Promotion of row 1–specific tip complex condensates by Gpsm2-Gai provides insights into row identity of the tallest stereocilia

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The mechanosensory stereocilia in hair cells are organized into rows of graded height, a property crucial for auditory perception. Gpsm2-Gai-Whirlin-Myo15-Eps8 complex at tips of the tallest stereocilia is proposed to define hair bundle row identity, although the underlying mechanism remains elusive. Here, we find that Gpsm2 could undergo phase separation. Moreover, row 1–specific Gpsm2-Gai complex significantly promotes the formation of the five-component tip complex density (5xTCD) condensates. The 5xTCD condensates display much stronger actin-bundling ability than those without Gpsm2-Gai, which may provide critical insights into how Gpsm2-Gai specifies the tallest stereocilia. A deafness-associated mutation of Gpsm2 leads to impaired formation of the 5xTCD condensates and consequently reduced actin bundling, providing possible clues for etiology of hearing loss in patients with Chudley-McCullough syndrome.

INTRODUCTION

Sound detection relies on the epithelial mechanosensory hair cells in the cochlea of inner ear. The mechanically sensitive organelles of hair cells are stereocilia, a cluster of actin-rich protrusions at their apical surface (1, 2). Stereocilia are interconnected by several kinds of hair bundle links such as tip links (3, 4). A unique structural feature of stereocilia is that they are organized into rows of graded height (5) (Fig. 1A). Such a staircase-like pattern is critical for mechanosensory transduction. Mechanical force evoked by sound waves results in stereocilia deflection toward the tallest row, which, in turn, activates the mechanoelectrical transducer channels located in the shorter stereocilia, thus achieving the mechanoelectrical transduction (1, 6).

Proper development of stereocilia is essential for auditory perception, as manifested by severe hearing loss caused by genetic mutations of genes that regulate stereocilia growth (2, 4). Many of these genes encode actin cytoskeleton regulatory proteins that locate at the electron-dense areas of the distal tips of stereocilia (7, 8), playing crucial roles in promoting actin polymerization and bundling. Examples are Whirlin, myosin 15 (Myo15), and epidermal growth factor receptor pathway substrate 8 (Eps8) (9–11). Mice deficient in any of these three genes exhibited similar stereocilia stunting and profound deafness (10, 12, 13). Mechanistically, we and others demonstrated that the three proteins can directly interact with one another to form a tight complex (14, 15) (Fig. 1B). These interactions allow Myo15 motor to transport protein cargos (Whirlin and Eps8) within developing stereocilia where Eps8, an evolutionarily conserved cytoskeletal effector, executes its actin-capping and actin-bundling activities (9). Moreover, the ternary complex forms liquid–liquid phase separation (LLPS) in vitro and in living cells and facilitates actin bundling, indicating that the electron-dense areas located at the tips of stereocilia may form via LLPS of the tip complex in all rows (14).

However, it still remains unclear how stereocilia adopt such a staircase-like architecture and what determines the row identity of the tallest stereocilia. Given the planar asymmetry in the staircase pattern of stereocilia, recent studies on planar cell polarity may provide crucial clues for the possible underlying mechanism (16–20). A complex of Insuteable (Insc), G protein signaling modulator 2 (Gpsm2; also known as LGN), and heterotrimeric G protein Gαi (i.e., mInsc-Gpsm2-Gαi complex) defines an apical compartment (termed as the “bare zone”) with a sharp microvilli exclusion boundary along the mediolateral planar polarity axis to create the V-shaped hair bundles (16). Most notably, Gpsm2-Gai is then restrictedly enriched at the distal tips of stereocilia abutting the bare zone that later become the tallest row (row 1), colocalized with Whirlin, Myo15, and Eps8 (18, 21) (Fig. 1A). Mice deficient in Gpsm2 or Gaii (encoding Gαi) showed similar stereocilia stunting and severe hearing loss as whirler mice (inactivating Whirlin), shaker-2 mice (lacking Myo15), and Eps8-knockout mice did (18, 22–24). Recently, mutations in Gpsm2 were found in patients with Chudley-McCullough syndrome (CMCS; Online Mendelian Inheritance in Man: 604213), a rare autosomal recessive disease characterized by severe to profound sensorineural hearing loss and partial agenesis of the corpus callosum (25–27). Moreover, Gpsm2 was found to interact with Whirlin directly, colocalized into a highly dense nanodomain with a cap-like structure at the tips of the tallest stereocilia (23). Therefore, the five components (Myo15, Eps8, Whirlin, Gpsm2, and Gaii) at the tips of the tallest stereocilia may operate in the same pathway to determine the row identity, although the underlying mechanism is largely unclear.

