degeneration, see [4,5]). The vast majority of genes on the sex chromosomes are not directly involved in sex determination, and development as a male

or female depends on the presence of a single master sex-determining locus, the *Sry* gene, on the male-limited Y chromosome. Expression of *Sry* early in embryonic development initiates testis differentiation by activating male-specific developmental networks, while in its absence, ovaries develop. The first visible signs of sexual differentiation of the ovary and testis occur by the sixth week of gestation in humans [6], and sex hormones initiate further sexual differentiation in nongonadal tissues and organs [7]. When this developmental process goes awry, the effects can be catastrophic, causing everything from ambiguous external genitalia (which occurs in up to one in 4,500 infants) to sterility (which is more cryptic and difficult to diagnose but may be far more common).

Like humans and most mammals, other genetic model systems, such as *Drosophila melanogaster* flies and *Caenorhabditis elegans* nematodes, harbor sex chromosomes, and their commonalities have led to general assumptions about the conservation of sex determination mechanisms. However, these model organisms present a false impression of stability in how sex is determined, and their commonalities mask the diversity and turnover in sex determination mechanisms that is readily

## Introduction

Sex—the mixing of genomes via meiosis and fusion of gametes—is nearly universal to eukaryotic life and encompasses a diverse array of systems and mechanisms [1]. One major role of sex is to bring together alleles carried by different individuals, revealing beneficial genetic

Essays articulate a specific perspective on a topic of broad interest to scientists.

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apparent when taking a broader taxonomic view. In this article, we address three common myths about sex determination and then deconstruct them based on a broad taxonomic survey of animals and plants.

## Myths of Sex Determination

### Myth 1: Sex is typically determined by X and Y chromosomes

Many biologists are habituated to thinking about sex determination through the familiar examples of mammals and *D. melanogaster*, and assume that sex determination by sex chromosomes is the norm, that males are XY and females are XX, and that sex chromosomes are a stable component of the genome. While biologists are generally aware of other modes of sex determination (such as female heterogamety in birds, temperature-dependent sex determination in reptiles, or development of males from unfertilized eggs in bees), these alternatives are often viewed as strange and aberrant [8].

### Myth 2: Sex is controlled by one master-switch gene

Sex determination in model species suggests that a master-switch gene (e.g. *Sry* in mammals, *Sxl* in *D. melanogaster*, and *xol-1* in *C. elegans*) acts as the main control element to trigger either male or female sexual development. Changes in the sex determination pathways across taxa are assumed to involve adding a new master-switch gene to this molecular pathway (as in some fly taxa; [9]), with little change to downstream elements of the sex determination pathway [10]. A few genes are thought to have the capacity to take on the role of sex determination genes, and these have been co-opted as master-switch genes independently in different lineages (for example, *dmrt1* in several vertebrates [11–14] and *tra* in insects [15–17]).

### Myth 3: Sex chromosome differentiation and degeneration is inevitable

Sex chromosomes originate from identical autosomes by acquiring a sex determination gene (for example, the origin of the *Sry* gene in mammals approximately 180 million years ago or *Sxl* in the *Drosophila* genus >60 million years ago). They are then thought to differentiate through an inevitable and irreversible process in which recombination between X and Y chromosomes is shut down and the Y degenerates (see Figure 1). Ultimately, Y chromosomes are fated to disappear entirely (“born to be destroyed,”

[18]). Thus, sex chromosomes that are morphologically similar (homomorphic) must be evolutionarily young, and in time they too will degenerate.

## The Myths Deconstructed

These myths do not survive a survey of sex determination systems across the tree of life. To deconstruct these myths, we first provide background on the evolution of separate sexes. We then summarize the diversity of sex-determining mechanisms found among animals and plants and discuss the evolutionary forces that drive transitions among systems (Myth 1 revisited). This is followed by a summary of more recent findings on the underlying molecular genetics of sex determination (Myth 2 revisited) and a deconstruction of common misconceptions of sex chromosome evolution in humans and other species (Myth 3 revisited). We conclude with an outlook for future research that might improve our understanding of how and why sex determination evolves so rapidly in many animals and plants.

## The Evolution of Separate Sexes

While the evolution of anisogamy led to the evolution of male and female functions, the evolution of separate sexes is not inevitable across lineages. Indeed, most flowering plants (94%, [19]) have both male and female sex organs within a single individual and often within the same flower. By contrast, hermaphroditism is rare among animals considered as a whole (about 5% of all species), which is largely due to the absence of hermaphrodites in the species-rich insects, but it is common in many other animal taxa, including fish and many invertebrates (most snails, corals, trematodes, barnacles, and many echinoderms) [20]. Hermaphrodites can mate with each other and benefit from the advantages of sex by mixing their genomes, but when mates are difficult to find, hermaphrodites can also escape the need for a reproductive partner by self-fertilization (which, however, may produce low-fitness offspring due to “inbreeding depression;” see below). This advantage of reproductive assurance is particularly pronounced in sessile animals—like corals—and plants, which cannot move to find a mate [21,22]. Thus there is a clear advantage to combining both male and female functions within an individual, especially in taxa with low mobility.

However, in some plants and most animals, species are driven to separate the sexes. This can be achieved in several ways. One partial solution is the spatial

separation of male and female gonads in the same individual, as in monoecious plants with separate male and female flowers (e.g., maize) and in most hermaphroditic animals. Alternatively, male and female function can be separated in time within an individual, as found in many plants (“dichogamy,” [23]) and some animals (“sequential hermaphroditism,” [24]); slipper shells, for example, are born male and become female later in life. Finally, male and female reproductive organs can be segregated into different individuals, as in some plants (such as papaya and cannabis) and most animals.

Separate sexes have evolved independently many times among plants and animals, which suggests that there must be an evolutionary cost to hermaphroditism, at least in some groups. Two major hypotheses have been proposed to explain the evolution of separate sexes. The first hypothesis is that there are trade-offs between male and female function, such as when mating displays enhance male fitness but decrease female fitness. In this case, individuals can gain reproductive advantages by specializing as a male or female [25]. Direct evidence for the trade-off hypothesis is sparse [26], and observations consistent with it come from hermaphroditic great pond snails, which reallocate resources to female function when sperm production is experimentally abolished [27], and from strawberries, in which increased pollen production comes at the cost of reduced seed set [28]. Indirect evidence of a trade-off comes from the fact that many asexual animals [29] and plants [30] that still have residual sperm/pollen production evolve reduced investment in male gametes over time, suggesting that doing so increases female function. The second major hypothesis is that separate sexes evolve as a means to avoid self-fertilization, which can produce low-fitness offspring because of the exposure of recessive deleterious alleles (“inbreeding depression”) [31]. Empirical evidence for inbreeding depression is widespread in animals and plants [32,33]; for instance, in the Hawaiian endemic plant genus *Scheidia*, high inbreeding depression promotes the evolution of dioecy [34].

When separate sexes are favored, the transition can occur via several evolutionary pathways. Separate sexes may evolve from hermaphrodites either by gradual increases in sex-specific investment or rapidly by the appearance of male- or female-sterility mutations (Figure 2). The latter occurs regularly in plants, often generating mixed sexual systems, such as

### Box 1. From Mating Types to Sexes

Meiotic sex likely has a single origin, which dates back to the origin of eukaryotes [144,145]). While most eukaryotes display some form of meiotic sex, many lack differentiated male and female gametes—a situation referred to as isogamy. Even with isogamy, however, mating is often not random but requires that joining cells differ at a mating type (MAT) locus. Mating types might have evolved to orchestrate the developmental transition from the haploid to the diploid phase of the life cycle [146,147]: *plus* and *minus* gametes express complementary transcription factors, encoded by different alleles at the MAT locus; these combine in the zygote into heterodimers that silence the genes expressed in the haploid phase and switch on the diploid program.

Isogamy permits a theoretically unlimited number of mating types; high numbers increase the probability that randomly mating partners display complementarity. Most basidiomycete fungi, for instance, present two independent MAT loci (and are therefore said to be tetrapolar, because a single meiosis can produce cells of four distinct mating types); each locus can be multiallelic, resulting in up to thousands of different mating types. Alternatively, a low probability of encountering complementary partners might have driven a transition to homothallism observed in many ascomycete fungi, which refers to a mating compatibility between genetically identical individuals. Homothallism evolved via genic capture: a single genome harbors complementary mating-type alleles, which are differentially expressed in *plus* and *minus* gametes. Mating-type switching in yeasts allows different cells from the same clone to express complementary mating types, and thus enter the diploid phase of their life cycle.

Anisogamy (small male and large female gametes) evolved independently in many eukaryotic lineages, including several different groups of protists (such as red algae, brown algae, apicomplexa, dinoflagellates, and ciliates; [148]), as well as most plants and animals. The transition towards anisogamy is thought to result from disruptive selection [1,149,150]: given opposing pressures to simultaneously maximize the number of gametes, their encounter rate, as well as the mass and ensuing survival of resulting zygotes, the fitness of both partners is often maximized when one interacting gamete is small and mobile, while its large and sessile partner provides the resources required for zygote development. Intermediate gametes do worse than small ones in terms of mobility and numbers, and worse than large ones in terms of provisioning. Such constraints largely explain why sexes (at the gametic level) are two and only two, and why anisogamy independently evolved in many lineages. At the molecular level, one route to anisogamy is by the incorporation of genes controlling gamete size into the MAT region [151]. Further extensions of the MAT region, possibly involving additional sex-antagonistic genes, led to the U and V chromosomes characterizing male and female gametophytes, as found, e.g., in mosses and liverworts [152].

Importantly, the evolution of anisogamy does not require the evolution of separate sexes, because hermaphrodites can produce both sperm and eggs. Similarly, several unicellular organisms that are anisogamous, such as apicomplexa and dinoflagellates, can make cells that produce sperm as well as cells that produce eggs. The evolution of completely separate sexes, where individuals cannot give rise to both sperm and egg descendants, is thought to be fairly derived and is found primarily among multicellular organisms with rare unicellular exceptions (e.g., the ciliate *Vorticella* [153] and several dioecious diatoms [154]).

into fertile males and females is a fundamental developmental process. Contrary to Myth 1, however, diverse mechanisms are used to determine sex [3,4] (Figure 3, Figure 4; Box 2). All crocodiles, most turtles, and some fish exhibit temperature-dependent sex determination; *Wolbachia* infections override existing sex determination systems in many arthropod species and either kill/sterilize males or transform them into phenotypic females; male scale insects eliminate their father's genome after fertilization; marine worm Bonellidae larvae develop as males only if they encounter a female; and many plants and animals—including some snails and fish—change sex during their lifetime in response to environmental or social cues [3,37].

In fact, sex determination is a rapidly evolving trait in many lineages (Figure 3), and sometimes closely related species, or populations of the same species, have different modes of sex determination [3,4,38]. Houseflies, for example, normally have XY sex chromosomes, but dominant masculinizing and feminizing alleles on other chromosomes exist in some populations that override sex determination by the XY chromosomes [39]. This variety has stimulated investigation into what evolutionary forces drive the turnover of sex determination mechanisms, what molecular mechanisms underlie the different modes of sex determination, and why sex determination is labile in some taxa and not in others.

### Genotypic versus environmental sex determination

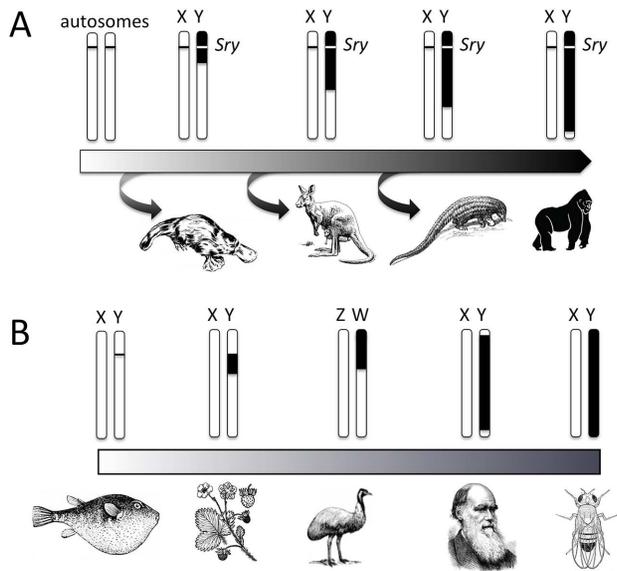
With *genotypic sex determination* (GSD), which occurs in the majority of species with known sex-determining mechanisms, genetic elements specify whether individuals are female or male. In many animals and some plants, however, the switch to develop into a female or male does not lie in the genes. With *environmental sex determination* (ESD), external stimuli control sex determination, such as temperature in reptiles [40], photoperiod in marine amphipods and some barnacles [41,42], and social factors in many coral-reef-dwelling fish and limpets [43,44]. Exactly how the environment triggers sex development has remained an open question, although a recent study found that methylation provided the link in European sea bass [45]. In many species, the line between GSD and ESD is blurred, with certain environments altering the (otherwise genetically determined) sex of developing offspring [46]. For example, tongue sole have differentiated ZW sex chromosomes, but

gynodioecy (mixtures of females and hermaphrodites) and androdioecy (mixtures of males and hermaphrodites). Figure 2 highlights the possible pathways for the evolution of separate sexes from a hermaphrodite ancestor and illustrates their relation to sex chromosome evolution. While we have emphasized the evolutionary transition from hermaphroditism to separate sexes, several examples are known where the opposite transitions

occur (e.g., [35,36]), indicating that the conditions favoring the separation of male and female function are not always present.

### Myth 1 Revisited—Sex-Determining Mechanisms Are Diverse and Can Evolve Rapidly

In animals and plants that have evolved separate sexes, accurate differentiation



**Figure 1. Sex chromosome differentiation.** **A.** Reconstructed evolutionary path of sex chromosome differentiation in humans. Sex chromosomes originate from autosomes that acquired a sex-determining function (the *Sry* gene) after their split from monotremes. Suppression of recombination between the sex chromosomes, associated with degeneration of the non-recombining region of the Y chromosome, results in the morphological and genetic differentiation of sex chromosomes. Recombination suppression occurred in multiple episodes along the human X and Y chromosome, forming so-called evolutionary strata. The oldest stratum is shared between eutherian mammals and marsupials, while the youngest stratum of humans is primate-specific. **B.** The degree of sex chromosome differentiation ranges widely across species, spanning the entire spectrum of homomorphic to heteromorphic sex chromosomes, from a single sex-determining locus, as seen in pufferfish, a small differentiated region (strawberry and emu), most of the sex chromosomes apart from short recombining regions (humans), to the entire sex chromosome pair, as seen in *Drosophila*. Note that the sex chromosomes are not drawn to scale.

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ZW embryos develop into males when incubated at high temperatures, and sex reversal is accompanied with substantial methylation modification of genes in the sex determination pathway [47].

ESD is favored over GSD when specific environments are more beneficial to one sex [3], selecting for sex-determining mechanisms that match each sex to its best environment. For example, in some gobies and wrasses, nest sites are limited, and a male's ability to defend his nest depends on body size; individuals tend to start life as females, and only become males once they are sufficiently large to successfully defend a nesting site [48]. The reverse transition, from ESD to GSD, is thought to be favored when the environment is unpredictable or not variable enough, in which case ESD could produce strongly skewed sex ratios or intersex individuals [3]. Indeed, snow skinks, which are small, live-bearing lizards, have different sex-determining mechanisms in different environments. ESD occurs at low altitudes where early birth is advantageous for females and the variance in temperature

between years is low. GSD predominates at high altitudes where there is no advantage for early-born females and between-year variance in temperature is high [49]. In this situation, ESD produces optimal sex ratios at low elevations, while GSD prevents extreme sex ratios at high altitudes. Importantly, global climate change poses a threat to species with temperature-dependent sex determination if they cannot adapt rapidly enough to avoid biased sex ratios [50]. Another threat to ESD systems comes from within: they may be prone to invasion by genetic elements that favor biased sex ratios (see below).

