

Article

Phase separation-mediated condensation of Whirlin-Myo15-Eps8 stereocilia tip complex

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SUMMARY

Stereocilia, the mechanosensory organelles on the apical surface of hair cells, are necessary to detect sound and carry out mechano-electrical transduction. An electron-dense matrix is located at the distal tips of stereocilia and plays crucial roles in the regulation of stereocilia morphology. Mutations of the components in this tip complex density (TCD) have been associated with profound deafness. However, the mechanism underlying the formation of the TCD is largely unknown. Here, we discover that the specific multivalent interactions among the Whirlin-myosin 15 (Myo15)-Eps8 complex lead to the formation of the TCD-like condensates through liquid-liquid phase separation. The reconstituted TCD-like condensates effectively promote actin bundling. A deafness-associated mutation of Myo15 interferes with the condensates formation and consequently impairs actin bundling. Therefore, our study not only suggests that the TCD in hair cell stereocilia may form via phase separation but it also provides important clues for the possible mechanism underlying hearing loss.

INTRODUCTION

Usher syndrome (USH) is an autosomal recessive genetic disease characterized by deafness and vision impairment (Boughman et al., 1983; Keats and Corey, 1999; Mathur and Yang, 2019). A lot of USH causative genes have been identified during the last several decades (Michalski and Petit, 2019; Petit and Richardson, 2009). These genetic data reveal that the abnormal development of hair cells may underlie the pathogenesis of the syndrome (Barr-Gillespie, 2015). Hair cells are specialized mechanoreceptor cells that detect sound and then carry out mechano-electrical transduction (MET). The mechanosensory organelles of hair cells are hair bundles or stereocilia, a cluster of actin-rich protrusions located at the apical surface of polarized hair cells (Barr-Gillespie, 2015; Gillespie and Müller, 2009; Figure 1A). Stereocilia are organized into rows of graded heights forming precisely uniform staircase patterns, which are interconnected by various extracellular hair bundle links, such as tip links (Gillespie and Müller, 2009; Kachar et al., 2000; Richardson and Petit, 2019; Sakaguchi et al., 2009; Figure 1A). Electron microscopy (EM) analysis has revealed that the lower and upper insertion sites of tip links are electron-dense areas that are referred to as lower tip link density (LTLD) and upper tip link density (UTLD),

respectively (Furness and Hackney, 1985; Gillespie and Müller, 2009; Sahin et al., 2005; Figure 1A).

Proper development and maintenance of stereocilia are essential for normal hair cell functions (Barr-Gillespie, 2015; Richardson and Petit, 2019). It is not fully understood, however, how stereocilia length is controlled. Intriguingly, several proteins encoded by USH causative genes have been involved in the regulation of actin cytoskeleton dynamics in stereocilia (Drummond et al., 2012). Deafness-associated mutations of these actin cytoskeleton regulatory proteins perturb stereocilia growth and consequently lead to hearing loss (Sahin et al., 2005). Specifically, a set of protein complexes located at the distal tips of the tallest stereocilia (also known as “tip complex”), which includes Whirlin, myosin 15 (Myo15), Eps8, LGN, and Gzi, plays crucial roles in promoting actin polymerization and bundling (Belyantseva et al., 2003, 2005; Mauriac et al., 2017; Sahin et al., 2005; Zampini et al., 2011; Figure 1A).

Myo15, an unconventional myosin expressed in cochlear hair cells, is thought to transport protein cargos within developing stereocilia (Belyantseva et al., 2005; Delprat et al., 2005). Mutations of Myo15 cause profound deafness both in humans and mice (Probst et al., 1998; Wang et al., 1998). Mice deficient in Myo15 (*shaker-2* mice) displayed very short stereocilia without





Figure 1. Characterization of interactions between Whirlin, Myo15, and Eps8

(A) A schematic diagram illustrating the organization of hair cell stereocilia.