In this study, we characterize the Gpsm2-Whirlin interaction in detail through a combination of biochemical and biophysical approaches. We found that Gpsm2 itself could form LLPS in vitro and in cells. Gpsm2-Gaii significantly promotes the formation of the five-component tip complex density (5xTCD) condensates (Gpsm2-Gaii-Whirlin-Myo15-Eps8). The promotion effect is most likely due to Gpsm2 LLPS, as a mutant of Gpsm2 that disrupts its LLPS largely impairs the 5xTCD condensates formation. Consistent with their
LLPS capabilities, 5xTCD condensates display much stronger actin-bundling ability than Myo15-Eps8 and Whirlin-Myo15-Eps8 do, which may provide critical insights into how Gpsm2-Gαi specifies the tallest stereocilia and defines hair bundle row identity. Notably, a CMCS-associated mutation of Gpsm2 interferes with the condensate formation and consequently impairs actin bundling, offering possible clues for the etiology of CMCS-related hearing loss.

RESULTS
Characterization of the Gpsm2-Whirlin interaction
Gpsm2-Gαi requires Whirlin as an adaptor to reach the tips of the tallest stereocilia and form the Myo15-Eps8-Whirlin-Gpsm2-Gαi complex specific to row 1 (21). We first set to dissect the interaction between Gpsm2 and Whirlin. Previous report showed that a fragment of Whirlin [amino acids 672 to 810] interacted with N-terminal domain of Gpsm2 (23). Gpsm2 contains eight tetratricopeptide repeats (TPRs) at its N terminus [hereafter Gpsm2 TPR (amino acids 15 to 350)] and four GoLoco (GL) motifs at its C terminus (Fig. 1B). During mitotic spindle orientation of asymmetric cell division, Gpsm2 serves as an adaptor linking the spindle orientation machinery with the cortical polarity cue(s) by binding to mInsc and the nuclear mitotic apparatus (NuMA) via its TPRs and Gαi via its GLs, respectively (28).

We confirmed the Gpsm2-Whirlin interaction by showing that glutathione S-transferase (GST)–Whirlin 673–809 effectively bound to purified Gpsm2 TPR (Fig. 1C). Further truncation-based binding assays indicated that Whirlin 717–746 (hereafter Whirlin GBD, short for Gpsm2-binding domain) was sufficient and necessary for Gpsm2 TPR binding, as further deletion of Whirlin GBD either at its N terminus (Whirlin 737–746) or C terminus (Whirlin 717–733) diminished the binding (Fig. 1C). Moreover, isothermal titration calorimetry (ITC)–based assay using highly purified proteins showed that Gpsm2 TPR binds to WhirlinGBD with a dissociation constant (K_d) of ~1.4 μM (Fig. 1D).

Gpsm2-Whirlin complex structure
To gain insights into the assembly principles of the Gpsm2-Whirlin complex, we tried to crystallize a chimeric construct in which WhirlinGBD was fused C-terminally to Gpsm2TPR. We were able to obtain crystals of the chimera diffracting to 2.6-Å resolution. We solved the complex structure by molecular replacement and refined
it to an $R_{\text{free}}$ of 28.3 and an $R_{\text{work}}$ of 23.1 (table S1). The final structure includes the entire length of Gpsm2TPR and amino acids 729 to 743 of Whirlin.

In the complex structure, Gpsm2TPR folds as a right-handed super-helix creating an inner concave groove that holds the extended WhirlinGBD peptide in an antiparallel manner (Fig. 2A and B). Each TPR repeat consists of two helices, $\alpha$A and $\alpha$B, connected by a short loop (Fig. 2A). The eight TPRs pack together to form a curved $\alpha$-helical solenoid, with $\alpha$A helices facing the inner surface contacting the elongated WhirlinGBD peptide (Fig. 2A). It is noted that the overall conformation of the assembly is reminiscent of those of Gpsm2TPR in complex with other partners such as NuMA, Insc, and Afadin (fig. S1) (29–32).

**The Gpsm2-Whirlin interface**

The Gpsm2TPR-WhirlinGBD interface is mainly mediated by electrostatic interactions and hydrogen bonding interactions. R221/R236 (E to R) form salt bridges with E732 and static interactions and hydrogen bonding interactions. R236 (E to R) forms a hydrogen bond with the main chain of P733Whirlin (Fig. 2C). N100Gpsm2 and R136Gpsm2 form a hydrogen bond and a salt bridge with the main chain and side chain of E732Whirlin, respectively (Fig. 2D). D81Gpsm2 forms additional electrostatic interaction with R739Whirlin (Fig. 2D). It should be mentioned that the hydrophobic interaction between L741Whirlin and L18/A21/L22/F40/A56/I57 from Gpsm2TPR further stabilizes the complex assembly (Fig. 2E). Consistently, mutations that disrupt these polar interactions weakened or even abolished the Gpsm2TPR-WhirlinGBD interaction in ITC-based assays (Fig. 2F).