### Genomic conflict and transitions in sex determination

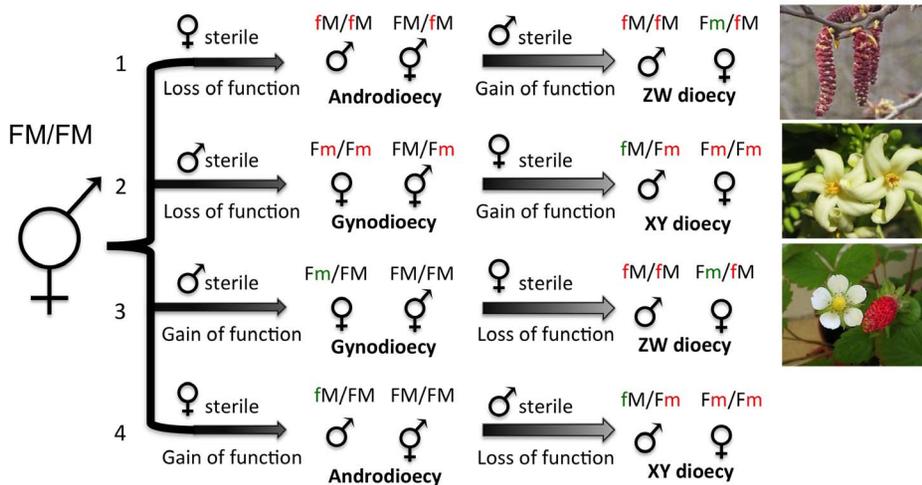
More generally, selection on the sex ratio can trigger transitions between and among different ESD and GSD systems [3]. Sex-biased inheritance patterns of different genetic elements—such as sex chromosomes, organelles, or endosymbionts—create a conflict over which sex is preferred, and can drive the evolution of

male- or female-biased sex ratios. In populations with a skewed sex ratio, selection on autosomal genes typically favors equal investment in males and females [51,52], and a new GSD or ESD system can become established if it restores a more even sex ratio. An equal number of males and females is, however, not always favored, even among autosomal genes (e.g., with local mate competition, [53]). In this case, selection for biased sex ratios can favor the establishment of a new sex-determining system [54].

Many examples are known of sex chromosomes, organelles, and endosymbionts that bias the primary sex ratio. Meiotic drive, where genetic elements bias the proportion of gametes that carry them, can create male-biased sex ratios if they are located on the Y or Z chromosomes (as seen in many *Drosophila* species [55]), whereas driving X or W chromosomes create female-biased sex ratios (found in *D. simulans* [56], stalk-eyed flies [57], and rodents [58]); autosomal genes that restore unbiased sex ratios are known in many systems. Cyto-nuclear conflict arises because cytoplasmic factors such as mitochondria or chloroplast are often inherited only through the mother, and they favor production of females, while autosomal genes are inherited through both sexes and favor more equal sex ratios. Cytoplasmic male sterility encoded by mitochondria has been widely reported in plants, including maize, petunia, rice, common bean, and sunflower [59], as have nuclear-encoded male fertility restorer genes [60]. Likewise, cellular endosymbionts are only transmitted through the mother and can create maternally inherited female-biased sex ratios; examples include male-killing bacteria in butterflies and *Drosophila* [61,62]. Recurrent invasions of sex ratio distorters and their suppressors can result in rapid transitions among sex determination mechanisms between species, and may be a major force contributing to the diversity of sex-determining mechanisms observed across the tree of life.

### Turnover of sex chromosomes

In species with genotypic sex determination, the chromosome pair that determines sex can change rapidly over time. Transitions are particularly likely when the ancestral sex chromosome exhibits little genetic differentiation, since WW or YY combinations are then less likely to be lethal (Figure 5). New sex-determining genes (or copies of the original gene in a new location) can lead to transitions within and between different XY and ZW systems (Figure 5). Invasions of sex-deter-



**Figure 2. Evolutionary pathways from hermaphroditism to separate sexes.** Shown are two-step pathways involving intermediate male- and female-sterile individuals. Loss-of-function mutations (red) are assumed to be recessive, while gain-of-function mutations (green) are assumed to be dominant. Ancestral alleles are in black. M, male fertility allele; m, male sterility mutation; F, female fertility allele; f, female sterility mutation. Because loss of function mutations (red) are almost 50 times more frequent than gain of function mutations (green) in flowering plants, we would expect pathways 1 (e.g., some poplar species) or 2 (e.g., papaya) to arise earlier. Furthermore, transitions through gynodioecy, pathways 2 and 3 (e.g., strawberry) allow females to completely avoid inbreeding depression, while transitions through androdioecy are more costly because males must compete with hermaphrodites for fertilization and do not have any of their own ovules to fertilize. These theoretical arguments help to account for the prevalence of gynodioecy and the XY chromosome system (via pathway 2) observed in plants; nevertheless, all four pathways may be biologically relevant, although no known examples for pathway 4 currently exist.  
doi:10.1371/journal.pbio.1001899.g002

mining genes are facilitated when the new sex-determining gene (or a gene closely linked to it) has beneficial effects on fitness [63].

Sexually antagonistic selection, which occurs when a mutation is beneficial to one sex but detrimental to the other, can also drive transitions between sex determination by different pairs of chromosomes [64,65]. For example, if an allele of an autosomal gene is beneficial to males but harmful to females and becomes linked to a dominant masculinizing mutation, then chromosomes that carry both the male-beneficial and male-dominant alleles create a novel Y that can replace the ancestral mechanisms. Conversely, alleles that benefit females and harm males can create novel W chromosomes when linked to feminizing mutations. Turnover of sex chromosomes can also be triggered by the degeneration of the Y and W chromosome, which commonly follows the cessation of recombination [66,67], and will result in the replacement of a low-fitness Y or W chromosome with a nondegenerate one [68].

### Sex determination by the whole genome

In many animals, sex determination involves the entire genome. With haplodiploidy (found in about 12% of animal species, including all ants, wasps, and bees) and paternal genome elimination (found in

scale insects), males only transmit their maternal set of genes (see Figure 4; Box 2: Glossary). The loss of the paternal genome in sons benefits mothers but not fathers because these uniparental sons transmit more of a mother's genome to grandchildren than do biparental sons [3]. Females also experience a selective advantage from haplodiploidy (but not paternal genome elimination) because unfertilized eggs can develop and contribute to fitness when mating opportunities are rare.

Despite numerous theoretical predictions for how and why sex determination mechanisms change, many hypotheses remain untested. Only a small proportion of taxa have actually been characterized for their sex determination mechanisms, hindering the use of comparative methods to assess the factors associated with transitions between them. However, because sex determination changes so rapidly in many clades, we can catch these transitions *in action* to test theoretical predictions in a direct, experimental way.

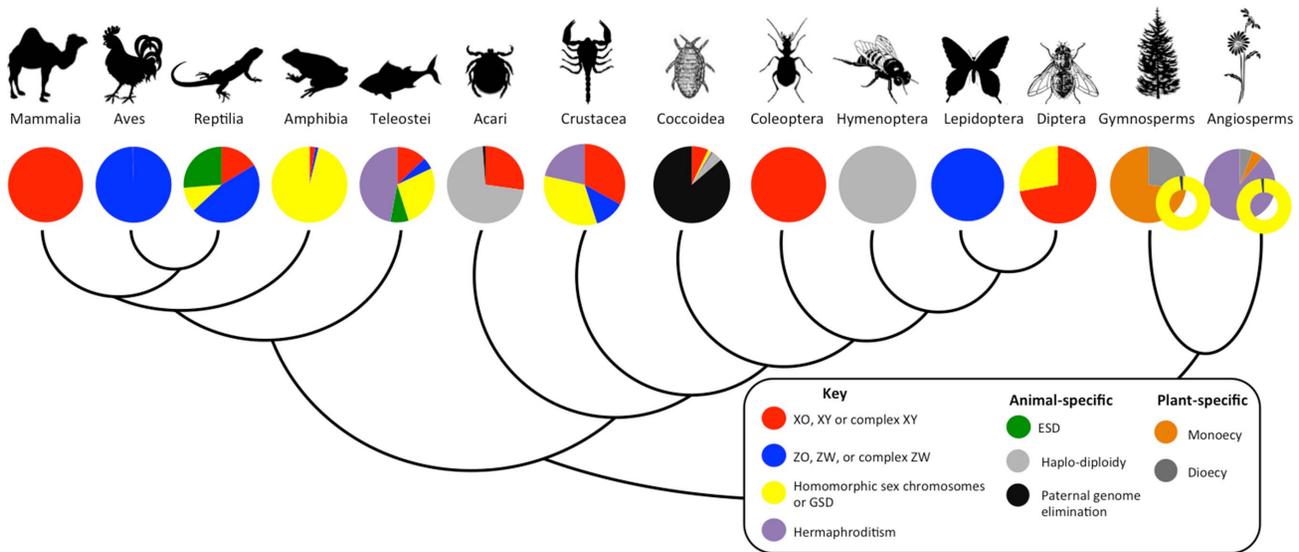
### Myth 2 Revisited—Multiple and Various Genes Can Determine Sex

The pathways that control sexual development have been well characterized at

the molecular level in *D. melanogaster*, *C. elegans*, and mammals. All three involve a master-switch sex-determining gene, which led to the birth of Myth 2. Although the simplicity of a single master-switch is alluring, this archetype of sex determination is clearly not universal. Below we discuss systems where sex is determined by multiple genes, recent molecular data on the nature and evolution of sex-determining genes, and how sex determination can vary in different parts of the body.

### Polygenic sex determination

In some species, sex determination is polygenic. For example, in zebrafish (*Danio rerio*), a key developmental model organism, sex is not controlled by a single master regulator but is instead a quantitative threshold trait with either a male or female outcome, which is determined by multiple regions in the genome [69–71]. While some of those regions contain genes known to play a role in sex determination in other organisms [70], there is an enduring mystery as to how these multiple loci and the environment interact to control downstream sexual differentiation in zebrafish. Zebrafish gonads develop as testes in the absence of signals from germ line cells, suggesting that the factors determining sex may regulate germ cell proliferation [72]. Sex as a threshold trait has been inferred in several other fish [73–75]



**Figure 3. Diversity of sex determination systems for representative plant and animal clades.** The bubble insert graph for the plant clades represents the relative proportion of species with documented sex chromosomes within plants with separate sexes. Vertebrates: Mammalia (placental, marsupial, and monotreme mammals), Aves (birds), Reptilia (turtles, snakes, crocodiles, lizards), Amphibia (frogs, toads, salamanders), and Teleostei (bony fishes). Invertebrates: Acari (mites and ticks), Crustacea (shrimps, barnacles, crabs), and Insects, which include Coccoidea (scale insects), Coleoptera (beetles), Hymenoptera (ants, bees, and wasps), Lepidoptera (butterflies), and Diptera (flies). Plants: Gymnosperms (non-flowering plants) and Angiosperms (flowering plants). doi:10.1371/journal.pbio.1001899.g003

and invertebrates [76], and further examples of multiple interacting loci controlling sex determination are no doubt waiting to be described. Indeed, in taxa where separate sexes evolved recently from a hermaphrodite ancestor, as is common in plants, multiple sex-determining loci are in fact expected, since at least two independent mutations—one suppressing male function, one suppressing female function—are necessary to produce separate sexes from a hermaphrodite (Figure 2). If separate sexes evolve by gradual increase in sexual investment from a hermaphrodite, sex determination may also be due to polygenic inheritance.

### The nature and evolution of sex-determining genes and pathways

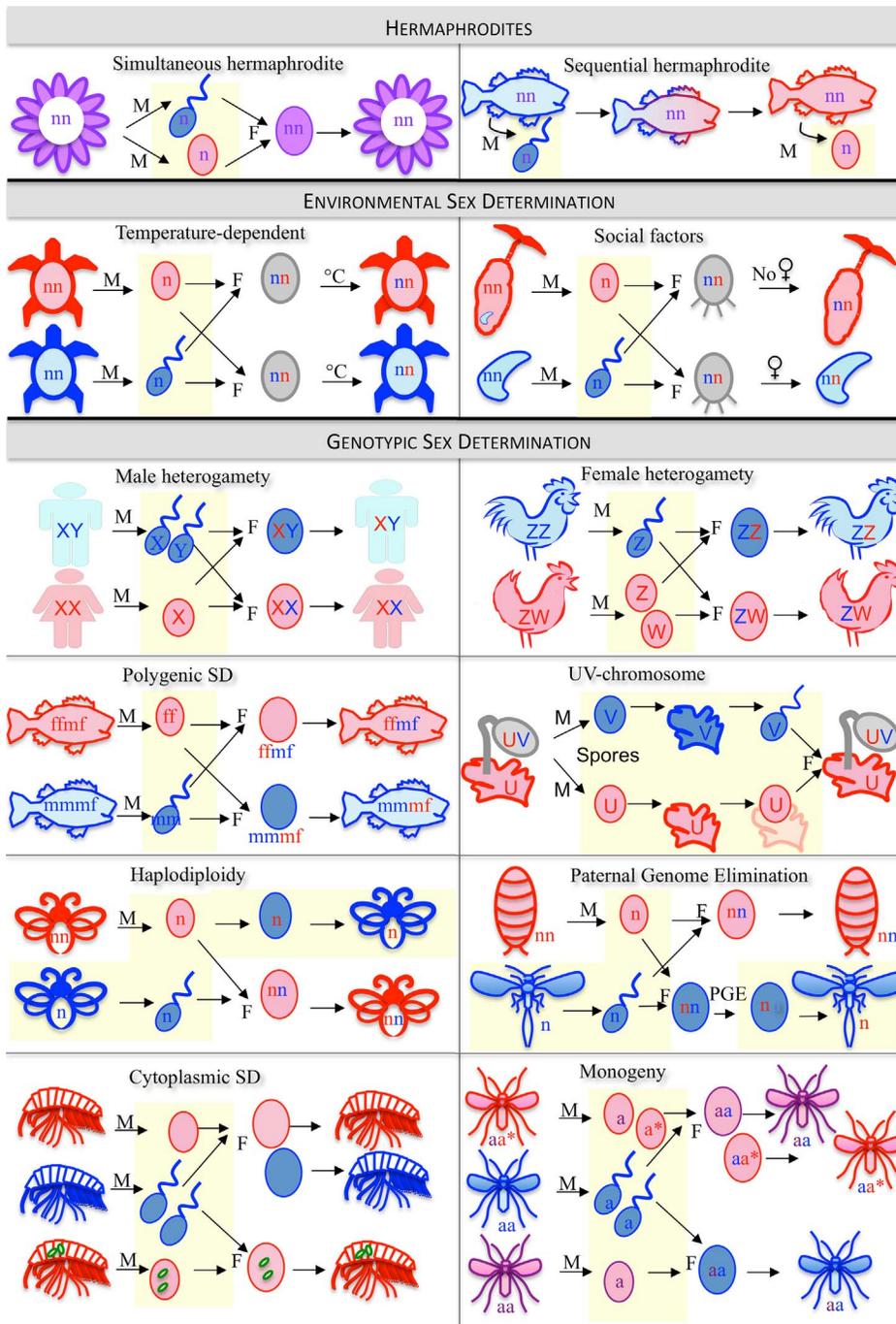
Some taxa have master-switch sex-determining genes that are highly conserved, such as the *Sry* gene in nearly all mammals [77]. In other lineages, such as fish from the genus *Oryzias* [78–80], the master-switch gene differs among closely-related species (Table 1). There is some empirical evidence for the repeated use of the same master sex determination switch genes in animals. For example, in vertebrates other than mammals, *dmrt1* (a DM family gene) and its paralogs act as the primary sex determination signal in African clawed frog (*Xenopus laevis*) [13],

chicken (*Gallus gallus*) [12], medaka fish (*Oryzias latipes*) [78,79], and possibly the smooth tongue sole (*Cynoglossus semilaevis*) [14]. In insects, paralogs of *transformer* (*tra*), a key gene in the sex determination cascade of *Drosophila*, have evolved as the primary switch in houseflies *Musca domestica* [17], as well as the haplodiploid wasp *Nasonia vitripennis* [15] and the honeybee *Apis mellifera* [16].