(B) Domain organizations of key components of the tip complex in stereocilia. FERM, 4.1/ezrin/radixin/moesin; HHD, Harmonin homology domain; Myth4, myosin

tip links (Probst et al., 1998). Whirlin (encoded by *WHRN* gene) is a USH protein that sequentially consists of N-terminal harmonin homology domain 1 (HHD1), a tandem of PDZ domains (PDZ1–2), HHD2, and the PDZ3 domain directly followed by a C-terminal PDZ-binding motif (PBM) (Mburu et al., 2003; Figure 1B). Whirlin acts as an organizer of the tip complex that control the coordinated actin dynamics of stereocilia (Kikkawa et al., 2005). Mutations of *WHRN* have been found in patients with either non-syndromic deafness (DFNB31) (Mburu et al., 2003) or USH of type 2 (USH2D) (van Wijk et al., 2006). Mice deficient in Whirlin (*whirler* mice) were profoundly deaf and showed short stereocilia (Mburu et al., 2003). Eps8 (epidermal growth factor receptor pathway substrate 8) is an evolutionarily conserved regulator of actin dynamics and can control actin-based motility by capping the barbed ends of actin filaments and promoting actin bundling (Disanza et al., 2004, 2006). In addition, Eps8 can directly interact with monomeric actin (Hertzog et al., 2010). Like *shaker-2* and

whirler mice, *Eps8* knockout mice also exhibit similar stereocilia stunting and severe hearing loss (Zampini et al., 2011), indicating a strong genetic interaction between these three deafness-associated genes (Behloul et al., 2014). Physically, Myo15 interacts with both Whirlin and Eps8 (Manor et al., 2011). These interactions are required for Myo15-mediated trafficking of Whirlin and Eps8 to the very tips of stereocilia (Belyantseva et al., 2003, 2005). Whirlin was also found to interact with Eps8 in the tip complex (Manor et al., 2011). Expression of both Eps8 and Myo15 at stereocilia tips was reduced in Whirlin-deficient mice (Manor et al., 2011), suggesting a critical scaffolding role of Whirlin in the assembly of the stable Whirlin-Myo15-Eps8 tip complex.

Most notably, transmission EM (TEM) images have shown that an electron-dense structure is located at the distal tips of stereocilia and covers the barbed ends of the actin protrusions (Engström and Engström, 1978; Furness and Hackney, 1985;

Hackney and Furness, 2013; Mogensen et al., 2007; Rzadzinska et al., 2004; Figure 1A). This dense structure exhibits properties reminiscent of the postsynaptic density (PSD) in neurons (Palay, 1956). PSD contains various proteins and forms a densely packed structure to regulate actin cytoskeleton dynamics in accordance with distinct neuronal stimuli (Zhu et al., 2016). Similarly, the components of the tip complex may act as key molecular organizers of this tip complex density (TCD) (Gillespie and Müller, 2009; Sahin et al., 2005). For example, the tallest stereocilia showed very strong Myo15 staining covering their oblate tips, whereas the *shaker-2* mice exhibited a rounded tip without the characteristic TCD and displayed bundle defects (Rzadzinska et al., 2004). The electron density at the tips of stereocilia was markedly patchy and irregular in the *whirler* mice (Mogensen et al., 2007). Intriguingly, the electron-dense material-free areas appeared to coincide with a lack of actin filament at the tip (Mogensen et al., 2007). These observations also suggest a tight correlation between the TCD formation and stereocilia growth. Despite the crucial roles of the tip complex-mediated formation of TCD in orchestrating the architecture and function of hair cell stereocilia, several key questions remain to be answered. What is the molecular basis underlying the assembly of the tip complex? How could the tip complex contribute to form electron-dense TCD assembly? What is the relationship between TCD formation and the pathogenesis of USH?

In this study, we characterize the interactions between Whirlin, Myo15, and Eps8 in detail and provide the high-resolution structure of the Whirlin-Myo15 complex. Importantly, we demonstrate that the formation of the Whirlin-Myo15-Eps8 complex leads to autonomous condensation of the complex both *in vitro* and in cells via liquid-liquid phase separation (LLPS). Our biochemical reconstitution experiments suggest that the key components of the tip complex are sufficient to form the densely packed TCD-like condensates. We further demonstrate that the reconstituted TCD-like condensates effectively promote actin bundling. A deafness-associated mutation of Myo15 impairs the formation of the condensates and thus leads to defects in actin bundling.