In particular, substitution of E732 of WhirlinGBD with Ala totally abolished the interaction (Fig. 2F), suggesting that E732 is essential for the interaction, which explains why Whirlin734–746 did not interact with Gpsm2TPR in the GST pull-down assay (Fig. 1C). Notably, key residues in WhirlinGBD that contribute to the interface are absolutely conserved among different species (Fig. 2G), implying the indispensable function of the Gpsm2-Whirlin complex throughout the evolution.

**Gpsm2 undergoes phase separation in vitro and in living cells**

During our investigation of the Gpsm2-Whirlin interaction in cells, we found an interesting discovery that red fluorescent protein (RFP)–Gpsm2 formed condensed puncta in the cytoplasm when expressed in cells (Fig. 3A). In a fluorescence recovery after photobleaching (FRAP) experiment, RFP signals rapidly recovered after photobleaching over a short period of time (Fig. 3, A and B), indicating that RFP-Gpsm2 could freely exchange between the puncta and the surrounding cytoplasm. We next purified full-length Gpsm2 protein and found that it became turbid at room temperature and turned clear after cooling on ice. The turbid solution formed many spherical liquid-like droplets under differential interference contrast (DIC) microscopy (Fig. 3C). Furthermore, Cy3-labeled Gpsm2 formed spherical droplets in a concentration-dependent manner under fluorescence microscopy (Fig. 3C). These droplets could fuse with each other over time (Fig. 3D). Quantification of distribution of droplet sizes both in cells and in vitro revealed that the diameter of most droplets was within the range of 1 to 3 μm (fig. S2). The aforementioned biophysical features of Gpsm2 are reminiscent of biomolecular condensates formed via LLPS in many systems, suggesting that Gpsm2 can autonomously undergo LLPS in vitro and in living cells. Notably, the number of Gpsm2 droplets decreased when the salt concentration was elevated (Fig. 3E), indicating that electrostatic interaction may contribute to LLPS of Gpsm2.

To figure out the mechanism of phase separation of Gpsm2, we made a variety of truncation and deletion of Gpsm2 and investigated their LLPS abilities (Fig. 3F). Deletion of either TPR domain (TPR1 to TPR8) or GL motifs significantly disrupted LLPS (Fig. 3G). Deletion of a linker region (amino acids 421 to 476) between TPR and GL domains also disrupted LLPS. A unique characteristic of this linker is its enrichment with lysine residue (Lys, K) (this linker is hereafter named as “poly-K loop”) (Fig. 3F and fig. S3). We replaced all the lysine residues with Ala (referred to as the “KA” mutant) and found that the Gpsm2KA mutant failed to form LLPS (Fig. 3G). These results, together with the fact that LLPS of Gpsm2 is highly sensitive to salt concentration, indicated that Gpsm2 LLPS may form via multivalent electrostatic interactions between the poly-K loop and other negative charged surface(s) of Gpsm2. Intriguingly, careful sequence analysis revealed that the poly-K loop is absent in activator of G protein signaling 3 (AGS3; also known as Gpsm1), a close
homolog of Gpsm2, although the two proteins share highly conserved functional domains (fig. S3). Given the essential role of the poly-K loop in LLPS, AGS3, which lacks this sequence, did not form LLPS as expected (Fig. 3G).

A CMCS-associated mutation interferes with Gpsm2 LLPS

Notably, a mutation of Gpsm2, p.R318RfsX8, was found in patients with CMCS (23). p.R318RfsX8 encodes a truncated protein lacking the poly-K loop and all four GL motifs (Fig. 3F). Because key elements required for LLPS are missing in this truncated protein, Gpsm2 R318RfsX8 failed to form LLPS as expected (Fig. 3G).

Cocondensation of Gpsm2 and Whirlin

We recently showed that Whirlin could form phase-separated condensation both in vitro and in living cells (14). Because Gpsm2 physically interacts with Whirlin, we next wondered whether the Gpsm2 condensates and the Whirlin condensates can coexist with each other. Green fluorescent protein (GFP)–Whirlin perfectly colocalized with RFP-Gpsm2 in the bright puncta when they were coexpressed in human embryonic kidney (HEK) 293T cells (Fig. 4A). Both GFP and RFP signals in the puncta could recover after photobleaching within a short period of time (Fig. 4B). We also found that mixing equal amounts of Cy5-labeled Gpsm2 and Cy3-labeled Whirlin gave rise to condensed liquid-like droplets in solution (Fig. 4C). FRAP assay showed that both proteins could exchange between the condensed phase and the aqueous solution, although the recovery rates were slower than those in cells (Fig. 4D). Collectively, these results indicated that Gpsm2 and Whirlin could form cocondensates both in vitro and in cells.