These data suggest that there are constraints on the types of genes that can be co-opted as master sex determination genes [81]. Nevertheless, there are several cases of switch genes with no homologs in closely related taxa. These include an immunity-related gene in rainbow trout (*Oncorhynchus mykiss*) [82] and *Sxl* in *Drosophila* [83], whose ortholog has a non-sex-related function in mRNA splicing in other flies [84]. The primary master sex-determining gene in the silkworm *Bombyx mori* is a W-derived female-specific piRNA (produced from a piRNA precursor termed *Fem*) that targets a Z-linked gene (named *Masc*), and silencing of *Masc* mRNA by *Fem* piRNA is required for female development [85]. Undoubtedly, many other sex determination genes remain to be found, making it unclear at present whether there truly are constraints on the types of genes that could evolve to be master control switches.

No master sex determination gene has been identified in dioecious plants, but genes that affect flower sex determination have been found [86,87]. Indeed, many genes may serve as potential targets for sex determination in plants, given that male or female sterility can evolve in various ways [86]. For example, 227 male-sterility genes have been identified in rice, with at least one male-sterility gene found on each of rice's 12 chromosomes—hence each autosome could, in principle, evolve a sex-determining function [88]. This abundance and diversity within a single species indicates that the initial evolution of separate sexes is unlikely to be limited to a scant handful of master genes.

In sharp contrast with the diversity of primary sex-determining signals, some key regulatory genes play conserved roles in the molecular pathways leading to male or female gonad development across invertebrates and vertebrates, such as the *doublesex-mab3* (DM) family genes [89,90]. Despite profound differences in the mode of sex determination and the identity of the master-switch genes, DM genes are specifically expressed in the developing gonads of almost all animals, including vertebrates (mammals [91], birds [92], turtles and alligators [93–95], amphibians [96], and fish [97]) and invertebrates (*Drosophila* [98], hymenoptera [99], crustaceans



**Figure 4. Schematic overview of some sex determination (SD) mechanisms.** M refers to meiosis, F to fertilization. Haploid stages (n) are indicated as shaded areas and diploid stages (nn) are unshaded. **Hermaphrodites:** Most flowering plants (and gastropods and earthworms) simultaneously contain both male and female sexual organs (*simultaneous hermaphrodites*). Many fish and some gastropods and plants are *sequential hermaphrodites*; clownfish, for example, are born males and change into females, while many wrasses or gobies begin life as females and then change to males. **Environmental Sex Determination:** In turtles and some other reptiles, sex is determined by incubation temperature of the eggs (*temperature-dependent sex determination*). **Social factors** can act as primary sex-determining cues: sexually undifferentiated larvae of the marine green spongeworm that land on unoccupied sea floor develop into females (and grow up to 15 cm long), while larvae that come into contact with females develop into tiny males (1–3 mm long) that live inside the female. **Genotypic Sex Determination:** Almost all mammals and beetles, many flies and some fish have *male heterogamety* (XY sex chromosomes), while *female heterogamety* (ZW sex chromosomes) occurs in birds, snakes, butterflies, and some fish. In mosses or liverworts, separate sexes are only found in the haploid phase of the life cycle of an individual (*UV sex chromosomes*). In some flowering plants and fish, such as zebrafish, sex is determined by multiple genes (*polygenic sex determination*). In bees, ants, and wasps, males develop from unfertilized haploid eggs, and females from fertilized diploid eggs (*haplodiploidy*), while males of many scale insects inactivate or lose their paternal chromosomes (*paternal genome elimination*). In some species, sex is under the control of cytoplasmic elements, such as intracellular parasites (e.g., *Wolbachia*) in many insects or mitochondria in many flowering plants (*cytoplasmic sex determination*). In some flies and crustaceans, all offspring of a particular individual female are either exclusively male or exclusively female (*monogeny*). doi:10.1371/journal.pbio.1001899.g004

[100,101], and mollusks [102,103]). Thus, the evolution of sex-determining pathways, at least in animals, appears to occur by the recruitment of new master-switches controlling sexual fate, while the downstream developmental pathways that regulate gonadal differentiation are retained [10,81,104], although the function of some of these downstream elements appears to diverge among lineages [105]. Characterization of polygenic sex determination systems and identification of master sex determination genes across kingdoms will provide insight into the mechanistic constraints limiting the evolution of sex determination pathways.

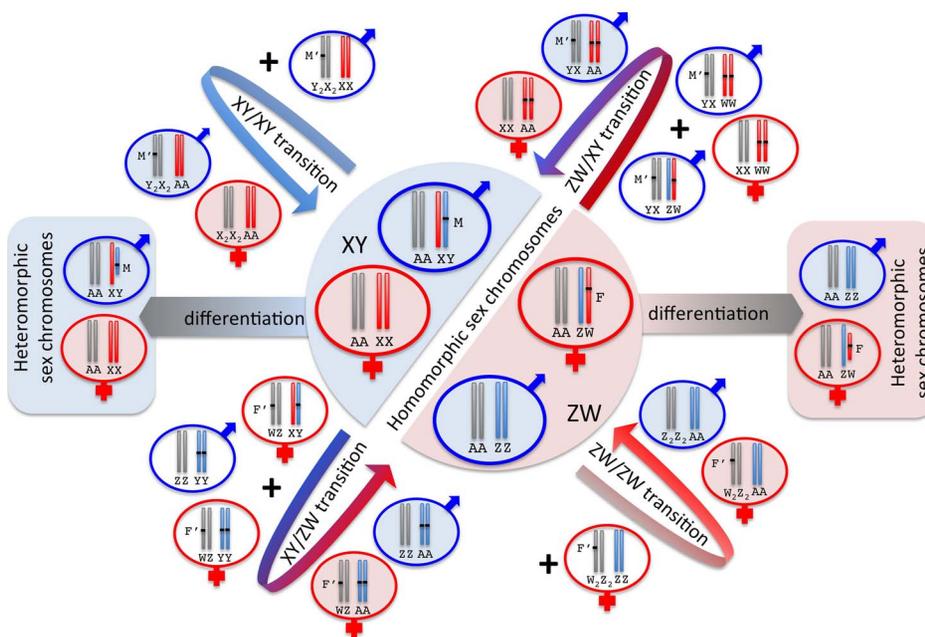
### Sex determination: soma vs. germ line

Sex determination can also differ with respect to where in the body sex is determined. In humans, sex is determined in the developing gonad, and gonadal sex hormones in turn trigger sex determination and differentiation in nongonadal tissues. By contrast, in birds, *Drosophila*, and nematodes [106–109], sexual differentiation is a cell-autonomous process, although secreted signaling molecules can play a role in generating sexual dimor-

phism in somatic tissues. Studies in *Drosophila* have shown that only a subset of cells express the genes of the sex determination cascade and have a sexual identity [106]. Cell-autonomous sex determination can result in the formation of gynandromorphs—individuals that contain both male and female characteristics, found in birds and many insects, including butterflies and beetles. Sex determination can also be regulated differently in the soma versus the germ line of the same species [110,111]. In houseflies [112] and some frogs [113] and fish [114–116], transplantation experiments indicate that the sex of germ cells depends on the surrounding soma, i.e., XX germ cells transplanted into male soma form sperm, and XY germ cells transplanted in a female soma form oocytes. In contrast, germ cells in *Drosophila* [117] and mammals [118] receive signals from the surrounding somatic gonad, but they also make an autonomous decision during germ line sexual development; this may also be true for chickens [107]. In these animals, the “sex” of the soma must match the “sex” of the germ cells for proper gametogenesis to occur. If XX germ cells are transplanted into male soma they do not form sperm, and XY germ cells transplanted into female soma fail to form oocytes.

### Myth 3 Revisited—Sex Chromosomes’ Eternal Youth

Heteromorphic sex chromosomes evolve from autosomes that are initially identical but then stop recombining and differentiate. Recombination suppression is favored when it links together sexually antagonistic alleles and sex-determining loci (i.e., a male-beneficial allele and a male-determining gene on a Y chromosome, or a female-beneficial allele and a female-determining gene on a W chromosome). A side effect of repressed recombination on Y and W chromosomes is that natural selection is inefficient (reviewed in [4,5]), which can result in the loss of most of their genes. Y or W degeneration has occurred in many animal taxa with heteromorphic sex chromosomes, including mammals [119], many birds [120], snakes [121], and many insects [122,123], along with some plants, including *Rumex* [124]. In the most extreme cases, the Y or W is entirely lost, resulting in so-called XO and ZO systems. According to Myth 3, differentiation of sex chromosomes is evolutionarily inevitable, and the degree of heteromorphism reflects their age (Figure 5). However, as we explain below, evidence from a broad array of organisms indicates that the link between sex chro-



**Figure 5. Transitions versus differentiation of sex chromosomes.** Transitions between homomorphic sex chromosomes result from new masculinizing (M') or feminizing (F') mutations that invade an existing XY or ZW system to create a new chromosome pair (in grey) that harbors the sex-determining gene (additional transitional karyotypes are indicated by unshaded circles). XY→XY transitions result in the loss of the ancestral Y (and ZW→ZW transitions cause loss of the ancestral W). Transitions between XY and ZW systems result in some offspring that are homozygous for the Y (blue) or W (red) chromosome and are thus more likely if the chromosomes have similar gene content but become increasingly difficult if these chromosomes have degenerated (side boxes on left and right), causing YY and WW individuals to be less fit. doi:10.1371/journal.pbio.1001899.g005

## Box 2. Glossary of Sex-Determining Mechanisms

- Hermaphrodites: individuals that contain both male and female sex organs.
- Simultaneous hermaphroditism: male and female sexual organs coexist in one individual (e.g., most flowering plants and earthworms, many terrestrial gastropods).
- Sequential hermaphroditism: individuals change sex at some point during their life (e.g., many fish, snails, and some plants).
- Dioecy (plants) or gonochorism (animals): individuals are either male or female throughout their life.
- Environmental sex determination: sex is triggered by environmental cues, such as temperature, pH, social interactions, and seasonality (e.g., many reptiles and some fish).
- Genotypic sex determination: an individual's sex is established by its genotype (e.g., mammals, birds, amphibians, most insects, some reptiles and fish, and some plants).
- Male heterogamety: type of genotypic sex determination in which males are heterozygous for the sex-determining locus (termed X and Y, as seen in therian mammals and *Drosophila*).
- Female heterogamety: type of genotypic sex determination in which females are heterozygous for the sex-determining locus (termed Z and W, as seen in birds, snakes, butterflies, and ginkgo trees).
- UV sex determination: separate sexes are only found in the haploid phase of the life cycle of an individual (e.g., mosses or liverworts).
- Polygenic sex determination: sex is determined by multiple genes (e.g., some fish and flowering plants).
- Haplodiploidy: males develop from unfertilized, haploid eggs, and females from fertilized, diploid eggs (e.g., bees, ants, and wasps).
- Paternal genome elimination: paternal chromosomes in males are inactivated or lost after fertilization, leaving males functionally haploid (e.g., many scale insects).
- Cytoplasmic sex determination: sex is under the control of cytoplasmic elements, such as intracellular parasites (e.g., *Wolbachia* in many insects) or mitochondria (e.g., cytoplasmic male sterility in flowering plants).
- Monogeny: all offspring of a particular individual female are either exclusively male or exclusively female (e.g., some flies and crustaceans).
- Sexual reproduction: the mixing of genomes via meiosis and fusion of gametes.
- Sex: the sexual phenotype of an individual.
- Sex determination: the mechanism by which the sexual phenotype of an individual is established in a given species.
- Sex chromosome: a chromosome involved with determining the sex of an individual.
- Autosome: a chromosome not involved with determining the sex of an individual (i.e. any chromosome that is not a sex chromosome).
- Y degeneration: the loss of genetic information on the non-recombining Y chromosome.
- Homomorphic sex chromosomes: sex chromosomes that are morphologically indistinguishable.
- Heteromorphic sex chromosomes: sex chromosomes that are morphologically distinct.
- Sexually antagonistic selection: selection for a trait that benefits one sex to the detriment of the other sex.
- Gynodioecy: a breeding system that consists of a mixture of females and hermaphrodites.
- Androdioecy: a breeding system that consists of a mixture of males and hermaphrodites.
- Meiotic drive (also called segregation distortion): a system in which genetic elements termed segregation distorters bias the proportion of gametes that carry them, resulting in over- or under-representation of one gametic type (i.e. non-mendelian segregation).
- Nucleo-cytoplasmic conflict: conflict in inheritance patterns between the nuclear genome and organelle genomes that are transmitted only maternally.
- Gynandromorphs: individuals that contain both male and female characteristics.

mosome heteromorphism and age is often far from direct.

### Not all sex chromosomes become differentiated

Differentiation is often seen as the default path of sex chromosome evolution, but contrary to Myth 3, some ancient sex chromosomes recombine and are undifferentiated over most of their length. Examples are found in python snakes and ratite birds, whose homomorphic sex chromosomes are about 140 and 120

million years old, respectively [121,125,126], i.e. almost as old as the heteromorphic sex chromosomes of therian mammals (about 180 million years old).

How do some ancient sex chromosomes avoid differentiation? One hypothesis is that occasional X-Y recombination purges deleterious alleles on the Y. A mechanism proposed for tree frogs is that XY embryos are occasionally sex-reversed, and so the X and Y recombine when these embryos develop into females [127,128]. Second, some taxa may have few genes under

sexually antagonistic selection on their sex chromosomes and thus avoid selection to suppress recombination between the X and Y [129]. Third, sexually antagonistic selection can be resolved by other means, such as the evolution of sex-specific expression [130]. Sexually antagonistic alleles can accumulate along the sex chromosomes, and sex-specific expression will confine the product of such alleles to the sex they benefit, thereby eliminating the selective pressure for recombination suppression. Consistent with this last

**Table 1.** Known master sex-determining genes in vertebrates and insects, and their paralogs.

Species	Master sex determining gene	Sex-determining mechanisms	Gene paralog	Paralog function	Reference
mammals	<i>Sry</i>	sex-determining Y	<i>Sox3</i>	HMG-box transcription factor	[77]
chicken ( <i>Gallus gallus</i> )	<i>dmrt1</i>	dose-dependent Z	-	SD pathway transcription factor	[12]
African clawed frog ( <i>Xenopus laevis</i> )	<i>dmW</i>	sex-determining W	<i>dmrt1</i>	SD pathway transcription factor	[13]
medaka ( <i>Oryzias latipes</i> )	<i>dmrt1Y</i>	sex-determining Y	<i>dmrt1</i>	SD pathway transcription factor	[78,79]
( <i>Oryzias luzonensis</i> )	<i>gsdfY</i>	sex-determining Y	<i>gsdf</i>	secretory protein in SD pathway	[80]
Patagonian pejerrey ( <i>Odontesthes hatcheri</i> )	<i>amhY</i>	sex-determining Y	<i>amh</i>	anti-Mullerian hormone	[155]
rainbow trout ( <i>Oncorhynchus mykiss</i> )	<i>sdY</i>	sex-determining Y	<i>lrf9</i>	interferon regulatory factor	[82]
tiger pufferfish ( <i>Takifugu rubripes</i> )	<i>amhr2</i>	dose-dependent X	<i>amhr</i>	anti-Mullerian hormone receptor	[156]
smooth tongue sole ( <i>Cynoglossus semilaevis</i> )	<i>dmrt1</i>	dose-dependent Z	-	SD pathway	[14]
fruit flies ( <i>Drosophila</i> )	<i>Sxl</i>	dose-dependent X	CG3056	mRNA splicing, non-sex specific	[83,84]
housefly ( <i>Musca domestica</i> )	<i>F</i>	sex-determining W	<i>tra</i>	SD pathway switch splice factor	[17]
silkworm ( <i>Bombyx mori</i> )	<i>Fem</i>	sex-determining W	-	piRNA	[85]
honeybee ( <i>Apis mellifera</i> )	<i>csd</i>	haplodiploid	<i>tra</i>	SD pathway switch splice factor	[16]
wasp ( <i>Nasonia vitripennis</i> )	<i>Nvtra</i>	haplodiploid	<i>tra</i>	SD pathway switch splice factor	[15]

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possibility, the recombining, non-differentiated region along the sex chromosomes of the emu (a ratite bird) contains an excess of genes whose expression is sex-biased, relative to autosomes [126].