RESULTS

Characterization of the interactions between Whirlin, Myo15, and Eps8

We wanted to dissect the interactions among Whirlin, Myo15, and Eps8. A previous study reported that Whirlin binds to the N-terminal region (amino acids [aa] 1–535) of Eps8 (Manor et al., 2011). We verified this interaction by showing that glutathione S-transferase (GST)-Eps8^{1–529}, but not GST-Eps8^{530–821}, bound robustly to GFP-tagged Whirlin in the GST-pull down assays (Figure 1C). We further found that Eps8^{207–464} (referred to as Whirlin-binding domain [WBD]) also bound effectively to Whirlin, whereas Eps8 phosphotyrosine binding (PTB) (aa 33–207) did not (Figure 1C). In line with our biochemical data, both RFP-Eps8^{1–529} and RFP-Eps8 WBD could colocalize with GFP-Whirlin into the cytoplasmic puncta-like structures in cells, whereas red fluorescent protein (RFP)-Eps8 PTB could not (Figure S1A). Next, we tin

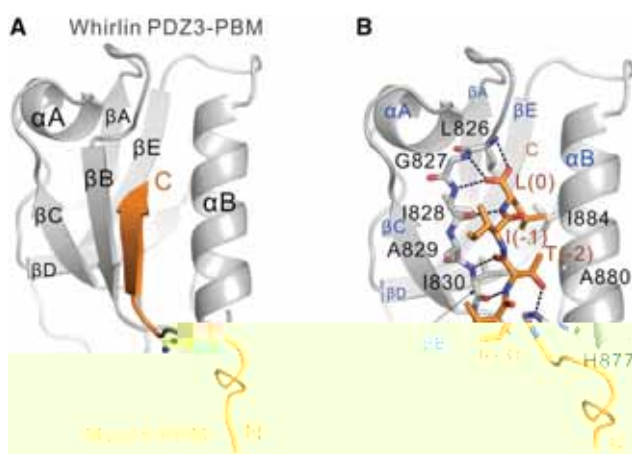


Figure 2. Crystal structure of Whirlin-Myo15 complex
(A) Overall structure of the Whirlin PDZ3-PBM-Myo15 PBM complex.
(B) Detailed interface between Whirlin PDZ3-PBM and Myo15 PBM. Residues involved in protein interaction are shown with the stick model. Hydrogen bonds are shown as black dashed lines.
See also Figure S2 and Table S1.

formed by L826, I828, I830, and I884 from Whirlin PDZ3. The carboxyl group of L(0) of Myo15 PBM forms hydrogen bonds with the backbone amides of L826, G827, and I828 (the “GLGF” motif of PDZ domain) of Whirlin PDZ3. The side chain of T(-2) of Myo15 PBM interacts with the imidazole ring of H877 at the α B helix of Whirlin PDZ3 (Figure 2B).

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

An interesting observation during our cellular imaging study was that GFP-Whirlin formed bright puncta in cytosol when expressed in cells (Figure 3A). A fluorescence recovery after photobleaching (FRAP) experiment showed that GFP-Whirlin signals could be recovered after photobleaching over a short period of time (Figures 3A and 3B), suggesting that GFP-Whirlin could exchange between the puncta and the surrounding cytoplasm. These data indicated that the puncta were reversible liquid droplets. We purified the recombinant full-length Whirlin protein and found that Whirlin protein became turbid at room temperature when the concentration was $>20 \mu\text{M}$ and turned clear again after being cooled on ice. Under differential interference contrast (DIC) microscopy, the turbid solution formed many spherical liquid droplets (Figure 3C). Moreover, Cy3-labeled Whirlin protein forms bright droplets in solution in a concentration-dependent manner under fluorescence microscopy (Figure 3C). These droplets could fuse with one another into larger spherical droplets over time (Figure 3D). The FRAP experiment showed that the recombinant the Cy3 signals could recover after photobleaching as well, suggesting that Whirlin could exchange between the droplets and the surrounding solution (Figure 3E). We further tested the ability of Whirlin to form liquid droplets in buffers with different salt concentrations. Significantly, both the size and number of the droplets were reduced when the NaCl concentration was