Reconstitution of 5xTCD condensates

Myo15, Eps8, Whirlin, Gpsm2, and Gαi interact with each other to form a five-component tip complex specific to the tallest stereocilia (Fig. 1B). Thus, we reasonably hypothesized that Myo15, Eps8, and Gαi could be recruited to the Gpsm2–Whirlin cocondensates because of the interaction network. We tried to verify this point by

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**Fig. 3. Gpsm2 undergoes phase separation in vitro and in living cells.** (A) Representative images showing expression of RFP-Gpsm2 in human embryonic kidney (HEK) 293T cells produced many spherical puncta. FRAP experiments showing that RFP-Gpsm2 signals recovered quickly after photobleaching within a short period of time. The photobleached area is indicated with a yellow dotted box. Scale bar, 5 μm. (B) Quantitative results for FRAP experiments of RFP-Gpsm2 in puncta and cytoplasm of HEK293T cells. Time 0 refers to the time point of the photobleaching pulse. All data are represented as means ± SEM from five droplets (n = 5) in cells. (C) Representative DIC and fluorescence images showing that purified full-length Gpsm2 protein underwent phase separation at indicated concentrations. Gpsm2 was sparsely labeled by Cy3 at 1%. Scale bar, 5 μm. (D) Representative images showing the Gpsm2 liquid-like droplets (indicated by arrows) fused with each other over time. Scale bar, 1 μm. (E) Fluorescence images showing that the number of the Gpsm2 droplets were reduced with increased NaCl concentration. Scale bar, 5 μm. (F) Schematic diagram showing the domain organization of Gpsm2. The deafness-associated mutation Gpsm2R318RfsX8 was also indicated. Note that there is a lysine-rich region (poly-K loop) in the linker region between TPR and GL domains. (G) The percentage of cells showing spherical puncta with various Gpsm2 constructs. All data are expressed as means ± SEM. Five batches of cultures with 30 cells counted in each batch.
reconstitution of the five-component complex in vitro. We were able to purify all these proteins except full-length Myo15 (14). Instead, we successfully purified a truncated form of Myo15 (i.e., C-terminal myosin tail homology 4–4.1/ezrin/radixin/moesin–PDZ binding motif, amino acids 2972 to 3511) that includes both binding sites for Whirlin and Eps8 (Fig. 1B) (14). In the following reconstitution assays, we refer this truncated protein as Myo15\textsuperscript{CTD}. Notably, although we were able to purify full-length Eps8 (Eps8\textsuperscript{FL}), however, the amount of Eps8\textsuperscript{FL} was relatively low, which is not suitable for extensive phase separation study. Instead, we were able to obtain a large amount of a truncated Eps8 (Eps8\textsuperscript{NTD}, Fig. 1B) which includes both the binding sites for Whirlin and Myo15. Notably, Eps8\textsuperscript{NTD} and Eps8\textsuperscript{FL} showed comparable effect on assembly of TCD condensates (fig. S4). Therefore, we used Eps8\textsuperscript{NTD} in the following in vitro reconstitution assay, which was in accordance with our previous work (14). All the protein used in the reconstitution assays were freshly purified and behaved well (fig. S5).

We intended to study LLPS of the five-component complex using labeled proteins under fluorescence microscopy. However, because of the limitation of the fluorescence microscopy technology, we were only able to simultaneously label four proteins at a time. Therefore, we decided to stain for different combinations of the five components. First, we chose not to label Myo15\textsuperscript{CTD} for it could be recruited to the Whirlin condensates together with Eps8\textsuperscript{NTD} in our previous study (14). When mixing fluorescently labeled Eps8\textsuperscript{NTD} (Alexa Fluor 488), Whirlin (Cy3), Gpsm2 (Cy5), and G\alpha\textsubscript{i3} (Alexa Fluor 405) at a 1:1:1:1 molar ratio at their individual concentration of 5 μM, we readily observed micrometer-sized, liquid-phase droplets with spherical shapes under fluorescence microscopy (Fig. 5A). Then, we chose not to label Whirlin and found that Eps8\textsuperscript{NTD} (Alexa Fluor 488), Myo15\textsuperscript{CTD} (Cy3), Gpsm2 (Cy5), and G\alpha\textsubscript{i3} (Alexa Fluor 405) formed codroplets as well (Fig. 5A). Fluorescence images showed that each droplet was highly enriched with each component of the tip complex (Fig. 5A).

These data strongly indicated that the five-component complex can form liquid-phase condensation. To further verify this point, we treated the 5xTCD condensates with 1,6-hexanediol, which has been widely used for LLPS study. Treatment of 1,6-hexanediol greatly impaired the LLPS (Fig. 5B), indicating the reversible character of the 5xTCD condensates. Formation of five-component tip complex condensates was highly specific as they cannot form droplets with an unrelated protein, thioredoxin (Trx) (fig. S6).