### Y chromosomes are not doomed

Y chromosome degeneration has prompted the suggestion that the human Y will eventually disappear [131–133], a claim based on the naïve assumption of a constant rate of gene loss from the Y over time. However, theory predicts that the rate of gene decay on the Y decreases over evolutionary time and should halt on an old, gene-poor Y chromosome [67,134]. Recent comparative genomic studies support this hypothesis as the gene content of the primate Y chromosome has been stable over the last 25 million years, suggesting that an equilibrium gene content has been reached in humans [135]. Moreover, old gene-poor Y chromosomes that are tens of millions of years old, such as the *Drosophila* Y [136], actually show a net rate of gene gain rather than gene loss [137]. Thus, the Y chromosome can be a stable and important component of the

genome in many species, and may even prevent turnover of sex-determining mechanisms (see below).

### Evolutionary traps and conserved sex-determining systems

In contrast to the lability of sex determination mechanisms in some groups, eutherian mammals, birds and many insects exhibit virtually no variation in how sex is determined (Figure 3). This stability could be due to an absence of genetic variation, particularly when multiple genetic steps are required for a transition to a new sex-determining system (Figure 2). Mutations are known, however, that override sex determination (Table 1) [138], suggesting that the conservation is not due to a lack of genetic variation. Instead, evolutionary traps may stabilize sex-determining systems for long spans of evolutionary time.

Heteromorphic sex chromosomes may act as just such a trap. Transitions between XY and ZW systems that create YY or WW individuals are prevented when Y or W chromosomes lack essential genes (Figure 5). Also, if the Y (or W) chromosome

has evolved sex-essential genes, such as spermatogenesis genes located on the human and *Drosophila* Y, sex chromosome transitions are only possible if these genes are moved to another chromosome, since the old Y, along with its genes, is lost during the transition (Figure 5). Overall, phylogenetic patterns in vertebrates or insects [3,139] are consistent with the notion that heteromorphic sex chromosomes constrain shifts in sex determination mechanism, but several notable exceptions exist in both groups. In rodents, for example, many species with unusual sex-determining systems can be found: XY females in some lemming species, X0 females or XX males in vole species, and X0 females and males in some Japanese spiny rats and mole voles [140]. Likewise, some insect groups are known that harbor variation in sex chromosome karyotype among species; in grasshoppers, fusions between sex-chromosomes and autosomes combined with Y-degeneration and/or Y-loss have created much variation in sex chromosome karyotype, including species with multiple X or Y

chromosomes [141]; true fruit flies (Tephritidae) that contain both XY and ZW species [142]; or blowfly species that have secondarily lost their heteromorphic sex chromosomes [143].

How much sex chromosome heteromorphism is required to create a trap, and how strong this trap is, remains unknown. To date, only one example of the reversal of an ancient sex chromosome back to an autosome has been characterized. Specifically, all *Drosophila* species contain an autosome that was formerly an X chromosome: the dot chromosome. This chromosome still has a minor feminizing role during sex determination, is targeted by a chromosome-specific regulatory mechanism similar to dosage compensation of the X, and its patterns of biased gene expression during early embryogenesis, oogenesis, and spermatogenesis resemble that of the current X in *Drosophila* [136]. The retention of the specialized genomic architecture of highly differentiated sex chromosomes on the dot chromosome illustrates the numerous barriers to sex chromosome turnover that exist in highly heteromorphic systems, even though there are some cases where these are overcome.

Haplodiploidy also appears to be an evolutionary trap. While it has arisen a few dozen times, the reverse transition has not been reported [3]. Transitions from or to haplodiploidy require changes in genetic architecture and meiotic mechanisms,

which are likely more complex than a simple change in a master-switch sex-determining gene. Furthermore, females switching from haplodiploidy would lose the fitness benefit associated with producing uniparental sons.

Systems that involve interacting somatic and germ line sex determination mechanisms may also limit transitions of sex-determining mechanisms, since changes in either germ line sex or somatic sex alone may produce infertile individuals [111]. Thus, while sex determination is generally characterized by diversity and turnover, some sex-determining systems appear to be more evolutionarily stable than others [3].

## Outlook

Studying the forces that drive the evolution of sex determination has mainly come from theoretical works, with little empirical data. However, the genomic revolution has allowed researchers to address scientific questions and tackle novel biological systems at the molecular level. As new genomic approaches increase the pace of discovery and characterization of sex determination in non-model organisms, we anticipate that comparative phylogenetic methods will be key to examining the roles of various ecological and genetic factors that drive changes in sex determination mechanisms. Additionally, genomic data make it increasingly possible to map sex-determining loci

from closely related species and to identify the evolutionary mechanisms hypothesized to cause transitions among sex-determining systems. Finally, comparative and functional genomic data will allow researchers to address how new master sex determination genes are incorporated into existing genetic networks controlling sexual development. A full understanding of the diversity of sex determination mechanisms will require that we expand the taxonomic breadth of study systems well beyond classic model organisms. Promising models include dipteran insects, such as houseflies or chironomids; teleost fish; and reptilian clades, including turtles and lizards; as well as plant genera, such as strawberries, that show variation within and between species in how sex (or gender in plants) is determined. Integrative and interdisciplinary approaches across the tree of life will illuminate the diversity of sex determination and yield exciting new insights of how and why sex determination evolves in animals and plants.

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## References

- Bell G (1982) The masterpiece of nature. Berkeley: University of California.
- Otto SP (2009) The evolutionary enigma of sex. *Am Nat* 174 Suppl 1: S1–S14.
- Bull JJ (1983) Evolution of Sex Determining Mechanisms. Menlo Park, CA: Benjamin Cummings.
- Charlesworth B (1996) The evolution of chromosomal sex determination and dosage compensation. *Curr Biol* 6: 149–162.
- Bachtrog D (2013) Y-chromosome evolution: emerging insights into processes of Y-chromosome degeneration. *Nat Rev Genet* 14: 113–124.
- Eggers S, Sinclair A (2012) Mammalian sex determination—insights from humans and mice. *Chromosome Res* 20: 215–238.
- Ono M, Harley VR (2013) Disorders of sex development: new genes, new concepts. *Nat Rev Endocrinol* 9: 79–91.
- Campbell NA (1996) Biology. Menlo Park, CA: Benjamin/Cummings Publishing Co.
- Pane A, Salvemini M, Delli Bovi P, Polito C, Saccone G (2002) The transformer gene in *Ceratitis capitata* provides a genetic basis for selecting and remembering the sexual fate. *Development* 129: 3715–3725.
- Wilkins AS (1995) Moving up the hierarchy: a hypothesis on the evolution of a genetic sex determination pathway. *Bioessays* 17: 71–77.
- Vollf JN, Nanda I, Schmid M, Schartl M (2007) Governing sex determination in fish: regulatory putches and ephemeral dictators. *Sex Dev* 1: 85–99.
- Smith CA, Roeszler KN, Ohnesorg T, Cummins DM, Farlie PG, et al. (2009) The avian Z-linked gene DMRT1 is required for male sex determination in the chicken. *Nature* 461: 267–271.
- Yoshimoto S, Okada E, Umemoto H, Tamura K, Uno Y, et al. (2008) A W-linked DM-domain gene, DM-W, participates in primary ovary development in *Xenopus laevis*. *Proc Natl Acad Sci U S A* 105: 2469–2474.
- Chen S, Zhang G, Shao C, Huang Q, Liu G, et al. (2014) Whole-genome sequence of a flatfish provides insights into ZW sex chromosome evolution and adaptation to a benthic lifestyle. *Nat Genet* 46: 253–260.
- Verhulst EC, Beukeboom LW, van de Zande L (2010) Maternal control of haplodiploid sex determination in the wasp *Nasonia*. *Science* 328: 620–623.
- Beye M, Hasselmann M, Fondrk MK, Page RE, Omholt SW (2003) The gene *csd* is the primary signal for sexual development in the honeybee and encodes an SR-type protein. *Cell* 114: 419–429.
- Hediger M, Henggeler C, Meier N, Perez R, Saccone G, et al. (2010) Molecular characterization of the key switch F provides a basis for understanding the rapid divergence of the sex-determining pathway in the housefly. *Genetics* 184: 155–170.
- Steinemann S, Steinemann M (2005) Y chromosomes: born to be destroyed. *Bioessays* 27: 1076–1083.
- Renner SS, Ricklefs RE (1995) Dioecy and its correlates in the flowering plants. *Am J Botany* 82: 596–606.
- Jarne P, Auld JR (2006) Animals mix it up too: the distribution of self-fertilization among hermaphroditic animals. *Evolution* 60: 1816–1824.
- Eppley SM, Jesson LK (2008) Moving to mate: the evolution of separate and combined sexes in multicellular organisms. *J Evol Biol* 21: 727–736.
- Ghiselin MT (1974) The Economy of Nature and the Evolution of Sex. Berkeley, CA: The University of California Press.
- Bertin RI, Newman CM (1993) Dichogamy in angiosperms. *Bot Rev* 59: 112–152.
- Munday PL, Buston PM, Warner RR (2006) Diversity and flexibility of sex-change strategies in animals. *Trends Ecol Evol* 21: 89–95.
- Charnov EL, Maynard Smith J, Bull JJ (1976) Why Be an Hermaphrodite. *Nature* 263: 125–126.
- Schärer L (2009) Tests of sex allocation theory in simultaneously hermaphroditic animals. *Evolution* 63: 1377–1405.
- De Visser JAM, Ter Maat A, Zonneveld C (1994) Energy budgets and reproductive allocation in the simultaneous hermaphrodite pond snail, *Lymnaea stagnalis*: a trade-off between male and female function. *Am Nat* 144: 861–867.
- Ashman T-L (2003) Constraints on the evolution of males and sexual dimorphism: Field estimates of genetic architecture of reproductive traits in

- three populations of gynodioecious *Fragaria virginiana*. *Evolution* 57: 2012–2025.
29. Weinzierl RP, Berthold K, Beukeboom LW, Michiels NK (1998) Reduced Male Allocation in the Parthenogenetic Hermaphrodite *Dugesia polychoa*. *Evolution* 52: 109–115.
  30. Whitton J, Sears CJ, E.J. B, Otto SP (2008) The dynamic nature of apomixis in the angiosperms. *Int J Plant Sci* 169: 169–182.
  31. Charlesworth B, Charlesworth D (1978) Model for Evolution of Dioecy and Gynodioecy. *Am Nat* 112: 975–997.
  32. Dufay M, Billard E (2012) How much better are females? The occurrence of female advantage, its proximal causes and its variation within and among gynodioecious species. *Ann Bot* 109: 505–519.
  33. Charlesworth D, Willis JH (2009) The genetics of inbreeding depression. *Nat Rev Genet* 10: 783–796.
  34. Sakai AK, Karoly K, Weller SG (1989) Inbreeding Depression in *Schiedea globosa* and *S. salicaria* (Caryophyllaceae), Subdioecious and Gynodioecious Hawaiian Species. *Am J Bot* 76: 437–444.
  35. Pannell JR (2002) The evolution and maintenance of androdioecy. *Ann Rev Ecol Evol Syst* 33: 397–425.
  36. Schaefer H, Renner SS (2010) A three-genome phylogeny of Momordica (Cucurbitaceae) suggests seven returns from dioecy to monoecy and recent long-distance dispersal to Asia. *Mol Phylogenet Evol* 54: 553–560.
  37. Valenzuela N, Lance VA (2004) Temperature Dependent Sex Determination in Vertebrates. Washington, DC: Smithsonian Books.
  38. Ming R, Bendahmane A, Renner SS (2011) Sex chromosomes in land plants. *Annu Rev Plant Biol* 62: 485–514.
  39. Dubendorfer A, Hediger M, Burghardt G, Bopp D (2002) *Musca domestica*, a window on the evolution of sex-determining mechanisms in insects. *Int J Dev Biol* 46: 75–79.
  40. Merchant-Larios H, Diaz-Hernandez V (2013) Environmental sex determination mechanisms in reptiles. *Sex Dev* 7: 95–103.
  41. Guler Y, Short S, Kile P, Ford AT (2012) Integrating field and laboratory evidence for environmental sex determination in the amphipod, *Echinogammarus marinus*. *Mar Biol* 159: 2885–2890.
  42. Walker G (2005) Sex determination in the larvae of the parasitic barnacle *Heterosaccus lunatus*: an experimental approach. *J Exp Mar Bio Ecol* 318: 31–38.
  43. Kobayashi Y, Nagahama Y, Nakamura M (2013) Diversity and plasticity of sex determination and differentiation in fishes. *Sex Dev* 7: 115–125.
  44. Warner RR, Fitch DL, Standish JD (1996) Social control of sex change in the shelf limpet, *Crepidula norrisianus*: size-specific responses to local group composition. *J Exp Mar Bio Ecol* 204: 155–167.
  45. Navarro-Martín L, Viñas J, Ribas L, Díaz N, Gutiérrez A, et al. (2011) DNA methylation of the gonadal aromatase (*cyp19a*) promoter is involved in temperature-dependent sex ratio shifts in the European sea bass. *PLoS Genet* 7: e1002447.
  46. Sarre SD, Georges A, Quinn A (2004) The ends of a continuum: genetic and temperature-dependent sex determination in reptiles. *Bioessays* 26: 639–645.
  47. Shao C, Li Q, Chen S, Zhang P, Lian J, et al. (2014) Epigenetic modification and inheritance in sexual reversal of fish. *Genome Res* 24: 604–615.
  48. Munday PL, Buston PM, Warner RR (2006) Diversity and flexibility of sex-change strategies in animals. *Trends Ecol Evol* 21: 89–95.
  49. Pen I, Uller T, Feldmeyer B, Harts A, While GM, et al. (2010) Climate-driven population divergence in sex-determining systems. *Nature* 468: 436–438.
  50. Mitchell NJ, Janzen FJ (2010) Temperature-dependent sex determination and contemporary climate change. *Sex Dev* 4: 129–140.
  51. Fisher RA (1930) *The Genetical Theory of Natural Selection*. Oxford: Oxford University Press.
  52. Kozielska M, Weissing FJ, Beukeboom LW, Pen I (2010) Segregation distortion and the evolution of sex-determining mechanisms. *Heredity* 104: 100–112.
  53. Hamilton WD (1967) Extraordinary sex ratios. *Science* 156: 477–478.
  54. Kocher TD (2004) Adaptive evolution and explosive speciation: The cichlid fish model. *Nat Rev Genet* 5: 288–298.
  55. Tao Y, Masly JP, Araripe L, Ke Y, Hartl DL (2007) A sex-ratio meiotic drive system in *Drosophila simulans*. I: an autosomal suppressor. *PLoS Biol* 5: e292.
  56. Montchamp-Moreau C (2006) Sex-ratio meiotic drive in *Drosophila simulans*: cellular mechanism, candidate genes and evolution. *Biochem Soc Trans* 34: 562–565.
  57. Presgraves DC, Severance E, Wilkinson GS (1997) Sex chromosome meiotic drive in stalk-eyed flies. *Genetics* 147: 1169–1180.
  58. Cocquet J, Ellis PJ, Mahadevaiah SK, Affara NA, Vaiman D, et al. (2012) A genetic basis for a postmeiotic X versus Y chromosome intragenomic conflict in the mouse. *PLoS Genet* 8: e1002900.
  59. Saumitou-Laprade P, Cuguen J, Vernet P (1994) Cytoplasmic male sterility in plants: molecular evidence and the nucleocytoplasmic conflict. *Trends Ecol Evol* 9: 431–435.
  60. Caruso CM, Case AL, Bailey MF (2012) The evolutionary ecology of cytonuclear interactions in angiosperms. *Trends Plant Sci* 17: 638–643.
  61. Jiggins FM, Hurst GDD, Majerus MEN (1998) Sex ratio distortion in *Acraea eumedon* (Lepidoptera: Nymphalidae) is caused by a male-killing bacterium. *Heredity* 81: 87–91.
  62. Sheeley SL, McAllister BF (2009) Mobile male-killer: similar *Wolbachia* strains kill males of divergent *Drosophila* hosts. *Heredity* (Edinb) 102: 286–292.
  63. Lande R, Seehausen O, van Alphen JJM (2001) Mechanisms of rapid sympatric speciation by sex reversal and sexual selection in cichlid fish. *Genetica* 112: 435–443.
  64. van Doorn GS, Kirkpatrick M (2007) Turnover of sex chromosomes induced by sexual conflict. *Nature* 449: 909–912.
  65. van Doorn GS, Kirkpatrick M (2010) Transitions between male and female heterogamety caused by sex-antagonistic selection. *Genetics* 186: 629–645.
  66. Otto SP, Pannell JR, Peichel CL, Ashman T-L, Charlesworth D, et al. (2011) About PAR: The distinct evolutionary dynamics of the pseudoautosomal region. *Trends Genet* 27: 358–367.
  67. Bachtrog D (2008) The temporal dynamics of processes underlying Y chromosome degeneration. *Genetics* 179: 1513–1525.
  68. Blaser O, Gossen C, Neuenschwander S, Perrin N (2013) Sex-chromosome turnovers induced by deleterious mutation load. *Evolution* 67: 635–645.
  69. Anderson JL, Rodriguez Mari A, Braasch I, Amores A, Hohenlohe P, et al. (2012) Multiple sex-associated regions and a putative sex chromosome in zebrafish revealed by RAD mapping and population genomics. *PLoS ONE* 7: e40701.
  70. Bradley KM, Breyer JP, Melville DB, Broman KW, Knapik EW, et al. (2011) An SNP-Based Linkage Map for Zebrafish Reveals Sex Determination Loci. *G3* (Bethesda) 1: 3–9.
  71. Liew WC, Bartfai R, Lim Z, Sreenivasan R, Siegfried KR, et al. (2011) Polygenic sex determination system in zebrafish. *PLoS ONE* 7: e34397.
  72. Siegfried KR, Nüsslein-Volhard C (2008) Germ line control of female sex determination in zebrafish. *Dev Bio* 324: 277–287.
  73. Parnell NF, Streebman JT (2013) Genetic interactions controlling sex and color establish the potential for sexual conflict in Lake Malawi cichlid fishes. *Heredity* (Edinb) 110: 239–246.
  74. Ser JR, Roberts RB, Kocher TD (2010) Multiple interacting loci control sex determination in lake Malawi cichlid fish. *Evolution* 64: 486–501.
  75. Vandeputte M, Dupont-Nivet M, Chavanne H, Chatain B (2007) A polygenic hypothesis for sex determination in the European sea bass *Dicentrarchus labrax*. *Genetics* 176: 1049–1057.
  76. Yusa Y (2007) Nuclear sex-determining genes cause large sex-ratio variation in the apple snail *Pomacea canaliculata*. *Genetics* 175: 179–184.
  77. Foster JW, Graves JAM (1994) An Sry-Related Sequence on the Marsupial X-Chromosome - Implications for the Evolution of the Mammalian Testis determining Gene. *Proc Natl Acad Sci U S A* 91: 1927–1931.
  78. Matsuda M, Nagahama Y, Shinomiya A, Sato T, Matsuda C, et al. (2002) DMY is a Y-specific DM-domain gene required for male development in the medaka fish. *Nature* 417: 559–563.
  79. Nanda I, Kondo M, Hornung U, Asakawa S, Winkler C, et al. (2002) A duplicated copy of DMRT1 in the sex-determining region of the Y chromosome of the medaka, *Oryzias latipes*. *Proc Natl Acad Sci U S A* 99: 11778–11783.
  80. Myosho T, Otake H, Masuyama H, Matsuda M, Kuroki Y, et al. (2012) Tracing the Emergence of a Novel Sex-Determining Gene in Medaka, *Oryzias luzonensis*. *Genetics* 191: 163–170.
  81. Graves JAM, Peichel CL (2010) Are homologies in vertebrate sex determination due to shared ancestry or to limited options? *Genome Biol* 11: 205.
  82. Yano A, Guyomard R, Nicol B, Jouanno E, Quillet E, et al. (2012) An Immune-Related Gene Evolved into the Master Sex-Determining Gene in Rainbow Trout, *Oncorhynchus mykiss*. *Curr Biol* 22: 1–6.
  83. Maine EM, Salz HK, Cline TW, Schedl P (1985) The Sex-lethal gene of *Drosophila*: DNA alterations associated with sex-specific lethal mutations. *Cell* 43: 521–529.
  84. Cline TW, Dorsett M, Sun S, Harrison MM, Dines J, et al. (2010) Evolution of the *Drosophila* feminizing switch gene Sex-lethal. *Genetics* 186: 1321–1336.
  85. Kiuchi T, Koga H, Kawamoto M, Shoji K, Sakai H, et al. (2014) A single female-specific piRNA is the primary determinant of sex in the silkworm. *Nature* 509: 633–636.
  86. Diggle PK, Di Stilio VS, Gschwend AR, Golenberg EM, Moore RC, et al. (2011) Multiple developmental processes underlie sex differentiation in angiosperms. *Trends Genet* 27: 368–376.
  87. Martin A, Troadec C, Boualem A, Rajab M, Fernandez R, et al. (2009) A transposon-induced epigenetic change leads to sex determination in melon. *Nature* 461: 1135–U1237.
  88. Cui X, Wang Q, Yin W, Xu H, Wilson ZA, et al. (2012) PMRD: a curated database for genes and mutants involved in plant male reproduction. *BMC Plant Biol* 12: 215.
  89. Haag ES, Doty AV (2005) Sex determination across evolution: connecting the dots. *PLoS Biol* 3: e21.
  90. Kopp A (2012) Dmrt genes in the development and evolution of sexual dimorphism. *Trends Genet* 28: 175–184.
  91. Raymond CS, Murphy MW, O'Sullivan MG, Bardwell VJ, Zarkower D (2000) *Dmrt1*, a gene related to worm and fly sexual regulators, is required for mammalian testis differentiation. *Genes Dev* 14: 2587–2595.
  92. Chue J, Smith CA (2011) Sex determination and sexual differentiation in the avian model. *FEBS J* 278: 1027–1034.

