

Forming genitals in boys requires complex dance between genes and hormones

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The most common congenital disorders of all, especially in baby boys, are differences in a newborn's sexual anatomy that is not standard female or male. In boys, they include undescended testicles, misplaced urethras, and improperly developed internal organs.



Some of these disorders may be treated with surgery and some also with hormonal supplementation. Scientists are troubled because the incidence of these disorders is increasing, especially in developed countries. While causes are unknown, researchers suspect that <u>endocrine disruptors</u> from environmental contaminants may be involved.

"It's hard to know how something goes wrong if you don't know how it goes right," says Joan Jorgensen, a professor of comparative biosciences in the University of Wisconsin–Madison School of Veterinary Medicine.

So, Jorgensen, lead author Anbarasi Kothandapani, and others have set out to understand how functional reproductive systems are formed. Over several years, they studied how one gene, GLI3, coordinates the precisely timed development of the male reproductive organs. Using a <u>mouse model</u>, they discovered that GLI3 promotes the growth of the only cells capable of producing <u>male hormones</u> in a fetus. Without enough of those cells, development goes awry.

The researchers published their findings June 4 in the journal *PLOS Genetics*. Their works helps us understand how <u>genes</u> and hormones interact to develop male reproductive systems, a step toward researching why disorders in these organs have become more common and how we might prevent them.

The Jorgensen lab has long studied male and female sex differentiation in developing mice. Altogether, a male fetus must successfully have testes descend toward the scrotum, differentiate external genitals, and develop internal organs such as the prostate. Much of this process is controlled by a genetic pathway whimsically named Hedgehog—after the gene was discovered in flies covered in bristles—and hormones. The two most important hormones are testosterone and another named INSL3.



The GLI3 gene is part of the Hedgehog gene pathway and mutations in the gene are linked to reproductive tract disorders in humans. To study how GLI3 works, the Jorgensen lab turned to a line of mice carrying a mutation that models human mutations in the gene. In mice, the mutation is lethal before birth, so they had to study its effects in developing embryos.

"The first thing we wanted to do was see if they had the same reproductive organ phenotypes as human patients," says Jorgensen. These problems include undescended testicles and disorders in urethra placement.

The <u>mutant mice</u> had several defects similar to humans with mutations in the gene. They experienced delays in the descent of their testicles within their abdomens And in about half of the mutants, the urethra opened at the base of the penis instead of at the tip.

Suspecting a problem with the hormone-producing cells called Leydig cells, the researchers examined the testes. Although mutant mice had typical-looking gonads, Jorgensen's team saw their testes had fewer Leydig cells and these cells even grew weaker as the embryos developed.

"We concluded that the cells started off okay, but they weren't able to maintain their identity and they lost their function," says Jorgensen. "Over time, they weren't able to increase the amount of androgen synthesis needed to make sure everything develops properly."

The researchers discovered that the Hedgehog pathway, through GLI3, stimulates the formation of Leydig cells, which produce the hormones necessary to develop male sex organs. And those hormones go on to reinforce the Hedgehog pathway, which is vital for developing the prostate and penis. They concluded that most of the defects in the mutant mice could be explained by dysfunctional Leydig <u>cells</u>.



Treating pregnant mice with extra testosterone wasn't enough to correct all the defects, because the GLI3 gene was itself necessary, especially for producing a normal penis. But when the researchers reintroduced the nonmutant GLI3 gene into the mutant testes in culture, they were able to restore production of genes necessary for the production of testosterone.

The upshot is that researchers discovered a complex dance between genes and hormones. Any misstep in the waltz can spell disaster.

"There are several environmental disruptors out there that will impact the Hedgehog pathway. When you put that together with genetic mutation, it could make some individuals more susceptible to developmental disorders than others," says Jorgensen.

This improved clarity in how the genes are supposed to function could help researchers understand how environmental toxins might be to blame for a rise in sex development disorders. That understanding is the first step toward prevention.

More information: Anbarasi Kothandapani et al. GLI3 resides at the intersection of hedgehog and androgen action to promote male sex differentiation, *PLOS Genetics* (2020). DOI: 10.1371/journal.pgen.1008810

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