Gpsm2-G\alpha\textsubscript{i} significantly promotes LLPS of Whirlin-Myo15\textsuperscript{CTD}-Eps8\textsuperscript{NTD} condensates

Because both Whirlin-Myo15\textsuperscript{CTD}-Eps8\textsuperscript{NTD} (3xTCD) condensates and 5xTCD condensates formed via LLPS, we intended to systematically compare the phase separation abilities among 3xTCD, 4xTCD\textsubscript{Gpsm2} (Gpsm2-Whirlin-Myo15\textsuperscript{CTD}-Eps8\textsuperscript{NTD}), 4xTCD\textsubscript{G\alpha\textsubscript{i3}} (G\alpha\textsubscript{i3}-Whirlin-Myo15\textsuperscript{CTD}-Eps8\textsuperscript{NTD}), and 5xTCD condensates. We found that addition of Gpsm2 (but not G\alpha\textsubscript{i3}) to the 3xTCD significantly increased the number of liquid-like droplets (Fig. 5C). Further addition of G\alpha\textsubscript{i3} into the resulting 4xTCD\textsubscript{Gpsm2} did not further increase the number (Fig. 5C). In addition, we also compared the threshold concentration for LLPS among different complexes. As we demonstrated previously (14), Eps8\textsuperscript{NTD} alone and Eps8\textsuperscript{NTD}-Myo15\textsuperscript{CTD} complex did not form LLPS even at their individual concentration of 30 μM. The 3xTCD formed LLPS at their individual concentration of 5 μM, while 4xTCD\textsubscript{Gpsm2} (but not 4xTCD\textsubscript{G\alpha\textsubscript{i3}}) and 5xTCD formed LLPS at much lower threshold concentration of ~1 μM (Fig. 5D). These data suggested that Gpsm2, but not G\alpha\textsubscript{i}, enhanced LLPS of 3xTCD. The promotion effect is most likely due to the phase separation of Gpsm2, as mutations of Gpsm2 (i.e., Gpsm2\textsuperscript{R318RfsX8}) that impaired the LLPS of Gpsm2 also showed defects in assembly of 5xTCD condensates (Fig. 5, C and D). Consistently, G\alpha\textsubscript{i} did not form LLPS by its own (fig. S7). It should be noted that the threshold concentration for 5xTCD was close to or
even lower than 1 μM, suggesting that the 5xTCD condensates may form at the physiological conditions.

5xTCD condensates induce robust actin bundling
How can the 5xTCD condensates contribute to additional actin elongation at the tallest stereocilia? We next wanted to investigate the correlation between the TCD condensates and actin cytoskeleton dynamics. To this end, we carried out the actin-bundling experiments in the presence of various TCD condensates (Fig. 6A). We used Eps8FL here because Eps8 CTD is necessary for actin bundling (Fig. 1B) (33). Therefore, Eps8 used in the following actin bundling experiments was its full-length form. As expected, Eps8FL alone could induce actin bundling, although with sporadic actin bundles (Fig. 6A).

Addition of Myo15CTD to Eps8FL did not change the activity much (Fig. 6, A and B). The 3xTCD condensates significantly produced more bundled actin filaments, as described in our previous work (14). Consistent with their phase separation abilities (Fig. 5, C and D), 4xTCD Gαi3 condensates exhibited much stronger actin-bundling ability than 3xTCD condensates did, while 5xTCD condensates displayed similar ability as 4xTCD Gαi3 condensates did (Fig. 6, A and B).

Moreover, we used transmission electron microscopy (TEM) to directly visualize actin bundles in the presence of the tip complex condensates. In line with the observation in fluorescent microscopy assays, thicker actin bundles were observed under TEM when 5xTCD or 4xTCD Gαi3 condensates were added in the system than those in 3xTCD condensates (Fig. 6, C and D). The five-component complex
formed codroplets at their individual concentration corresponding to those in the actin-bundling assay (Fig. S8). The 5xTCD components colocalized into small consecutive puncta along with F-actin bundles (Fig. 6E), suggesting that the 5xTCD condensates may act along the F-actin bundles. To further prove that the robust actin-bundling activity of the 5xTCD condensates was attributed to LLPS, we took advantage of the Gpsm2 KA mutant characterized in Fig. 3 because this mutant is expected not to affect any of the interactions among the five-component complex, but instead, it significantly impaired LLPS of the 5xTCD condensates (Figs. 5C and 3F). Satisfyingly, the tip complex mixture with the Gpsm2 KA mutant showed limited actin-bundling capability compared with its wild-type counterpart (Fig. 6, A to D). As expected, the tip complex with the CMCS-associated mutation of Gpsm2, Gpsm2R318RfsX8, showed reduced actin bundles as well (Fig. 6, A to D), probably due to its weakened LLPS ability. These data suggested that formation of the 5xTCD condensates via LLPS is crucial for robust actin-bundling activity at the tips of the tallest stereocilia and consequently may define hair bundle row identity.