93. Sinclair A, Smith C, Western P, McClive P (2002) A comparative analysis of vertebrate sex determination. *Novartis Found Symp* 244: 102–111; discussion 111–104, 203–106, 253–107.
94. Shoemaker C, Ramsey M, Queen J, Crews D (2007) Expression of *Sox9*, *Mis*, and *Dmrt1* in the gonad of a species with temperature-dependent sex determination. *Dev Dyn* 236: 1055–1063.
95. Valenzuela N (2010) Multivariate expression analysis of the gene network underlying sexual development in turtle embryos with temperature-dependent and genotypic sex determination. *Sex Dev* 4: 39–49.
96. Yoshimoto S, Ito M (2011) A ZZ/ZW-type sex determination in *Xenopus laevis*. *FEBS J* 278: 1020–1026.
97. Herpin A, Schartl M (2011) *Dmrt1* genes at the crossroads: a widespread and central class of sexual development factors in fish. *FEBS J* 278: 1010–1019.
98. Hempel LU, Oliver B (2007) Sex-specific Doublesex<sup>3d</sup> expression in subsets of *Drosophila* somatic gonad cells. *BMC Dev Biol* 7: 113.
99. Cho S, Huang ZY, Zhang J (2007) Sex-specific splicing of the honeybee doublesex gene reveals 300 million years of evolution at the bottom of the insect sex-determination pathway. *Genetics* 177: 1733–1741.
100. Kato Y, Kobayashi K, Watanabe H, Iguchi T (2011) Environmental sex determination in the branchiopod crustacean *Daphnia magna*: deep conservation of a *Doublesex* gene in the sex-determining pathway. *PLoS Genet* 7: e1001345.
101. Zhang EF, Qiu GF (2010) A novel *Dmrt* gene is specifically expressed in the testis of Chinese mitten crab, *Eriocheir sinensis*. *Dev Genes Evol* 220: 151–159.
102. Klimbunga S, Amparyup P, Khamnamtong B, Hirono I, Aoki T, et al. (2009) Isolation and characterization of testis-specific *DMRT1* in the tropical abalone (*Haliotis asinina*). *Biochem Genet* 47: 66–79.
103. Naimi A, Martinez AS, Specq ML, Mrac A, Diss B, et al. (2009) Identification and expression of a factor of the DM family in the oyster *Crassostrea gigas*. *Comp Biochem Physiol A Mol Integr Physiol* 152: 189–196.
104. Verhulst EC, van de Zande L, Beukeboom LW (2010) Insect sex determination: it all evolves around transformer. *Curr Opin Genet Dev* 20: 376–383.
105. Valenzuela N, Neuwald JL, Liteman R (2013) Transcriptional evolution underlying vertebrate sexual development. *Developmental Dynamics* 242: 307–319.
106. Robinett CC, Vaughan AG, Knapp JM, Baker BS (2010) Sex and the single cell. II. There is a time and place for sex. *PLoS Biol* 8: e1000365.
107. Zhao D, McBride D, Nandi S, McQueen HA, McGrew MJ, et al. (2010) Somatic sex identity is cell autonomous in the chicken. *Nature* 464: 237–242.
108. Wolff JR, Zarkower D (2008) Somatic sexual differentiation in *Caenorhabditis elegans*. *Curr Top Dev Biol* 83: 1–39.
109. Ellis RE (2008) Sex determination in the *Caenorhabditis elegans* germ line. *Curr Top Dev Biol* 83: 41–64.
110. Steinmann-Zwicky M (1992) How do germ cells choose their sex? *Drosophila* as a paradigm. *Bioessays* 14: 513–518.
111. Murray SM, Yang SY, Van Doren M (2010) Germ cell sex determination: a collaboration between soma and germline. *Curr Opin Cell Biol* 22: 722–729.
112. Hilfiker-Kleiner D, Dübendorfer A, Hilfiker A, Nöthiger R (1994) Genetic control of sex determination in the germ line and soma of the housefly, *Musca domestica*. *Development* 120: 2531–2538.
113. Blackler AW (1965) Germ-cell transfer and sex ratio in *Xenopus laevis*. *J Embryol Exp Morphol* 13: 51–61.
114. Yoshizaki G, Ichikawa M, Hayashi M, Iwasaki Y, Miwa M, et al. (2010) Sexual plasticity of ovarian germ cells in rainbow trout. *Development* 137: 1227–1230.
115. Okutsu T, Suzuki K, Takeuchi Y, Takeuchi T, Yoshizaki G (2006) Testicular germ cells can colonize sexually undifferentiated embryonic gonad and produce functional eggs in fish. *Proc Natl Acad Sci U S A* 103: 2725–2729.
116. Shinomiya A, Shibata N, Sakaizumi M, Hamaguchi S (2002) Sex reversal of genetic females (XX) induced by the transplantation of XY somatic cells in the medaka, *Oryzias latipes*. *Int J Dev Biol* 46: 711–717.
117. Steinmann-Zwicky M, Schmid H, Nothiger R (1989) Cell-autonomous and inductive signals can determine the sex of the germ line of *Drosophila* by regulating the gene *Sxl*. *Cell* 57: 157–166.
118. Durcova-Hills G, Capel B (2008) Development of germ cells in the mouse. *Curr Top Dev Biol* 83: 185–212.
119. Skaletsky H, Kuroda-Kawaguchi T, Minx P, Cordum H, Hillier L, et al. (2003) The male-specific region of the human Y chromosome is a mosaic of discrete sequence classes. *Nature* 423: 825–837.
120. Nanda I, Schlegelmilch K, Haaf Y, Schartl M, Schmid M (2008) Synteny conservation of the Z chromosome in 14 avian species (11 families) supports a role for Z dosage in avian sex determination. *Cytogenet Genome Res* 122: 150–156.
121. Vicoso B, Emerson JJ, Zektser Y, Mahajan S, Bachtrog D (2013) Comparative sex chromosome genomics in snakes: differentiation, evolutionary strata, and lack of global dosage compensation. *PLoS Biol* 11: e1001643.
122. Carvalho AB, Koerich LB, Clark AG (2009) Origin and evolution of Y chromosomes: *Drosophila* tales. *Trends Genet* 25: 270–277.
123. White MJD (1973) *Animal Cytology and Evolution*: Cambridge University Press.
124. Navajas-Perez R, Schwarzacher T, Rejon MR, Garrido-Ramos MA (2009) Molecular cytogenetic characterization of *Rumex papillaris*, a dioecious plant with an XX/XY(1)Y(2) sex chromosome system. *Genetica* 135: 87–93.
125. Ohno S (1967) *Sex chromosomes and sex linked genes*. Berlin: Springer Verlag.
126. Vicoso B, Kaiser VB, Bachtrog D (2013) Sex-biased gene expression at homomorphic sex chromosomes in emus and its implication for sex chromosome evolution. *Proc Natl Acad Sci U S A* 110: 6453–6458.
127. Perrin N (2009) Sex Reversal: A Fountain of Youth for Sex Chromosomes? *Evolution* 63: 3043–3049.
128. Stöck M, Horn A, Grossen C, Lindtke D, Sermier R, et al. (2011) Ever-young sex chromosomes in European tree frogs. *PLoS Biol* 9: e1001062.
129. Rice WR (1987) The Accumulation of Sexually Antagonistic Genes as a Selective Agent Promoting the Evolution of Reduced Recombination between Primitive Sex-Chromosomes. *Evolution* 41: 911–914.
130. Jordan CY, Charlesworth D (2012) The potential for sexually antagonistic polymorphism in different genome regions. *Evolution* 66: 505–516.
131. Graves J (2004) The degenerate Y chromosome - can conversion save it? *Reprod Fertil Dev* 16: 527–534.
132. Aitken R, Marshall Graves J (2002) The future of sex. *Nature* 415: 963.
133. Graves J (2006) Sex chromosome specialization and degeneration in mammals. *Cell* 124: 901–914.
134. Engelstaedter J (2008) Muller's Ratchet and the Degeneration of Y Chromosomes: A Simulation Study. *Genetics* 180: 957–967.
135. Hughes JF, Skaletsky H, Brown LG, Pyntikova T, Graves T, et al. (2012) Strict evolutionary conservation followed rapid gene loss on human and rhesus Y chromosomes. *Nature* 483: 82–86.
136. Vicoso B, Bachtrog D (2013) Reversal of an ancient sex chromosome to an autosome in *Drosophila*. *Nature*: 499: 332–335.
137. Koerich LB, Wang X, Clark AG, Carvalho AB (2008) Low conservation of gene content in the *Drosophila* Y chromosome. *Nature* 456: 949–951.
138. Hodgkin J (2002) Exploring the envelope: Systematic alteration in the sex-determination system of the nematode *Caenorhabditis elegans*. *Genetics* 162: 767–780.
139. Pokorná M, Kratochvil L (2009) Phylogeny of sex-determining mechanisms in squamate reptiles: are sex chromosomes an evolutionary trap? *Zoological Journal of the Linnean Society* 156: 168–183.
140. Jimenez R, Barrionuevo FJ, Burgos M (2013) Natural exceptions to normal gonad development in mammals. *Sex Dev* 7: 147–162.
141. Castillo ER, Martí DA, Bidau CJ (2010) Sex- and neo-sex chromosomes in Orthoptera: a review. *J Orthoptera Research* 19: 213–231.
142. Bush GL (1966) Female Heterogamety in the Family Tephritidae. *Am Nat* 100: 119–126.
143. Ullerich FH (1963) Geschlechtschromosomen und Geschlechtsbestimmung bei einigen Calliphorinen (Calliphoridae, Diptera). *Chromosoma* 14: 45–110.
144. Malik SB, Fichtling AW, Stefaniak LM, Schurko AM, Logsdon JM (2008) An Expanded Inventory of Conserved Meiotic Genes Provides Evidence for Sex in *Trichomonas vaginalis*. *PLoS ONE* 3: e2879.
145. Lahr DJ, Parfrey LW, Mitchell EA, Katz LA, Lara E (2011) The chastity of amoebae: re-evaluating evidence for sex in amoeboid organisms. *Proc R Soc Lond B Biol Sci* 278: 2081–2090.
146. Haag ES (2007) Why two sexes? Sex determination in multicellular organisms and protistan mating types. *Semin Cell Dev Biol* 18: 348–349.
147. Perrin N (2012) What uses are mating types? The “developmental switch” model. *Evolution* 66: 947–956.
148. Dacks J, Kasinsky H (1999) Nuclear condensation in protozoan gametes and the evolution of anisogamy. *Comp Biochem Physiol* 124: 287–295.
149. Bulmer MG, Parker GA (2002) The evolution of anisogamy: a game-theoretic approach. *Proc Biol Sci* 269: 2381–2388.
150. Parker GA, Baker RR, Smith VG (1972) The origin and evolution of gamete dimorphism and the male-female phenomenon. *J Theor Biol* 36: 529–553.
151. Ferris P, Olson BJ, De Hoff PL, Douglass S, Casero D, et al. (2010) Evolution of an expanded sex-determining locus in *Volvox*. *Science* 328: 351–354.
152. Bachtrog D, Kirkpatrick M, Mank JE, McDaniel SF, Pires JC, et al. (2011) Are all sex chromosomes created equal? *Trends Genet* 27: 350–357.
153. Sleigh MA (1991) *Protozoa and other protists*. Cambridge: University of Cambridge.
154. Davidovich NA, Kaczmarek I, Ehrman JM (2010) Heterothallic and homothallic sexual reproduction in *Tabularia fasciculata* (Bacillariophyta). *Fottea* 10: 251–266.
155. Hattori RS, Murai Y, Oura M, Masuda S, Majhi SK, et al. (2012) A Y-linked anti-Mullerian hormone duplication takes over a critical role in sex determination. *Proc Natl Acad Sci U S A* 109: 2955–2959.
156. Kamiya T, Kai W, Tasumi S, Oka A, Matsunaga T, et al. (2012) A Trans-Species Missense SNP in *Amhr2* Is Associated with Sex Determination in the Tiger Pufferfish, *Takifugu rubripes* (Fugu). *PLoS Genet* 8: e1002798.