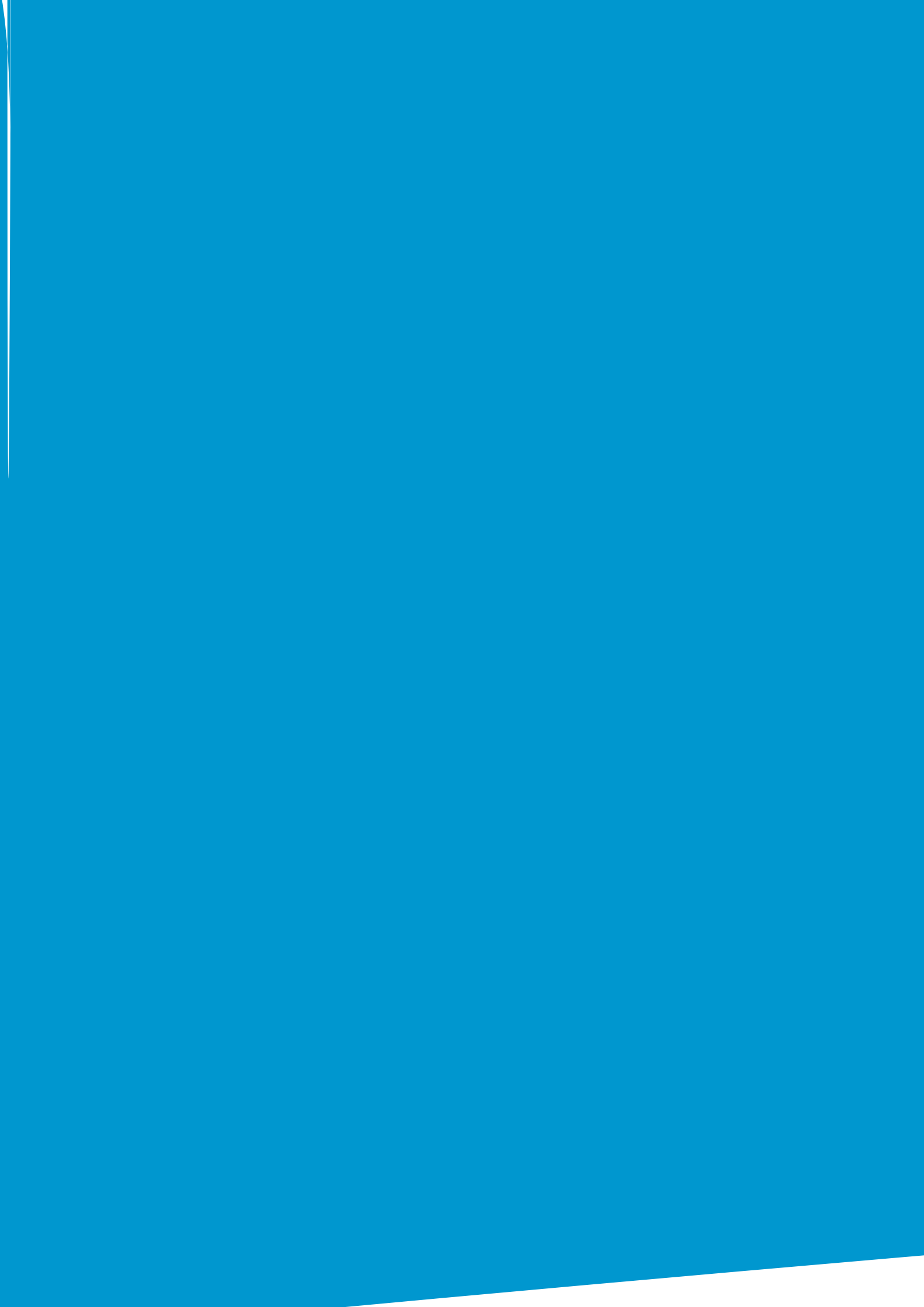
elevated (Figure 3F). The above biophysical features of Whirlin are reminiscent of biomolecular condensates formed via LLPS (Banani et al., 2017; Hyman et al., 2014; Shin and Brangwynne, 2017), suggesting that Whirlin can autonomously undergo phase separation *in vitro* and in living cells. Previous reports demonstrated that Whirlin can form oligomers (Chen et al., 2014; Delprat et al., 2005). More specifically, HHD1-PDZ1 was found to bind to itself as well as to PDZ2, and PDZ3 was also found to bind to itself (Chen et al., 2014). One would thus expect that these multivalent intermolecular interactions in Whirlin may contribute to its phase separation. In line with this hypothesis, a short isoform of Whirlin (Whirlin^{short}), which lacks the HHD1, PDZ1, and PDZ2 domains, could not form phase separation (Figure S4).