**DISCUSSION**

Staircase-like architecture of hair cell stereocilia is crucial for auditory perception, but little is known about how this planar asymmetry is achieved. Recently, several elegant studies on planar cell polarity may provide important clues. In the development of stereocilia, planar-polarized enrichment of Gpsm2-Gαi at a microvilli-free compartment in the apical surface of hair cells enables their specific localization at row 1 immediately adjacent to the microvilli exclusion zone. At the tips of the tallest stereocilia, Gpsm2-Gαi interacts
with the Whirlin-Myo15-Eps8 complex to build a five-component tip complex. It is believed that the complex greatly elongates row 1 and specifies the tallest stereocilia. Given that Gpsm2-Gai complex does not directly involve in actin dynamics, it is still puzzling how Gpsm2-Gai defines the hair bundle row identity.

An interesting observation of our present study may offer critical insights. We find that Gpsm2 can autonomously form LLPS in vitro and in cells. Moreover, the row 1–specific five-component tip complex could also form LLPS. The 5xTCD condensates display much stronger phase separation ability and actin-bundling capability than those without Gpsm2-Gai (Figs. 5 and 6). The promotion effect is mainly attributed to the LLPS of Gpsm2, as a mutant of Gpsm2 (Gpsm2<sup>SKA</sup>) that interferes with its LLPS largely impairs the 5xTCD condensates formation and actin bundling (Figs. 5 and 6). Notably, the fact that Gpsm2-Gai-Whirlin-Myo15<sup>CTD</sup> complex (5xTCD without Eps8) did not induce actin bundling at all demonstrates that Eps8 was the only actin-bundling factor among the complex (Fig. 6).

How does the robust actin bundling event contribute to the overelongation of the tallest stereocilia? Such an actin bundling and property of Eps8 may stem from the ability of actin bundling to stabilize the actin core of stereocilia that further facilitates the actin polymerization at the dynamic filament ends in the tallest stereocilia (34). Alternatively, Eps8-induced actin bundling may cause pronounced barbed-end elongation and thereby makes a longer bundle, as Espin (another actin-bundling protein) does in microvilli (35). It should be noted that the 5xTCD condensates might enhance actin polymerization/nucleation in addition to actin bundling, possibly by condensing higher amounts of Myo15 at row 1, especially when the motor domain of Myo15 was shown to directly accelerate actin filament polymerization by driving nucleation (36). Unfortunately, because of the technical challenges, we were unable to obtain qualified full-length Myo15 protein to prove this point in the current work.

However, our reconstitution experiment has demonstrated that the row 1–specific condensates can condense Myo15<sup>CTD</sup>; hence, it is reasonable to believe that full-length Myo15 would be enriched at row 1 to facilitate hair bundle elongation. Therefore, we propose that Gpsm2-Gai significantly promotes formation of LLPS-mediated 5xTCD condensates that, in turn, largely enrich Eps8 and Myo15 to induce robust actin elongation at the tips of the tallest stereocilia (Fig. 7). Consistently, a recent study has showed that, in Gpsm2<sup>−/−</sup> mutant hair cells, stereocilia are uniformly short, and Myo15-Eps8 is distributed in comparable amounts among all rows, which is in sharp contrast with that in wild-type mice where Myo15-Eps8 is predominantly localized at the tips of the tallest stereocilia (most likely via enrichment effect of row 1–specific TCD condensates) (21).

Compared with Gpsm2, Gai seems to be dispensable for promotion of LLPS. This is expected because Gai could not form LLPS by its own (Fig. S7) and therefore would not add valency into the system to further promote LLPS. We believe that cortical localized Gai may serve as an adaptor between the 5xTCD condensates and cortical membrane, thus acting as a cortical anchoring site for the actin elongation machinery (including but not limited to the 5xTCD condensates). Therefore, it would not be unexpected that depletion of Gai would lead to defects of stereocilia elongation and first row identity as reported previously because the actin elongation machinery would lose its cortical anchoring site in the absence of Gai. This is reminiscent of the scenario in asymmetric cell division where Gai functions as a linker between apical cortical membrane and spindle orientation machinery such as Gpsm2-NuMA complex (28).

Loss of Gai resulted in severe defective asymmetric cell divisions due to the mislocalization of the spindle orientation machinery during the process (37, 38).