## Klinefelter Syndrome (KS)

KS describes a set of physical, language, and social development symptoms in males who have an extra X chromosome. Its main feature is infertility. Outward signs of KS can be subtle, so symptoms often are not recognized, and may not be treated in a timely manner. The NICHD is one of many federal agencies and NIH Institutes working to understand KS, discover why it occurs, and identify and treat its symptoms.

### Common Name

- Klinefelter syndrome

### Medical or Scientific Names

- Klinefelter syndrome
- 47,XXY
- XXY syndrome or condition
- XXY trisomy
- 47,XXY/46,XY or mosaic syndrome (rare variation)
- Poly-X Klinefelter syndrome, including the following rare variations:
  - 48,XXYY (or tetrasomy)
  - 48,XXXY (or tetrasomy)
  - 49,XXXXY (or pentasomy)

## Klinefelter Syndrome (KS): Condition Information

### What is KS?

The term "Klinefelter (pronounced *KLAHYN-fel-ter*) syndrome," or KS, describes a set of features that can occur in a male who is born with an extra X chromosome (pronounced *KROH-muh-sohm*) in his cells. It is named after Dr. Henry Klinefelter, who identified the condition in the 1940s.<sup>1</sup>([/health/topics/klinefelter/conditioninfo/Pages/Default.aspx#f1](https://www.nichd.nih.gov/health/topics/klinefelter/conditioninfo/Pages/Default.aspx#f1)).

Usually, every cell in a male's body, except sperm and red blood cells, contains 46 chromosomes. The 45th and 46th chromosomes—the X and Y chromosomes—are sometimes called "sex chromosomes" because they determine a person's sex. Normally, males have one X and one Y chromosome, making them XY. Males with KS have an extra X chromosome, making them XXY.

KS is sometimes called "47,XXY" (47 refers to total chromosomes) or the "XXY condition." Those with KS are sometimes called "XXY males."

Some males with KS may have both XY cells and XXY cells in their bodies. This is called "mosaic" (*mo-ZAY-ik*). Mosaic males may have fewer symptoms of KS depending on the number of XY cells they have in their bodies and where these cells are located. For example, males who have normal XY cells in their testes may be fertile. <sup>2</sup>  
(</health/topics/klinefelter/conditioninfo/Pages/Default.aspx#f2>).

In very rare cases, males might have two or more extra X chromosomes in their cells, for instance XXXY or XXXXY, or an extra Y, such as XXYY. This is called poly-X Klinefelter syndrome, and it causes more severe symptoms.<sup>1</sup>  
(</health/topics/klinefelter/conditioninfo/Pages/Default.aspx#f1>).

## Citations

1. Klinefelter, H.F., Reifenstein, E.C., & Albright, F. (1942). Syndrome characterized by gynecomastia aspermatogenesis without A-Leydigism and increased excretion of follicle stimulating hormone. *Journal of Clinical Endocrinology & Metabolism*, 2, 615–627.
2. Bojesen, A., Juul, S., & Gravholt, C.H. (2003). Prenatal and postnatal prevalence of Klinefelter syndrome: A national registry study. *Journal of Clinical Endocrinology & Metabolism*, 88(2), 622–626.

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[What causes it? \(/health/topics/klinefelter/conditioninfo/Pages/causes.aspx\)](/health/topics/klinefelter/conditioninfo/Pages/causes.aspx) »

## What causes Klinefelter syndrome (KS)?

The extra chromosome results from a random error that occurs when a sperm or egg is formed; this error causes an extra X cell to be included each time the cell divides to form new cells. In very rare cases, more than one extra X or an extra Y is included.

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« [Condition Information \(/health/topics/klinefelter/conditioninfo/Pages/Default.aspx\)](/health/topics/klinefelter/conditioninfo/Pages/Default.aspx).

[How many people are affected/at risk?](#)

[\(/health/topics/klinefelter/conditioninfo/Pages/risk.aspx\)](/health/topics/klinefelter/conditioninfo/Pages/risk.aspx) »

# How many people are affected by or at risk for Klinefelter syndrome (KS)?

Researchers estimate that 1 male in about 500 newborn males has an extra X chromosome, making KS among the most common chromosomal disorders seen in all newborns.<sup>1</sup> (</health/topics/klinefelter/conditioninfo/Pages/risk.aspx#f1>). The likelihood of a third or fourth X is much rarer: [2./health/topics/klinefelter/conditioninfo/Pages/risk.aspx#f2](/health/topics/klinefelter/conditioninfo/Pages/risk.aspx#f2).

## Prevalence of Klinefelter syndrome variants

Number of extra X chromosomes	One (XXY)	Two (XXXY)	Three (XXXXY)
Number of newborn males with the condition	1 in 500	1 in 50,000	1 in 85,000 to 100,000

Scientists are not sure what factors increase the risk of KS. The error that produces the extra chromosome occurs at random, meaning the error is not hereditary (pronounced *huh-RED-i-ter-ee*) or passed down from parent to child. Research suggests that older mothers might be slightly more likely to have a son with KS. However, the extra X chromosome in KS comes from the father about one-half of the time.<sup>3</sup> (</health/topics/klinefelter/conditioninfo/Pages/risk.aspx#f3>).

## Citations

1. Nielsen, J., & Wohlert, M. (1991). Chromosome abnormalities found among 34,910 newborn children: Results from a 13-year incidence study in Aarhus, Denmark. *Human Genetics*, 87(1), 81–83.
2. Klinefelter, H.F., Reifenstein, E.C., & Albright, F. (1942). Syndrome characterized by gynecomastia aspermatogenesis without A-Leydigism and increased excretion of follicle stimulating hormone. *Journal of Clinical Endocrinology & Metabolism*, 2, 615–627.
3. National Human Genome Research Institute. Learning about Klinefelter Syndrome. Retrieved on June 5, 2012 from <http://www.genome.gov/19519068>

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« [What causes it? \(/health/topics/klinefelter/conditioninfo/Pages/causes.aspx\)](/health/topics/klinefelter/conditioninfo/Pages/causes.aspx)

[What are common symptoms?](#)

[\(/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx\)](/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx) »

## What are common symptoms of Klinefelter syndrome (KS)?

Because XXY males do not really appear different from other males and because they may not have any or have mild symptoms, XXY males often don't know they have KS.<sup>1</sup>

[\(/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f1\)](/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f1),<sup>2</sup>

[\(/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f2\)](/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f2).

In other cases, males with KS may have mild or severe symptoms. Whether or not a male with KS has visible symptoms depends on many factors, including how much testosterone his body makes, if he is mosaic (with both XY and XXY cells), and his age when the condition is diagnosed and treated.

KS symptoms fall into these main categories:

- Physical Symptoms  
[\(/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#physical\)](/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#physical)
- Language and Learning Symptoms  
[\(/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#language\)](/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#language)
- Social and Behavioral Symptoms  
[\(/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#social\)](/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#social)
- Symptoms of Poly-X KS  
[\(/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#polyx\)](/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#polyx)

### Physical Symptoms

Many physical symptoms of KS result from low testosterone levels in the body. The degree of symptoms differs based on the amount of testosterone needed for a specific age or developmental stage and the amount of testosterone the body makes or has available.

During the first few years of life, when the need for testosterone is low, most XXY males do not show any obvious differences from typical male infants and young boys. Some may have slightly weaker muscles, meaning they might sit up, crawl, and walk slightly later than average. For example, on average, baby boys with KS do not start walking until age 18 months.<sup>3</sup>[\(/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f3\)](/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f3)

After age 5 years, when compared to typically developing boys, boys with KS may be slightly:

- Taller
- Fatter around the belly
- Clumsier
- Slower in developing motor skills, coordination, speed, and muscle strength

Puberty for boys with KS usually starts normally. But because their bodies make less testosterone than non-KS boys, their pubertal development may be disrupted or slow. In addition to being tall, KS boys may have:

- Smaller testes and penis
- Breast growth (about one-third of teens with KS have breast growth)
- Less facial and body hair
- Reduced muscle tone
- Narrower shoulders and wider hips
- Weaker bones, greater risk for bone fractures
- Decreased sexual interest
- Lower energy
- Reduced sperm production

An adult male with KS may have these features:

- Infertility: Nearly all men with KS are unable to father a biologically-related child without help from a fertility specialist.<sup>4</sup>[\(/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f4\)](/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f4)
- Small testes, with the possibility of testes shrinking slightly after the teen years<sup>5</sup>[\(/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f5\)](/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f5)
- Lower testosterone levels, which lead to less muscle, hair, and sexual interest and function
- Breasts or breast growth (called gynecomastia, pronounced *GUY-nuh-kow-mast-ee-uh*).

In some cases, breast growth can be permanent, and about 10% of XXY males need breast-reduction surgery.<sup>6</sup>[\(/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f6\)](/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f6)

## Language and Learning Symptoms

Most males with KS have normal intelligence quotients (IQs)<sup>7</sup>

[\(/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f7\)](/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f7),<sup>8</sup>

[\(/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f8\)](/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f8) and successfully complete education at all levels. (IQ is a frequently used intelligence measure, but does not include emotional, creative, or other types of intelligence.) Between 25% and 85% of all males with KS have some kind of learning or language-related problem, which makes it more likely that they will need some extra help in school. Without this help or intervention, KS males might fall behind their classmates as schoolwork becomes harder.

KS males may experience some of the following learning and language-related challenges:<sup>9</sup>

[\(/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f9\)](/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f9)

- **A delay in learning to talk.** Infants with KS tend to make only a few different vocal sounds. As they grow older, they may have difficulty saying words clearly. It might be hard for

them to distinguish differences between similar sounds.

- **Trouble using language to express their thoughts and needs.** Boys with KS might have problems putting their thoughts, ideas, and emotions into words. Some may find it hard to learn and remember some words, such as the names of common objects.
- **Trouble processing what they hear.** Although most boys with KS can understand what is being said to them, they might take longer to process multiple or complex sentences. In some cases, they might fidget or "tune out" because they take longer to process the information. It might also be difficult for KS males to concentrate in noisy settings. They might also be less able to understand a speaker's feelings from just speech alone.
- **Reading difficulties.** Many boys with KS have difficulty understanding what they read (called poor reading comprehension). They might also read more slowly than other boys.

By adulthood, most males with KS learn to speak and converse normally, although they may have a harder time doing work that involves extensive reading and writing.

## Social and Behavioral Symptoms

Many of the social and behavioral symptoms in KS may result from the language and learning difficulties. For instance, boys with KS who have language difficulties might hold back socially and could use help building social relationships.

Boys with KS, compared to typically developing boys, tend to be:

- Quieter
- Less assertive or self-confident
- More anxious or restless
- Less physically active
- More helpful and eager to please
- More obedient or more ready to follow directions

In the teenage years, boys with KS may feel their differences more strongly. As a result, these teen boys are at higher risk of depression, substance abuse, and behavioral disorders. Some teens might withdraw, feel sad, or act out their frustration and anger.

As adults, most men with KS have lives similar to those of men without KS. They successfully complete high school, college, and other levels of education. They have successful and meaningful careers and professions. They have friends and families.

Contrary to research findings published several decades ago, males with KS are no more likely to have serious psychiatric disorders or to get into trouble with the law.<sup>10</sup>

([/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f10](http://health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f10)).

## Symptoms of Poly-X KS<sup>11</sup>

[\(/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f11\)](/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f11)

Males with poly-X Klinefelter syndrome have more than one extra X chromosome, so their symptoms might be more pronounced than in males with KS. In childhood, they may also have seizures, crossed eyes, constipation, and recurrent ear infections. Poly-KS males might also show slight differences in other physical features.

Some common additional symptoms for several poly-X Klinefelter syndromes are listed below.

### 48,XXYY

- Long legs
- Little body hair
- Lower IQ, average of 60 to 80 (normal IQ is 90 to 110)
- Leg ulcers and other vascular disease symptoms
- Extreme shyness, but also sometimes aggression and impulsiveness

### 48,XXXY (or tetrasomy)

- Eyes set further apart
- Flat nose bridge
- Arm bones connected to each other in an unusual way
- Short
- Fifth (smallest) fingers curve inward (clinodactyly, pronounced *KLAHY-noh-dak-tl-ee*)
- Lower IQ, average 40 to 60
- Immature behavior

### 49,XXXXY (or pentasomy)

- Low IQ, usually between 20 and 60
- Small head
- Short
- Upward-slanted eyes
- Heart defects, such as when the chambers do not form properly<sup>12</sup>  
[\(/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f12\)](/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx#f12)
- High feet arches
- Shy, but friendly
- Difficulty with changing routines

## Citations

1. Abramsky, L., & Chapple, J. (1997). 47, XXY (Klinefelter syndrome) and 47,XYY: Estimated rates of and indication for postnatal diagnosis with implications for prenatal counselling. *Prenatal Diagnosis*, 17(4), 363-368.  
[Case 3:21-cv-00490 Document 39-12 Filed 03/02/22 Page 8 of 21 PageID #: 955](#)

368.

2. Visootsak, J., Aylstock, M., & Graham, J.M. Jr. (2001). Klinefelter syndrome and its variants: An update and review for the primary pediatrician. *Clinical Pediatrics (Phila)*, 40(12), 639-651.
3. Simpson, J.L., de la Cruz, F., Swerdloff, R.S., Samanga-Sprouse, C., Skakkebaek, N.E., Graham, J.M. Jr., et al. (2003). Klinefelter syndrome: Expanding the phenotype and identifying new research directions. *Genetics in Medicine*, 5(6), 460-468.
4. Plotton, I., Brosse A., & Lejeune, H. (2010). Is it useful to modify the care of Klinefelter's syndrome to improve the chances of paternity? *Annales d'endocrinologie (Paris)*, 71(6), 494-504. French.
5. Smyth, C.M., & Brenner, W.J. (1998). Klinefelter syndrome. *Archives of Internal Medicine*, 158(12), 1309-1314.
6. Bock, R. (1993). Understanding Klinefelter syndrome: A guide for XXY males and their families (Adolescence section). NIH Pub. No. 93-3202. Office of Research Reporting. Retrieved June 5, 2012 from NICHD.
7. Geschwind, D.H., & Dykens, E. (2004). Neurobehavioral and psychosocial issues in Klinefelter syndrome. *Learning Disabilities Research & Practice*, 19(3), 166-173.
8. Linden, M.G., Bender, B.G., & Robinson, A. (2002). Genetic counseling for sex chromosome abnormalities. *American Journal of Medical Genetics*, 110(1), 3-10.
9. Visootsak, J., & Graham, J.M. Jr. (2009). Social function in multiple X and Y chromosome disorders: XXY, XYY, XXYY, and XXXY. *Developmental Disabilities Research Reviews*, 15(4), 328-332.
10. Ratcliffe, S. (1999). Long-term outcome in children of sex chromosome abnormalities. *Archives of Diseases in Children*, 80(2), 192-195.
11. Linden, M.G., Bender, B.G., & Robinson, A. (1995). Sex chromosome tetrasomy and pentasomy. *Pediatrics*, 96(4 Pt 1), 672-682.
12. Kassai, R., Hamada, I., Furuta, H., Cho, K., Abe, K., Deng, H. X., & Niikawa, N. (1991). Penta X syndrome: A case report with review of the literature. *American Journal of Medical Genetics*, 40(1), 51-56.

