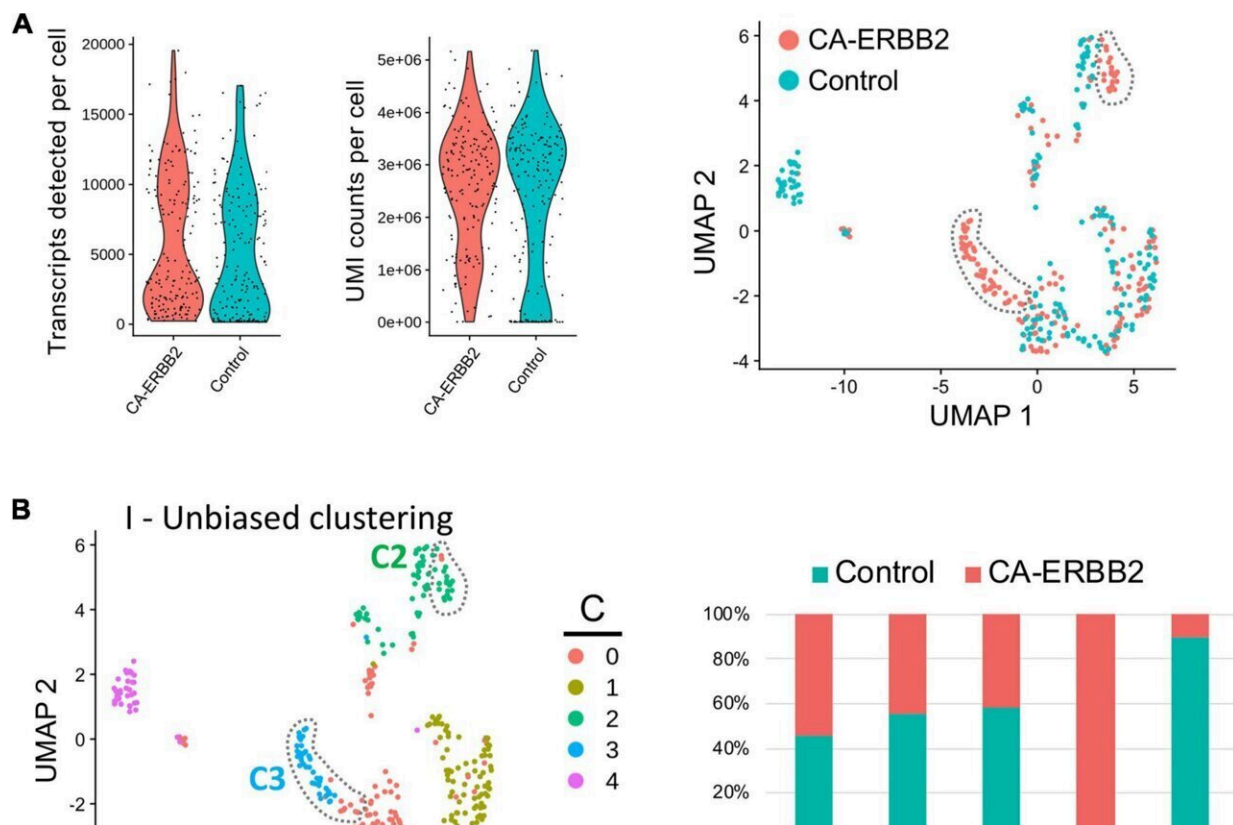


Can hearing loss be reversed? Research reveals clues that could regrow the cells that help us hear

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Unbiased Seurat clustering of cochlear SC transcriptomes from control and CA-ERBB2 samples. (A) Violin plots showing similar distributions observed for cells from control and CA-ERBB2 samples in the number of detected transcripts per cell and in UMI counts per cell used in generating sequencing libraries. On the right, preliminary UMAP plot showing distribution of CA-ERBB2 cells and control cells. Populations of CA-ERBB2 cells that shifted away from control

cells are depicted by dotted line. (B) Initial unbiased clustering showing differences between control cells and CA-ERBB2 cells in the number of clusters and in distribution of cells within the clusters. On right, bar graph showing proportion of control and CA-ERBB2 cells found in each cluster. (C) A second unbiased clustering distinguishes subpopulations within clusters C0, C1, and C2, generating ten unique clusters (S0–S9) with two clusters formed by CA-ERBB2 cells only (S4 and S7) and two clusters mostly composed of Control cells (S2 and S5). On right, bar graph showing proportion of Control and CA-ERBB2 cells found in each cluster. Statistical analyses were performed with Seurat package in R version 4.1.2. Credit: *Frontiers in Cellular Neuroscience* (2023). DOI: 10.3389/fncel.2022.1096872

Taking a bite of an apple is considered a healthy choice. But have you ever thought about putting in earplugs before your favorite band takes the stage?

Just like your future body will thank you for the apple, your future ears (specifically your cochlear [hair cells](#)) will thank you for protecting them. The most common cause of [hearing loss](#) is progressive because these hair cells—the primary cells to detect [sound waves](#)—cannot regenerate if damaged or lost. People who have repeated exposure to loud noises, like military personnel, construction workers, and musicians, are most at risk for this type of hearing loss. But it can happen to anyone over time (even concertgoers).

On the other hand, birds and fish can regenerate these hair cells, and now researchers at the Del Monte Institute for Neuroscience are getting closer to identifying the mechanisms that may promote this type of regeneration in mammals, as explained in research recently published in *Frontiers in Cellular Neuroscience*.

"We know from our previous work that expression of an active growth

gene, called ERBB2, was able to activate the growth of new hair cells (in mammals), but we didn't fully understand why," said Patricia White, Ph.D., professor of Neuroscience and Otolaryngology at the University of Rochester Medical Center. The [2018 study](#) led by Jingyuan Zhang, Ph.D., a postdoctoral fellow in the White lab at the time, found that activating the growth gene ERBB2 pathway triggered a cascading series of cellular events by which cochlear support cells began to multiply and activate other neighboring stem cells to become new sensory hair cells.

"This new study tells us how that activation is happening—a significant advance toward the ultimate goal of generating new cochlear hair cells in mammals," said White.

Using single-cell RNA sequencing in mice, researchers compared cells with an overactive growth gene (ERBB2 signaling) with similar cells that lacked such signaling. They found the growth gene—ERBB2—promoted stem cell-like development by initiating the expression of multiple proteins—including SPP1, a protein that signals through the CD44 receptor. The CD44 receptor is known to be present in cochlear-supporting cells. This increase in cellular response promoted mitosis in the supporting cells, a key event for regeneration.

"When we checked this process in adult mice, we were able to show that ERBB2 expression drove the protein expression of SPP1 that is necessary to activate CD44 and grow new hair cells," said Dorota Piekna-Przybylska, Ph.D., a staff scientist in the White Lab and first author of the study. "This discovery has made it clear that regeneration is not only restricted to the early stages of development. We believe we can use these findings to drive regeneration in adults."

"We plan to further investigation of this phenomenon from a mechanistic perspective to determine whether it can improve auditory function after damage in mammals. That is the ultimate goal," said

White.

Additional authors include Daxiang Na, Cameron Baker, and John Ashton, Ph.D., at the University of Rochester and Medical Center.

More information: Dorota Piekna-Przybylska et al, Single cell RNA sequencing analysis of mouse cochlear supporting cell transcriptomes with activated ERBB2 receptor indicates a cell-specific response that promotes CD44 activation, *Frontiers in Cellular Neuroscience* (2023). DOI: [10.3389/fncel.2022.1096872](https://doi.org/10.3389/fncel.2022.1096872)

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