How is the assembly/disassembly of the row 1–specific TCD condensates regulated? How are these regulatory processes correlated with stereocilia development? We believe that at least two issues should be taken into account: (i) timing of the interactions among the tip complex and (ii) protein level of the tip complex. Our data show that the protein concentration is critical for LLPS formation. The threshold concentration that promotes Gpsm2 LLPS is about 10 μM; the five-component tip complex could undergo LLPS at their individual concentration of ~1 μM. It should be mentioned that we used a truncation form of Myo15 in the current study. Full-length Myo15 may work as a dimer, which would further expand the valency of the system and lower the threshold concentration of LLPS. In the developing stereocilia, Myo15 and Eps8 formed an early tip complex already broadly enriched at E16.5 (21). By contrast, Whirlin, Gpsm2, and Gai were highly enriched at the exclusion zone at E16.5, and they were enriched at tips at later stages to form the five-component tip complex specific to row 1 (21). When the concentration of tip-accumulated five components reaches the threshold concentration of LLPS, the 5xTCD condensates form autonomously and trigger robust actin elongation event. In support of this idea, Krey et al. (39) recently found that the protein level of the TCD components were temporally regulated during development. Notably, enrichment of the above proteins at tips of the tallest stereocilia was highly correlated with the elongation of row 1 (39). At the mature stage (P21.5), the protein level sharply decreased (39), probably because of gene transcription and/or expression regulation.
or protein degradation system. It is most likely that accumulation of row 1–specific tip complex leads to formation of TCD condensates that promotes robust actin elongation event, while reduction of protein level disrupts LLPS and subsequently stops the elongation process.

We understand that direct investigation of the TCD condensates in hair cells is still challenging at current stage because of technical limit, and future work is definitely needed to demonstrate their critical roles in stereocilia development. Nevertheless, our work provides a proof of concept that condensation of row 1–specific tip complex via LLPS may underlie hair bundle row identity.

**MATERIALS AND METHODS**

**Protein expression and purification**

The coding sequences of Whirlin (GenBank: AB040959.1), Gpsm2 (GenBank: NM_029522.2), AGS3 (GenBank: NM_00135574.1), Gα13 (GenBank: NM_006496.3), Myo15 (GenBank: NM_010862.2), and Eps8 (GenBank: NM_007945.3) were cloned into a previously described modified pET32a vector with a Trx tag and a His6-tag at its N terminus (14). Primers of all constructs are listed in table S2. For the GST pull-down assays, various fragments of Whirlin were cloned into the pGEX-4T-1 vector. Mutations were created through site-directed mutagenesis method and confirmed by DNA sequencing. All freshly purified and labeled proteins were dissolved in the buffer containing 40 mM tris (pH 8.0), 150 mM NaCl, 100 mM NaHCO3 (pH 8.3), and 4 mM dithiothreitol (DTT) and 1 mM β-mercaptoethanol (β-ME). After protein labeling with fluorophore (Fisher Scientific; catalog no. A30000/A20000) were dissolved in dimethyl sulfoxide and incubated with the indicated proteins (molar ratio of 1:1) at room temperature for 1 hour. The labeling reaction was quenched by addition of the buffer of 200 mM tris (pH 8.2). The mixture was then purified with a HiTrap desalting column with the buffer containing 50 mM tris (pH 8.0), 300 mM NaCl, and 4 mM β-ME to remove the unlabelled fluorophore. Fluorescence labeling efficiency was determined by NanoDrop 2000 (Thermo Fisher Scientific).

**Protein labeling with fluorophore**

Freshly purified untagged proteins were labeled as described previously (14). In general, proteins were first dissolved in the buffer containing 300 mM NaCl, 100 mM NaHCO3 (pH 8.3), and 4 mM β-mercaptoethanol (β-ME). Cy-3/Cy5 N-hydroxysuccinimide (NHS) ester (AAT Bioquest; catalog no. 271/280) and Alexa Fluor 405/488 NHS ester (Thermo Fisher Scientific; catalog no. A30000/A20000) were dissolved in dimethyl sulfoxide and incubated with the indicated proteins (molar ratio of 1:1) at room temperature for 1 hour. The labeling reaction was quenched by addition of the buffer of 200 mM tris (pH 8.2). The mixture was then purified with a Hitrap desalting column with the buffer containing 50 mM tris (pH 8.0), 300 mM NaCl, and 4 mM β-ME to remove the unlabelled fluorophore. Fluorescence labeling efficiency was determined by NanoDrop 2000 (Thermo Fisher Scientific).

**ITC assay**

ITC assays were performed on a MicroCal iTC200 system (Malvern Panalytical, UK) at 25°C. Various Gpsm2TPR and WhirlinGRD fragments were dissolved in the buffer containing 50 mM tris (pH 8.0), 100 mM NaCl, 1 mM EDTA, and 1 mM DTT. The WhirlinGRD proteins (~500 μM) were loaded into syringe, and the Gpsm2TPR proteins (~50 μM) were loaded into the cell. In each titration, 2-μl aliquot of protein in the syringe was injected into the cell, and the time interval was 120 s to make sure that the titration peak returned to the baseline. Titration data were fitted with the one-site binding model using Origin 7.0.