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« [How many people are affected/at risk?](#)

[\(/health/topics/klinefelter/conditioninfo/Pages/risk.aspx\)](#)

[What are the treatments? \(/health/topics/klinefelter/conditioninfo/Pages/treatments.aspx\)](#) »

# What are the treatments for symptoms in Klinefelter syndrome (KS)?

It's important to remember that because symptoms can be mild, many males with KS are never diagnosed or treated.<sup>1</sup>(</health/topics/klinefelter/conditioninfo/Pages/treatments.aspx#f1>).

The earlier in life that KS symptoms are recognized and treated, the more likely it is that the symptoms can be reduced or eliminated.<sup>2</sup>

(</health/topics/klinefelter/conditioninfo/Pages/treatments.aspx#f2>) It is especially helpful to begin treatment by early puberty. Puberty is a time of rapid physical and psychological change, and treatment can successfully limit symptoms. However, treatment can bring benefits at any age.

The type of treatment needed depends on the type of symptoms being treated.

## Treating Physical Symptoms

(</health/topics/klinefelter/conditioninfo/Pages/treatments.aspx#physical>).

## Treating Language and Learning Symptoms

(</health/topics/klinefelter/conditioninfo/Pages/treatments.aspx#language>).

## Treating Social and Behavioral Symptoms

(</health/topics/klinefelter/conditioninfo/Pages/treatments.aspx#social>).

## Treating Physical Symptoms

### Treatment for Low Testosterone<sup>3</sup>

(</health/topics/klinefelter/conditioninfo/Pages/treatments.aspx#f3>)

About one-half of XXY males' chromosomes have low testosterone levels.<sup>4</sup>

(</health/topics/klinefelter/conditioninfo/Pages/treatments.aspx#f4>) These levels can be raised by taking supplemental testosterone. Testosterone treatment can:

- Improve muscle mass
- Deepen the voice
- Promote growth of facial and body hair
- Help the reproductive organs to mature
- Build and maintain bone strength and help prevent osteoporosis in later years
- Produce a more masculine appearance, which can also help relieve anxiety and depression
- Increase focus and attention

There are various ways to take testosterone:

- Injections or shots, every 2 to 3 weeks

- Pills
- Through the skin, also called transdermal (pronounced *tranz-DEEM-ul*); current methods include wearing a testosterone patch or rubbing testosterone gel on the skin

Males taking testosterone treatment should work closely with an endocrinologist (pronounced *en-doe-kren-AWL-oh-jist*), a doctor who specializes in hormones and their functions, to ensure the best outcome from testosterone therapy. For information on how to find an endocrinologist, see [the Resources and Publications section \(/health/topics/klinefelter/resources/Pages/patients.aspx\)](/health/topics/klinefelter/resources/Pages/patients.aspx).

*Is testosterone therapy right for every XXY male?*

Not all males with XXY condition benefit from testosterone therapy.

For males whose testosterone level is low to normal, the benefits of taking testosterone are less clear than for when testosterone is very low. Side effects, although generally mild, can include acne, skin rashes from patches or gels, breathing problems (especially during sleep), and higher risk of an enlarged prostate gland or prostate cancer in older age. In addition, testosterone supplementation will not increase testicular size, decrease breast growth, or correct infertility.

Although the majority of boys with KS grow up to live as males, some develop atypical gender identities. For these males, supplemental testosterone may not be suitable. Gender identity should be discussed with health care specialists before starting treatment.<sup>5</sup>  
[\(/health/topics/klinefelter/conditioninfo/Pages/treatments.aspx#f5\)](/health/topics/klinefelter/conditioninfo/Pages/treatments.aspx#f5)

## Treatment for Enlarged Breasts

No approved drug treatment exists for this condition of over-developed breast tissue, termed gynecomastia. Some health care providers recommend surgery—called mastectomy (pronounced *ma-STEK-tuh-mee*)—to remove or reduce the breasts of XXY males.

When adult men have breasts, they are at higher risk for breast cancer than other men and need to be checked for this condition regularly. The mastectomy lowers the risk of cancer and can reduce the social stress associated with XXY males having enlarged breasts.

Because it is a surgical procedure, mastectomy carries a variety of risks. XXY males who are thinking about mastectomy should discuss all the risks and benefits with their health care provider.

## Treatment for Infertility

Between 95% and 99% of XXY men are infertile because they do not produce enough sperm to fertilize an egg naturally. But, sperm are found in more than 50% of men with KS.<sup>6</sup>  
[\(/health/topics/klinefelter/conditioninfo/Pages/treatments.aspx#f6\)](/health/topics/klinefelter/conditioninfo/Pages/treatments.aspx#f6).

Advances in assistive reproductive technology (ART) have made it possible for some men with KS to conceive. One type of ART, called testicular sperm extraction with intracytoplasmic (pronounced *in-trah-sigh-toe-PLAZ-mick*) sperm injection (TESE-ICSI), has shown success for XXY males. For this procedure, a surgeon removes sperm from the testes and places one sperm into an egg.

Like all ART, TESE-ICSI carries both risks and benefits. For instance, it is possible that the resulting child might have the XXY condition. In addition, the procedure is expensive and is often not covered by health insurance plans. Importantly, there is no guarantee the procedure will work.

Recent studies suggest that collecting sperm from adolescent XXY males and freezing the sperm until later might result in more pregnancies during subsequent fertility treatments.<sup>7</sup>  
[\(/health/topics/klinefelter/conditioninfo/Pages/treatments.aspx#f7\)](/health/topics/klinefelter/conditioninfo/Pages/treatments.aspx#f7),<sup>8</sup>  
[\(/health/topics/klinefelter/conditioninfo/Pages/treatments.aspx#f8\)](/health/topics/klinefelter/conditioninfo/Pages/treatments.aspx#f8). This is because although XXY males may make some healthy sperm during puberty, this becomes more difficult as they leave adolescence and enter adulthood.

## Treating Language and Learning Symptoms

Some, but not all, children with KS have language development and learning delays. They might be slow to learn to talk, read, and write, and they might have difficulty processing what they hear. But various interventions, such as speech therapy and educational assistance, can help to reduce and even eliminate these difficulties. The earlier treatment begins, the better the outcomes.

Parents might need to bring these types of problems to the teacher's attention. Because these boys can be quiet and cooperative in the classroom, teachers may not notice the need for help.

Boys and men with KS can benefit by visiting therapists who are experts in areas such as coordination, social skills, and coping. XXY males might benefit from any or all of the following:

- **Physical therapists** design activities and exercises to build motor skills and strength and to improve muscle control, posture, and balance.
- **Occupational therapists** help build skills needed for daily functioning, such as social and play skills, interaction and conversation skills, and job or career skills that match

interests and abilities.

- **Behavioral therapists** help with specific social skills, such as asking other kids to play and starting conversations. They can also teach productive ways of handling frustration, shyness, anger, and other emotions that can arise from feeling "different."
- **Mental health therapists or counselors** help males with KS find ways to cope with feelings of sadness, depression, self-doubt, and low self-esteem. They can also help with substance abuse problems. These professionals can also help families deal with the emotions of having a son with KS.
- **Family therapists** provide counseling to a man with KS, his spouse, partner, or family. They can help identify relationship problems and help patients develop communication skills and understand other people's needs.

Parents of XXY males have also mentioned that taking part in **physical activities at low-key levels**, such as karate, swimming, tennis, and golf, were helpful in improving motor skills, coordination, and confidence.

With regard to education, some boys with KS will qualify to receive state-sponsored special needs services to address their developmental and learning symptoms. But, because these symptoms may be mild, many XXY males will not be eligible for these services. Families can contact a local school district official or special education coordinator to learn more about whether XXY males can receive the following free services:

- The Early Intervention Program for Infants and Toddlers with Disabilities (<https://www2.ed.gov/programs/osepeip/legislation.html>) is required by two national laws, the Individuals with Disabilities and Education Improvement Act (IDEIA) and the Individuals with Disabilities Education Act (IDEA). Every state operates special programs for children from birth to age 3, helping them develop in areas such as behavior, development, communication, and social play.
- An Individualized Education Plan (IEP) (<https://www2.ed.gov/parents/needs/speced/iepguide/index.html>) for school is created and administered by a team of people, starting with parents and including teachers and school psychologists. The team works together to design an IEP with specific academic, communication, motor, learning, functional, and socialization goals, based on the child's educational needs and specific symptoms.

## Treating Social and Behavioral Symptoms

Many of the professionals and methods for treating learning and language symptoms of the XXY condition are similar to or the same as the ones used to address social and behavioral symptoms.

For instance, boys with KS may need help with social skills and interacting in groups. Occupational or behavioral therapists might be able to assist with these skills. Some school districts and health centers might also offer these types of skill-building programs or classes.

In adolescence, symptoms such as lack of body hair could make XXY males uncomfortable in school or other social settings, and this discomfort can lead to depression, substance abuse, and behavioral problems or "acting out." They might also have questions about their masculinity or gender identity.<sup>9</sup>(</health/topics/klinefelter/conditioninfo/Pages/treatments.aspx#f9>). In these instances, consulting a psychologist, counselor, or psychiatrist may be helpful.

Contrary to research results released decades ago, current research shows that XXY males are no more likely than other males to have serious psychiatric disorders or to get into trouble with the law.<sup>10</sup>(</health/topics/klinefelter/conditioninfo/Pages/treatments.aspx#f10>).

## Citations

1. Bojesen, A., Juul S., & Gravholt, C.H. (2003). Prenatal and postnatal prevalence of Klinefelter syndrome: A national registry study. *Journal of Clinical Endocrinology & Metabolism*, 88(2), 622-626.
2. Dawson, 1997; Hurth, 1999; Rogers, 1989; Hoyson, 1984; Lovaas, 1987; Harris, 1991; McEachin, 1993; Greenspan, 1997; Smith, 1997; and Smith, 1998. As cited in Committee on Children with Disabilities, American Academy of Pediatrics. (2001). The pediatrician's role in the diagnosis and management of autistic spectrum disorder in children. *Pediatrics*, 107(5), 1221-1226.
3. Matsumoto, A.M., Yialamas, M., & Cunningham, G. (2017). Questions and answers: Low testosterone. Hormone Health Network/Endocrine Society. Retrieved September 11, 2019, from <https://www.hormone.org/diseases-and-conditions/low-testosterone>
4. Okada, H., Fujioka, H., Tatsumi, N., Kanzaki, M., Okuda, Y., Fujisawa, M., et al. (1999). Klinefelter's syndrome in the male infertility clinic. *Human Reproduction*, 14(4), 946-952.
5. Herlihy, A. S., & Gillam, L. (2011). Thinking outside the square: Considering gender in Klinefelter syndrome and 47,XXY. *International Journal of Andrology*, 34(5 Pt 2), e348-e34.
6. Paduch, D.A., Fine, R.G., Bolyakov, A., & Kiper, J. (2008). New concepts in Klinefelter syndrome. *Current Opinion in Urology*, 18(6), 621-627.
7. Plotton, I., Brosse, A., Group Fertipreserve, & Lejeune, H. (2011). Infertility treatment in Klinefelter syndrome. *Gynécologie, obstétrique & fertilité*, 39(9), 529-532. French.
8. Plotton I., Brosse A., & Lejeune, H. (2010). Is it useful to modify the care of Klinefelter's syndrome to improve the chances of paternity? *Annales d'endocrinologie (Paris)*, 71(6), 494-504. French.
9. Simpson, J.L., de la Cruz, F., Swerdloff, R.S., Samanga-Sprouse, C., Skakkebaek, N.E., Graham, J.M. Jr., et al. (2003). Klinefelter syndrome: Expanding the phenotype and identifying new research directions. *Genetics in Medicine*, 5(6), 460-468.
10. Ratcliffe, S. (1999). Long-term outcome in children of sex chromosome abnormalities. *Archives of Diseases in Children*, 80(2), 192-195.

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« [What are common symptoms?](/health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx)

(</health/topics/klinefelter/conditioninfo/Pages/symptoms.aspx>)

[How is it diagnosed?](/health/topics/klinefelter/conditioninfo/Pages/diagnosed.aspx) (</health/topics/klinefelter/conditioninfo/Pages/diagnosed.aspx>) »

## How do health care providers diagnose Klinefelter syndrome (KS)?

The only way to confirm the presence of an extra chromosome is by a karyotype (pronounced *care-EE-oh-type*) test. A health care provider will take a small blood or skin sample and send it to a laboratory, where a technician inspects the cells under a microscope to find the extra chromosome. A karyotype test shows the same results at any time in a person's life.

Tests for chromosome disorders, including KS, may be done before birth. To obtain tissue or liquid for this test, a pregnant woman undergoes chorionic villus (pronounced *KAWR-ee-on-ik vil-uhs*) sampling or amniocentesis (*am-nee-oh-sen-TEE-sis*).<sup>1</sup>

(</health/topics/klinefelter/conditioninfo/Pages/diagnosed.aspx#f1>). These types of prenatal testing carry a small risk for miscarriage and are not routinely conducted unless the woman has a family history of chromosomal disorders, has other medical problems, or is above 35 years of age.

## Factors that Influence when KS is Diagnosed

Because symptoms can be mild, some males with KS are never diagnosed.<sup>2</sup>

(</health/topics/klinefelter/conditioninfo/Pages/diagnosed.aspx#f2>).

Several factors affect whether and when a diagnosis occurs:

- Few newborns and boys are tested for or diagnosed with KS.
  - Although newborns in the United States are screened for some conditions, they are not screened for XXY or other sex-chromosome differences.
  - In childhood, symptoms can be subtle and overlooked easily. Only about 1 in 10 males with KS is diagnosed before puberty.<sup>1</sup>  
(</health/topics/klinefelter/conditioninfo/Pages/diagnosed.aspx#f1>).
  - Sometimes, visiting a health care provider will not produce a diagnosis. Some symptoms, such as delayed early speech, might be treated successfully without further testing for KS.
- Most XXY diagnoses occur at puberty or in adulthood.
  - Puberty brings a surge in diagnoses as some males (or their parents) become concerned about slow testes growth or breast development and consult a health care provider.
  - Many men are diagnosed for the first time in fertility clinics.<sup>3</sup>  
(</health/topics/klinefelter/conditioninfo/Pages/diagnosed.aspx#f3>). Among men seeking help for infertility, about 15% have KS;<sup>4</sup>(</health/topics/klinefelter/conditioninfo/Pages/diagnosed.aspx#f4>).

## Citations

1. Aksglaede, L., Skakkebaek, N.E., Almstrup, K., & Juul, A. (2011). Clinical and biological parameters in 166 boys, adolescents and adults with nonmosaic Klinefelter syndrome: A Copenhagen experience. *Acta*

*Paediatrica*, Jun;100(6), 793–806.