Formation of Whirlin-Myo15-Eps8 complex leads to TCD-like condensates

The fact that Whirlin, Eps8, and Myo15 interact specifically with one another may allow Whirlin-Myo15-Eps8 complex to form open oligomers (see below), thus promoting the formation of the liquid-like condensates. To verify this hypothesis, we decided to reconstitute the Whirlin-Myo15-Eps8 complex *in vitro*. We tried to purify recombinant full-length Myo15 and Eps8. However, we failed to obtain soluble full-length Myo15 protein. Alternatively, we successfully purified a truncated Myo15 protein (Myo15 CMF-PBM), which includes both binding sites for Whirlin and Eps8 (Figure 3A). Although we could purify full-length Eps8 protein, the amount was relatively limited, which is not suitable for extensive phase separation study. Instead, we were able to obtain large amounts of a truncated Eps8 (Eps8 PTB-WBD), which includes both binding sites for Whirlin and Myo15 (Figure 4A). For simplicity, we referred to these two truncated proteins as Myo15 and Eps8, respectively, during our subsequent reconstitution experiments.

When mixing fluorescently labeled Whirlin, Myo15, and Eps8 at a 1:1:1 molar ratio, we observed micrometer-sized, condensed droplets with spherical shapes under fluorescence microscopy (Figure 4B). Each droplet was enriched with all three proteins. Importantly, the number and size of the droplets progressively reduced when the concentration of the complex decreased (Figure 4B). Notably, the addition of Myo15 and Eps8 significantly lowered the threshold concentration of phase separation to $\leq 5 \mu\text{M}$ (Figures 4B versus 3C). The FRAP experiment suggested that each protein could exchange between the condensed phase and the aqueous solution (Figure 4C). Consistently, the three proteins could perfectly colocalize with each other into bright spherical puncta when they were co-expressed in cells (Figure 4D). The intensities of each protein in the cytoplasmic puncta could recover after photobleaching over a short period of time (Figure 4E). Notably, the recovery rate of Whirlin in the TCD-like condensates is much slower than that of Whirlin alone, suggesting that the TCD-like condensates may be highly dense and less dynamic.

Although the biomolecular condensates are in general driven by low-complexity sequences of intrinsic disordered regions (IDRs) (Banani et al., 2017; Shin and Brangwynne, 2017; Wright and Dyson, 2015), specific and multivalent protein-protein