**GST pull-down assay**

Freshly purified His6-Gpsm2TPR was incubated with various forms of GST-Whirlin fragments for 1 hour at 4°C. After centrifugation for 10 min at 4°C, the supernatant was loaded into 30-μl GSH-sepharose 4B slurry beads to incubate for 30 min at 4°C. After washing with PBS buffer three times, the bound proteins were eluted by boiling with 30 μl of 2 × SDS–polyacrylamide gel electrophoresis loading dye and detected by Western blot using anti-His antibody (Smart-lifesciences, catalog no. SLAB28; 1:5000).

**Crystallization, data collection, and structure determination**

To obtain stable Gpsm2TPR–WhirlinGRD complex, Gpsm2TPR (amino acids 15 to 350) was fused with WhirlinGRD (amino acids 717 to 746). The best crystals of the fusion protein (~10 mg/ml) were obtained by the hanging drop diffusion method at 16°C in the buffer containing 1.2 M ammonium tartrate dibasic and 0.1 M tris (pH 8.8). Crystals were cryoprotected in the corresponding reservoir solution with 25% glycerol before x-ray diffraction experiments. The diffraction data were collected at BL41XU at Spring-8 (Hyogo, Japan). The diffraction data were processed with XDS (40). The complex structure was solved by the molecular replacement method using the structure of GPSM2TPR in Gpsm2-NuMA complex (Protein Data Bank code: 3RO2) as the searching model through the software suits of Phaser (41). Further refinement was performed using Phenix (42) and Coot (43). The final refinement statistics of the complex structures are listed in table S1. Structural diagrams were prepared by PyMOL.

**In vitro phase separation assay**

All freshly purified and labeled proteins were dissolved in the buffer containing 50 mM tris (pH 8.0), 300 mM NaCl, and 4 mM β-ME. After centrifugation at 16,873g for 10 min at 4°C, samples were placed on ice before the phase separation assay. Each phase separation sample was made by mixing of indexed labeled proteins at indicated concentrations for 10 min at 25°C. Each sample was then injected into a homemade chamber as described previously (44) for fluorescence imaging (Leica TCS SP8).

**FRAP assay**

FRAP assay was performed as previously described (14). In brief, the assay was carried out on a Leica SP8 confocal microscope. For Cy3-labeled protein droplets, a circular region of interest (ROI) was bleached by a 561-nm laser beam at room temperature. For FRAP assay on liquid-phase droplets in living cell, HEK293T cells (American Type Culture Collection; catalog no. CRL-3216; Research Resource Identifiers: CVCL_0063) were cultured on glass-bottom dishes (MatTek) and transfected with the indicated plasmids. GFP and RFP signals were bleached with 488- and 561-nm laser beams at 37°C, respectively. For each experiment, the fluorescence intensity of a neighbor droplet with similar size to the bleached one was also recorded for intensity correction. Background intensity was subtracted before data analysis. The ROI intensity at time 0 s (right after the photo-bleaching) was set as 0%, and the prebleaching intensity was normalized to 100%.
Actin-bundling assay
To obtain preassembled F-actin filament, monomeric rabbit G-actin (Cytoskeleton) was induced to polymerize for 1 hour at room temperature in the polymerization buffer containing 50 mM tris (pH 8.0), 1 mM adenosine 5′-triphosphate, 0.5 mM DTT, 0.2 mM CaCl₂, 2 mM MgCl₂, and 50 mM KCl. Freshly purified wild-type and mutant form of five-component tip complex mixture and other tip complexes without Gpsm2-GαI3 were incubated with the above preassembled F-actin (~2 μM) at room temperature for 1 hour. Actin was then labeled with rhodamine-phalloidine (Cytoskeleton) for 15 min. The samples were carefully mounted between a slide and a coverslip and imaged by fluorescence microscopy (Leica TCS S8P). In these assays, the concentration of each component was shown as follows: Eps8 FL, 0.25 μM; Whirin, 0.5 μM; Myo15β(CTD), 0.5 μM; Gpsm2 WT/mutants, 0.5 μM; GαI3, 0.5 μM; F-actin, ~2 μM.

Samples for TEM (Tecnai G2 Spirit 120 kV) were adsorbed to glow-discharged, carbon-coated formvar films on copper grids for 1 min and negatively stained with 0.75% (m/v) uranium acetate for a few seconds.

Statistical analyses
All statistics [e.g., number of samples (n) and biological replicates (N) for all experiments] were described in the figure legends. All data were represented as means ± SEM using the two-tailed Student’s t test, using GraphPad Prism. All experiments were performed at least three times independently.

SUPPLEMENTARY MATERIALS
Supplemental material for this article is available at https://doi.org/10.1126/sciadv.abn4556

View request a protocol for this paper from Bio-protocol.

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Data and materials availability: All data needed to evaluate the conclusions in the paper are present in the paper and/or the Supplementary Materials. The atomic coordinate of the Gpsm2-Whirlin complex has been deposited to the Protein Data Bank under the accession code: 7EP7.

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