

# Discovery of a new genetic cause of hearing loss illuminates how inner ear works

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A gene called *GAS2* plays a key role in normal hearing, and its absence causes severe hearing loss, according to a study led by researchers in the Perelman School of Medicine at the University of Pennsylvania.

The researchers, whose findings are published online today in

*Developmental Cell*, discovered that the protein encoded by *GAS2* is crucial for maintaining the structural stiffness of support cells in the inner ear that normally help amplify incoming sound waves. They showed that inner ear support cells lacking functional *GAS2* lose their amplifier abilities, causing severe [hearing](#) impairment in mice. The researchers also identified people who have *GAS2* mutations and severe hearing loss.

"Anatomists 150 years ago took pains to draw these support cells with the details of their unique internal structures, but it's only now, with this discovery about *GAS2*, that we understand the importance of those structures for [normal hearing](#)," said study senior author Douglas J. Epstein, Ph.D., professor of genetics at Penn Medicine.

Two to three of every 1,000 children in the United States are born with hearing loss in one or both ears. About half of these cases are genetic. Although hearing aids and cochlear implants often can help, these devices seldom restore hearing to normal.

One of the main focuses of the Epstein laboratory at Penn Medicine is the study of genes that control the development and function of the inner ear—genes that are often implicated in congenital hearing loss. The inner ear contains a complex, snail-shaped structure, the cochlea, that amplifies the vibrations from sound waves, transduces them into nerve signals, and sends those signals toward the auditory cortex of the brain.

### **Unraveling the role of *Gas2* in hearing**

A few years ago, Epstein's team discovered that *Gas2*, the mouse version of human *GAS2*, is switched on in embryos by another gene known to be critical for inner ear development. To determine *Gas2*'s role in that development, the team developed a line of mice in which the gene had been knocked out of the genome and called them *Gas2*-knockout mice.

Alex Rohacek, Ph.D., a former graduate student in the Epstein lab, was puzzled to observe that the *Gas2*-knockout mice had inner ears with cells and structures that seemed quite normal. However, the animals, when tested, turned out to be severely hearing-impaired, with deficits at high sound frequencies of up to 50 decibels—equivalent to a loss of 99.999 percent of the normal acoustic energy.

Tingfang Chen, Ph.D., a postdoctoral fellow and co-first author on the study, determined that *Gas2* is normally active within inner-ear support cells called pillar cells and Deiters' cells. In these cells, the protein encoded by the gene binds to flexible, tube-like structures called microtubules in a way that bundles and stabilizes them, effectively stiffening the cells.

With help from the collaborating team of Benjamin L. Prosser, Ph.D., assistant professor of Physiology at Penn Medicine and an expert on microtubules, the researchers discovered that when pillar cells and Deiters' cells lack *Gas2*, their microtubule bundles tend to come apart, dramatically reducing the stiffness of the cells.

That turns out to have dire implications for hearing. Within the inner ear, pillar cells and Deiters' cells help form the basic structure of the cochlea and serve as physical supports for cells called outer hair cells. The outer hair cells move in response to incoming acoustical vibrations—essentially to provide a crucial amplification of that sound energy. The experiments revealed that the pillar and Deiters' cells' loss of stiffness, due to the absence of *Gas2*, severely degrades the sound-amplifying properties of the outer hair cells they support.

"We observed that some of Deiters' cells in the *Gas2*-knockout mice even buckled under the tension of the rapid movements of the [outer hair cells](#)," Epstein said.

The experiments included sophisticated imaging of propagating sound waves in the inner ears of live *Gas2*-knockout and normal mice, conducted by collaborator John Oghalai, MD, chair and professor of otolaryngology-head and neck surgery at the Keck School of Medicine of USC, and his team.

### ***GAS2* also causes human hearing loss**

Curiously, the researchers could find no reports of *GAS2*-associated congenital hearing loss in the medical literature. Even when they canvassed colleagues around the world who run hearing-loss clinics, they came up empty-handed.

Then one day, Hannie Kremer, Ph.D., professor and chair of molecular otogenetics at Radboud University Medical Center in the Netherlands, emailed Epstein. She and her team had been studying a Somalian family in which four of the siblings had severe hearing loss from early life. The affected family members had no mutations in known hearing-loss genes—but each carried two mutant copies of *GAS2*.

The study therefore establishes *GAS2* as a very probable new hearing loss gene in humans—the first one known to affect the mechanical properties of inner ear support cells.

The prevalence of hearing loss in people due to *GAS2* mutations remains to be determined, but Epstein noted that this type of congenital hearing loss is nevertheless an attractive target for a future gene therapy.

"In many genetic hearing loss conditions, the affected cells are permanently damaged or die, but in this one, the affected [cells](#) are intact and conceivably could be restored to normal or near-normal by restoring *GAS2* function," he said.

He added that such a gene therapy might be useful not only in more obvious cases of hearing loss in early childhood, but also in cases—perhaps more numerous—in which inherited mutations lead to a slower development of hearing loss in adulthood.

**More information:** *Developmental Cell* (2021).

[www.cell.com/developmental-cel ... 1534-5807\(21\)00355-5](https://www.cell.com/developmental-cell/1534-5807(21)00355-5)

Provided by Perelman School of Medicine at the University of Pennsylvania

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