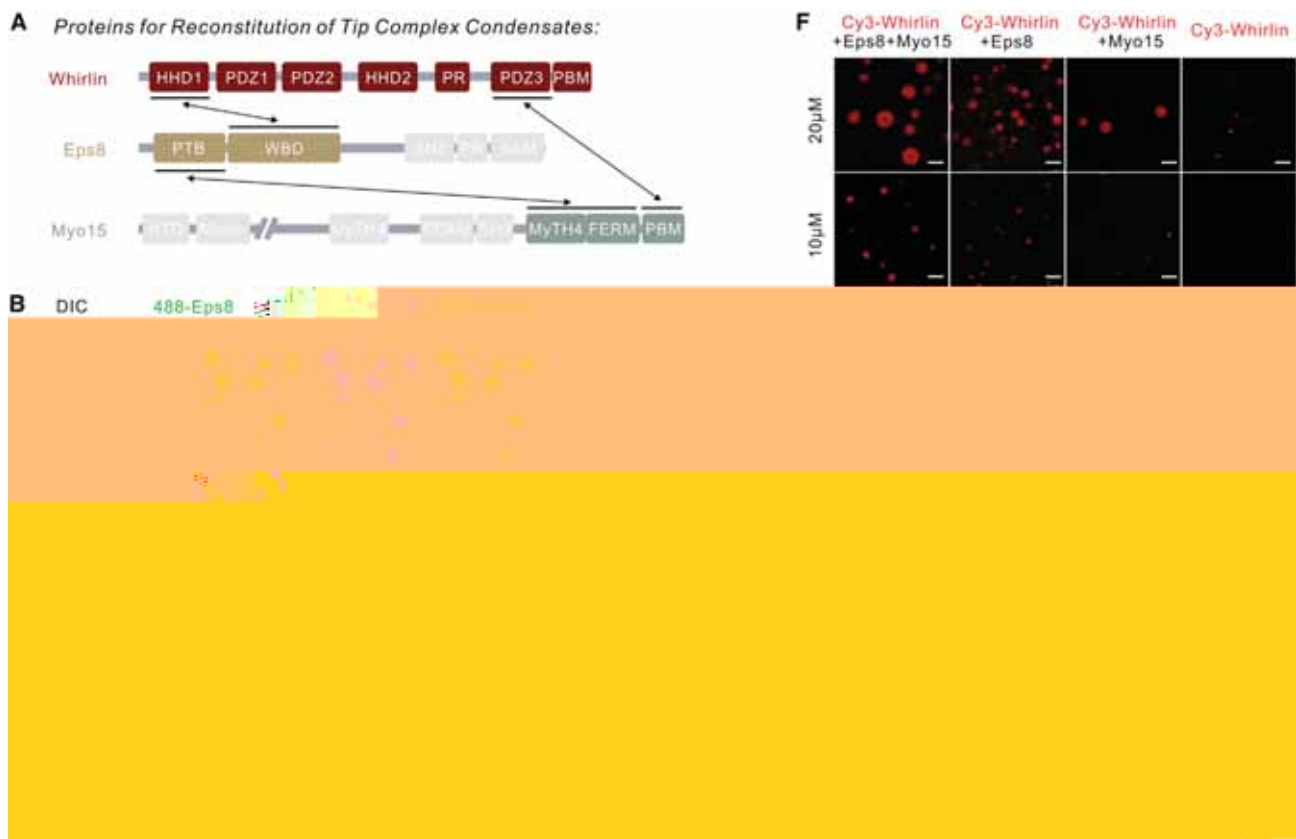


Figure 4. Formation of Whirlin-Myo15-Eps8 tip complex leads to TCD-like condensates

(A) Schematic diagram showing the components and the interactions among Whirlin, Myo15, and Eps8. Domains drawn in gray are removed from the corresponding proteins for technical reasons. Black arrows indicate the interactions between these proteins.

(B) DIC and fluorescence images showing that mixture of Whirlin, Myo15, and Eps8 at indicated concentrations led to formation of the liquid droplets. Whirlin, Myo15, and Eps8 were labeled with Cy3, Alexa 488, and Alexa-405, respectively, with each at 1% level. Scale bar: 5 μ m.

(C) FRAP experiment showing the exchange kinetics of each protein between the tip complex condensates and dilute solution. All data are represented as means \pm SDs from 5 independent experiments (N = 5).