2. Bojesen, A., Juul S., & Gravholt, C.H. (2003). Prenatal and postnatal prevalence of Klinefelter syndrome: A national registry study. *Journal of Clinical Endocrinology & Metabolism*, 88(2), 622–626.
  3. Forti, G., Corona, G., Vignozzi, L., Krausz, C., & Maggi, M. (2010). Klinefelter's syndrome: A clinical and therapeutic update. *Sexual Development*, Sep;4(4–5), 249–258.
  4. Ferlin, A., Arredi, B., & Foresta, C. (2006). Genetic causes of male infertility. *Reproductive Toxicology*, Aug;22(2), 133–141.
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« [What are the treatments? \(/health/topics/klinefelter/conditioninfo/Pages/treatments.aspx\)](/health/topics/klinefelter/conditioninfo/Pages/treatments.aspx).  
[Is there a cure? \(/health/topics/klinefelter/conditioninfo/Pages/cure.aspx\)](/health/topics/klinefelter/conditioninfo/Pages/cure.aspx) »

## Is there a cure for Klinefelter syndrome (KS)?

Currently, there is no way to remove chromosomes from cells to "cure" the XXY condition.

But many symptoms can be successfully treated, minimizing the impact the condition has on length and quality of life. Most adult XXY men have full independence and have friends, families, and normal social relationships.<sup>1</sup>[\(/health/topics/klinefelter/conditioninfo/Pages/cure.aspx#f1\)](/health/topics/klinefelter/conditioninfo/Pages/cure.aspx#f1)

They live about as long as other men, on average.<sup>2</sup>  
[\(/health/topics/klinefelter/conditioninfo/Pages/cure.aspx#f2\)](/health/topics/klinefelter/conditioninfo/Pages/cure.aspx#f2)

### Citations

1. Geschwind, D. H., & Dykens, E. (2004). Neurobehavioral and psychosocial issues in Klinefelter syndrome. *Learning Disabilities Research and Practice*, 19(3), 166–173.
2. Bojesen, A., Juul, S., Birkebaek, N., & Gravholt, C. H. (2004). Increased mortality in Klinefelter syndrome. *Journal of Clinical Endocrinology and Metabolism*, 89, 3830–3834.

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« [How is it diagnosed? \(/health/topics/klinefelter/conditioninfo/Pages/diagnosed.aspx\)](/health/topics/klinefelter/conditioninfo/Pages/diagnosed.aspx)  
[Other FAQs \(/health/topics/klinefelter/conditioninfo/Pages/faqs.aspx\)](/health/topics/klinefelter/conditioninfo/Pages/faqs.aspx) »

## Klinefelter Syndrome (KS): NICHD Research Goals

The NICHD has a long history of supporting research to learn more about Klinefelter syndrome. Early research included a study that examined the cells of more than 40,000 infants for extra X chromosomes. NICHD-supported research has also explored topics including the roles of sex chromosomes in development; symptoms that arise in KS such as infertility, low testosterone, and problems with language, learning, and behavior; and how best to treat males with these symptoms. Among the areas of research that hold hope for more successful intervention and prevention in Klinefelter syndrome are studies in the following areas:

- **Genetics of Klinefelter syndrome.** The full extent of the role of the X chromosome in development is not well understood. NICHD research into disorders of the X chromosome, such as Klinefelter, Turner, and Fragile X syndromes, will reveal more about how this chromosome functions and, ultimately, how to prevent or treat symptoms in individuals with an atypical number of X chromosomes. NICHD research also aims to improve understanding of processes that can go wrong in male germ cells before fertilization or right after it, when chromosomes conjugate and divide and can leave the resulting gamete with an unusual number of sex chromosomes.
- **Pathophysiological mechanisms of KS.** KS alters hormonal balance, especially reducing testosterone levels, and exactly how this leads to infertility is unclear. Researchers are studying the mechanisms behind sperm creation and how Leydig cells function, which could identify interventions that may help preserve or restore fertility in males with KS. Investigations also include those on gonadotropin-regulated genes involved in the progression of testicular gametogenesis, Leydig cell function, and other endocrine processes.
- **Treatment strategies for KS.** Research on early interventions has successfully limited the development and severity of symptoms in KS. The NICHD is gathering evidence to identify the best interventions for learning disabilities, osteoporosis (later in life), and infertility—all symptoms of KS.

## **Klinefelter Syndrome (KS): Research Activities and Scientific Advances**

- [Institute Activities and Advances](#)  
(</health/topics/klinefelter/researchinfo/Pages/activities.aspx#institute>)
- [Other Activities and Advances](#)  
(</health/topics/klinefelter/researchinfo/Pages/activities.aspx#other>)

### **Institute Activities and Advances**

KS can influence many aspects of a person's entire life, starting very soon after conception. Therefore, many branches, sections, and laboratories at NICHD conduct research that is relevant to males with XXY or poly-KS variations.

### **Investigating Sex Chromosomes**

KS arises from an unusual number of sex chromosomes, so research into these is important to finding ways to prevent or one day cure KS. Several components of the [Division of Intramural Research \(/about/org/dir/Pages/index.aspx\)](#) are studying these types of problems. The Section on Epigenetics and Development is studying how X chromosome genes influence brain, reproductive, metabolic, and immune system development. The Section on Gamete Development is studying the fruit fly for insight into early gamete cell division and how an additional X chromosome can become included. Other scientists are examining the formation of male germ cells, which are present before and after fertilization and can contain an extra X. In the Section on Clinical Genomics, scientists apply information gained from biochemical and genomic studies to clinical investigations, while also studying the biomechanical mechanisms that may contribute to genetic disorders.

### **Understanding KS Symptoms and Preventing or Treating Them**

Infertility is a key symptom in KS and many researchers at NICHD are involved in improving understanding of how sperm production fails, starting from early in development. In the Section on Clinical Genomics, scientists developed mouse models to analyze proteins that may be key in sperm production. Other research aims to explain the network of genes involved in the renewal and differentiation of spermatogonial stem cells, meiosis, and the post-meiotic differentiation of germ cells. Researchers are also exploring mechanisms behind sperm creation and the function of Leydig cells, which produce testosterone in the presence of luteinizing hormone, and searching for new gonadotropin-regulated genes involved in testicular gametogenesis, Leydig cell function, and other endocrine processes that are disrupted in KS.

Aside from infertility, scientists are working to find ways to treat other symptoms associated with KS. The Child Development and Behavior Branch (CDBB) (</about/org/der/branches/cdbb/Pages/overview.aspx>) is examining the behavioral, neurobiological, and genetic aspects of typical development and is focusing on factors that can threaten normal development. CDBB researchers are also studying prevention steps and, where intervention is needed, the most effective conditions and timing. Their findings will have implications for boys with KS, who can have some learning difficulties, such as in processing language.

Researchers in the Pediatric Growth and Nutrition Branch (</about/org/der/branches/pgnb/Pages/overview.aspx>) focus on nutritional science, childhood antecedents of adult disease, developmental endocrinology, developmental neuroendocrinology, and physical growth and body composition. Topics relevant to KS males include bone weakness and gender identity issues.

## Other Activities and Advances

The projects below also study aspects of health and infertility that might be related to KS.

- The Reproductive Medicine Network (RMN) (</research/supported/Pages/rmn.aspx>), founded in 1990, carries out large, multicenter clinical trials of diagnostic and therapeutic interventions for male and female infertility and reproductive diseases and disorders. The network is funded through the NICHD's Fertility and Infertility (FI) Branch (</about/org/der/branches/fi/Pages/overview.aspx>) and comprises seven research sites as well as a data coordinating center. The RMN currently has several ongoing clinical studies, including a clinical trial to determine a level of oxygen in culture media that improves live birth rates in couples undergoing *in vitro* fertilization.
- The National Centers for Translational Research in Reproduction and Infertility (NCTRI) (</research/supported/Pages/NCTRI.aspx>) (Formerly the Specialized Cooperative Centers Program in Reproduction and Infertility Research [SCCPIR]) is a national network of research-based centers, supported by the FI Branch, that aims to promote interactions between basic and clinical scientists with the goal of improving reproductive health.
- The Learning Disabilities Research Centers Consortium (</research/supported/Pages/ldrc.aspx>) includes four centers in Boulder, Houston, Tallahassee, and Seattle that conduct research on the causes and treatment of learning disabilities. Supported by the NICHD's CDBB, the centers emphasize, among other things, reading comprehension—how children understand what they read—which is difficult for some children with KS.
- The Biological Testing Facility, funded under contract with the Contraceptive Discovery and Development Branch (</about/org/der/branches/crb/Pages/overview.aspx>), has developed radioimmunoassay tests to accurately measure the impact of hormone

treatment given orally, subcutaneously, or transdermally. In individuals with KS taking testosterone, accurate testing helps determine the appropriate dose.

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Klinefelter Syndrome (KS) | NICHD - Eunice Kennedy Shriver National Institute of Child Health and Human Development

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# Turner Syndrome

Turner syndrome is a condition in which a girl or woman is partially or completely missing an X chromosome. It can cause infertility and heart problems and alter a female's appearance. NICHD plays a leading role in advancing research on Turner syndrome by supporting the investigation of its physical and emotional effects as well as potential therapies.

## About Turner Syndrome

Turner syndrome is a disorder caused by a partially or completely missing X chromosome. This condition affects only females.

Most people have 46 chromosomes in each cell—23 from their mother and 23 from their father. The 23rd pair of chromosomes are called the sex chromosomes—X and Y—because they determine whether a person is male or female. Females have two X chromosomes (XX) in most of their cells, and males have one X chromosome and one Y chromosome (XY) in most of their cells. A female with all of her chromosomes is referred to as 46,XX. A male is 46,XY.

Turner syndrome most often occurs when a female has one normal X chromosome, but the other X chromosome is missing (45,X). Other forms of Turner syndrome result when one of the two chromosomes is partially missing or altered in some way.<sup>1</sup>

## Citations

1. Turner Syndrome Society. (n.d.). About Turner Syndrome. <https://www.turnersyndrome.org/about-turnersyndrome> 

## What are the symptoms of Turner syndrome?

Turner syndrome causes a variety of symptoms in girls and women. For some people, symptoms are mild, but for others, Turner syndrome can cause serious health problems. In general, women with Turner syndrome have female sex characteristics, but these characteristics are underdeveloped compared to the typical female. Turner syndrome can affect:<sup>1</sup>

- **Appearance.** Features of Turner syndrome may include a short neck with a webbed appearance, low hairline at the back of the neck, low-set ears, hands and feet that are swollen or puffy at birth, and soft nails that turn upward.
- **Stature.** Girls with Turner syndrome grow more slowly than other children. Without treatment, they tend to have short stature (around 4 feet, 8 inches) as adults.
- **Puberty.** Most girls with Turner syndrome do not start puberty naturally.
- **Reproduction.** In most girls with Turner syndrome, the ovaries are missing or do not function properly. Without the estrogen made by their ovaries, girls with Turner syndrome will not develop breasts. Most women with Turner syndrome cannot become pregnant without assistive technology.<sup>2</sup>
- **Cardiovascular.** Turner syndrome can cause problems with the heart or major blood vessels. In addition, some women and girls with Turner syndrome have high blood pressure.
- **Kidney.** Kidney function is usually normal in Turner syndrome, but some people with this condition have kidneys that look abnormal.
- **Osteoporosis.** Women with Turner syndrome often have low levels of the hormone estrogen, which can put them at risk for osteoporosis. Osteoporosis can cause height loss and bone fractures.
- **Diabetes.** People with Turner syndrome are at higher risk for type 2 diabetes.
- **Thyroid.** Many people with Turner syndrome have thyroid issues. The most common one is hypothyroidism, or an underactive thyroid gland.
- **Cognitive.** People with Turner syndrome have normal intelligence. Some, however, have challenges learning mathematics or with visual-spatial coordination (such as determining the relative positions of objects in space).

## Citations

1. Turner Syndrome Society. (2017). Clinical practice guidelines for the care of women and girls with Turner syndrome. *European Society of Endocrinology*, 117:3, G1-G70. Retrieved 11/29/2017 from [http://docs.wixstatic.com/ugd/8fb9de\\_905ef4f4146a487a9f7031a319b85fe2.pdf](http://docs.wixstatic.com/ugd/8fb9de_905ef4f4146a487a9f7031a319b85fe2.pdf)  (PDF 1.4 MB).
2. Intersex Society of North America. (n.d.). *Turner syndrome*. Retrieved June 14, 2012, from <http://www.isna.org/faq/conditions/turner> 
3. Bondy, C. A. (2007). Care of girls and women with Turner syndrome: A guideline of the Turner Syndrome Study Group. *Journal of Clinical Endocrinology & Metabolism*, 92, 10-25.

## How many people are affected or at risk of Turner syndrome?

Turner syndrome affects about 1 of every 2,500 female live births worldwide.<sup>1</sup>

This disorder affects all races and regions of the world equally. There are no known environmental risks for Turner syndrome. Parents who have had many unaffected children can still have a child with Turner syndrome later on.

Generally, Turner syndrome is not passed on from mother to child. In most cases, women with Turner syndrome are infertile.

### Citations

1. Turner Syndrome Society. (n.d.). *What is Turner syndrome? Fact sheet*. Retrieved July 16, 2012, from <https://www.turnersyndrome.org/about-turnersyndrome> 

## What causes Turner syndrome?

Turner syndrome occurs when part or all of an X chromosome is missing from most or all of the cells in a girl's body. A girl normally receives one X chromosome from each parent. The error that leads to the missing chromosome appears to happen during the formation of the egg or sperm.

Most commonly, a girl with Turner syndrome has only one X chromosome. Occasionally, she may have a partial second X chromosome. Because she is missing part or all of a chromosome, certain genes are missing. The loss of these genes leads to the symptoms of Turner syndrome.<sup>1</sup>

Sometimes, girls with Turner syndrome have some cells that are missing one X chromosome (45,X) and some that are normal. This is because not every cell in the body is exactly the same, so some cells might have the chromosome, while others might not. This condition is called mosaicism (pronounced *moh-ZEY-uh-siz-uhm*). If the second sex chromosome is lost from most of a girl's cells, then it's likely that she will have symptoms of Turner syndrome. If the chromosome is missing from only some of her cells, she may have no symptoms or only mild symptoms.

## Citations

1. National Human Genome Research Institute. (2011). *Learning about Turner syndrome*. Retrieved June 14, 2012, from <https://www.genome.gov/Genetic-Disorders/Turner-Syndrome>

## How do healthcare providers diagnose Turner syndrome?

Healthcare providers use a combination of physical symptoms and the results of a genetic blood test, called a karyotype, to determine the chromosomal characteristics of the cells in a female's body. The test will show if one of the X chromosomes is partially or completely missing.

Turner syndrome also can be diagnosed during pregnancy by testing the cells in the amniotic fluid. Newborns may be diagnosed after heart problems are detected or after certain physical features, such as swollen hands and feet or webbed skin on the neck, are noticed. Other characteristics, like widely spaced nipples or low-set ears, also may lead to a suspicion of Turner syndrome. Some girls may be diagnosed as teenagers because of a slow growth rate or a lack of puberty-related changes. Still others may be diagnosed as adults when they have difficulty becoming pregnant.<sup>1</sup>

### Citations

1. National Human Genome Research Institute. (2011). *Learning about Turner syndrome*. Retrieved July 14, 2012, from <https://www.genome.gov/Genetic-Disorders/Turner-Syndrome>

## What are common treatments for Turner syndrome?

Although there is no cure for Turner syndrome, some treatments can help minimize its symptoms. These include<sup>1</sup>:

- **Human growth hormone.** If given in early childhood, hormone injections can often increase adult height by a few inches.
- **Estrogen replacement therapy (ERT).** ERT can help start the secondary sexual development that normally begins at puberty (around age 12). This includes breast development and the development of wider hips. Healthcare providers may prescribe a combination of estrogen and progesterone to girls who haven't started menstruating by age 15. ERT also provides protection against bone loss.

Regular health checks and access to a wide variety of specialists are important to care for the various health problems that can result from Turner syndrome.<sup>2</sup> These include ear infections, high blood pressure, and thyroid problems.

## Citations

1. National Human Genome Research Institute. (2011). *Learning about Turner syndrome*. Retrieved July 17, 2012, from <https://www.genome.gov/Genetic-Disorders/Turner-Syndrome>
2. Bondy, C. A. (2007). Care of girls and women with Turner syndrome: A guideline of the Turner Syndrome Study Group. *Journal of Clinical Endocrinology & Metabolism*, 92, 10-25.

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