(D) Representative images showing that co-expression of GFP-Myo15, RFP-Whirlin, and BFP-Eps8 in HEK293T cells produced multiple spherical puncta enriched with 3 proteins. Scale bar: 7.5 μ m.

(E) FRAP experiment showing the exchange kinetics of each protein between the condensed puncta and dilute cytoplasm. All data are represented as means \pm SDs from 5 droplets (n = 5) in cells.

(F) Valency- and protein concentration-dependent TCD-like condensates formation. Scale bar: 5 μ m.

(G) A model depicting the mechanism of the formation of the TCD-like condensates via phase separation.

See also [Figures S3, S4, and S7](#).

We next wanted to figure out how the deafness-associated Myo15 mutant affects the formation of the TCD-like condensates. We used a previously described sedimentation-based assay to quantify the distribution of the proteins in the bulk aqueous solutions (the “supernatant” fraction) and the condensed droplets (the “pellet” fraction) (Zeng et al., 2016; Zhu et al., 2020). In the wild-type tip complex mixture, ~95% of Whirlin and ~50% of Eps8 and Myo15 proteins were recovered from the condensed phase (Figures 5C and 5D). However, in the Myo15^{deaf} group, the number of proteins in the condensed droplets significantly decreased (only with ~50% of Whirlin and ~10% of Eps8 and Myo15 proteins) (Figures 5C and 5D). It is worth noting that all of the proteins used in the sedimentation-based assays were soluble and did not form aggregations in their

apo forms (Figure S4). In addition, compared with the wild-type tip complex condensates, the ones with either Myo15^{deaf} or Myo15 ^{Δ PBM} showed fewer liquid droplets under microscopy (Figures 5E and 5F). These data indicated that the deafness-associated mutation of Myo15 leads to defective TCD-like condensate formation, most likely due to the impairment of multivalent interactions in the tip complex.

The tip complex condensates promote actin bundling

Little is known about how the components of the tip complex work together to orchestrate the regulation of actin dynamics in stereocilia, especially when some of them do not possess actin regulatory activity. Our reconstituted TCD-like condensates may provide some clues. In the reconstitution system,

only Eps8 is able to facilitate actin bundling via its C-terminal region, including the SH3, the proline-rich motifs, and the sterile alpha motif (SAM) domain ([Disanza et al., 2004](#)). Full-length Eps8 (Eps8^{FL})

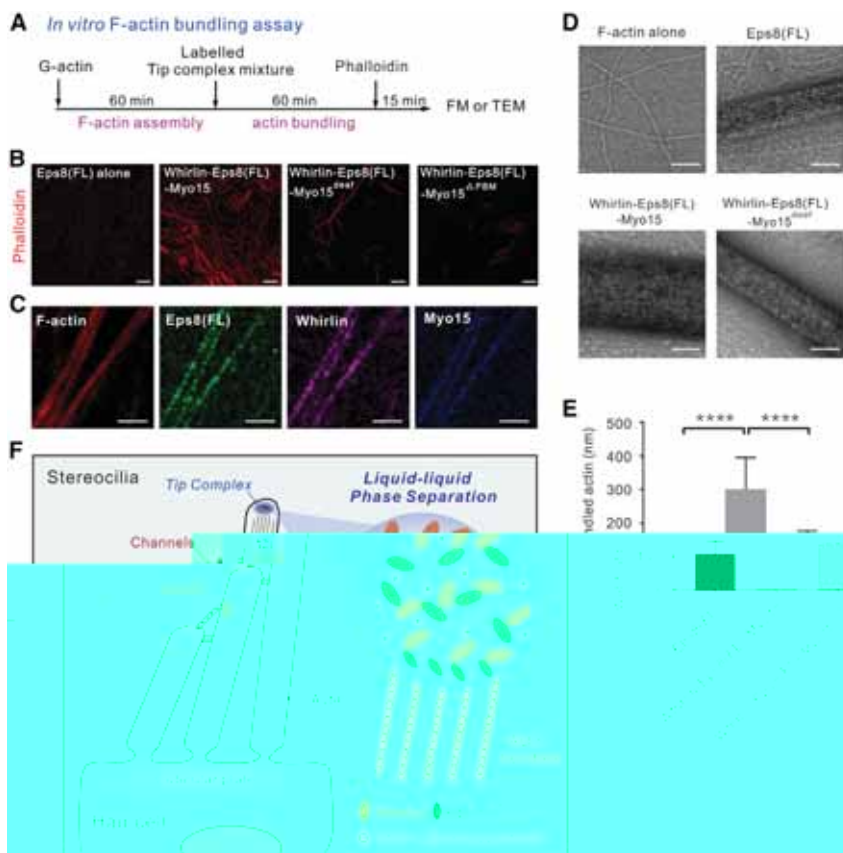


Figure 6. TCD-like condensates promote actin bundling

(A) A diagram showing the process of actin bundling experiment *in vitro*. FM, fluorescent microscopy; TEM, transmission electron microscopy.

(B) Representative images under FM showing that wild-type tip complex condensates significantly promote the actin bundling, whereas the mixture with Myo15^{deaf} or Myo15^{ΔPBM} could not. Scale bar: 5 μm.

(C) Representative images showing colocalization of Whirlin, Eps8, and Myo15 with F-actin bundles. Scale bar: 2.5 μm.

(D) Representative images of actin bundles induced by wild-type and mutant form of the tip complex mixture under TEM. Scale bar: 100 nm.

(E) Distribution of the width of actin bundles from the different groups of experiments. Statistics were performed by 2-tailed Student's t test. ****p < 0.0001.

(F) Proposed model for the tip density assembly mediated by phase separation of the tip complex.

The tip complex including but not limited 60del5-30193(th)TJ-1.0 tu

uncovered the molecular composition of these membrane-less electron densities. The TCD consists of various densely packed proteins (the tip complex), including scaffold proteins (e.g., Whirlin), molecular motors (e.g., Myo15, Myo3, Myo1), and actin regulatory proteins (e.g., Eps8, Espin) (Gillespie and Cyr, 2004; Salles et al., 2009). Recently, the polarity proteins LGN and Gαi were shown to form an ~200 nm nanodomain with Myo15 and Whirlin at the tips of the stereocilia (Mauriac et al., 2017; Tadenev et al., 2019). These proteins work together to regulate actin dynamics and stereocilia morphology. However, it remains unclear how this protein-dense semi-enclosed compartment is organized. Emerging evidence suggests that LLPS may be a general mechanism underlying the formation of membrane-less subcellular electron-dense plaques with broad physiological functions (Barnani et al., 2017; Brangwynne et al., 2009; Hyman et al., 2014). For example, the PSD and the active zone in post- and presynapses of neurons, respectively, may be assembled through LLPS (Wu et al., 2019; Zeng et al., 2016, 2018).

In the present study, we demonstrate that Whirlin, Myo15, and Eps8, three major proteins in the TCD, can autonomously undergo phase separation and form highly dense but dynamic TCD-like condensates both *in vitro* and in living cells. Therefore, we provide evidence to show that the TCD in hair cell stereocilia may be formed via LLPS. Intriguingly, the Myo7A-USH1C-USH1G complex, the core components of UTLD, was recently shown to undergo phase separation as well (He et al., 2019). It

like or solid state, which is difficult to investigate at this stage. Future work is needed to address this issue. Since Eps8 alone could not induce actin bundling at lower concentrations (Figure S5), it is possible that the TCD-like condensates facilitate the actin bundling by concentrating Eps8 in the mixture. Thus, it would not be surprising that the mutants of Myo15 that impair the formation of the TCD-like condensates led to defects in actin bundling activity (Figure 6). These data indicate that the LLPS-mediated TCD condensates may play critical roles in the regulation of stereocilia morphology by concentrating actin regulatory proteins in hair cells (Figure 6F). In accordance with this conclusion, it is generally believed that LLPS is used by cells to concentrate biomolecules at specific subcellular compartments (Banani et al., 2017; Shin and Brangwynne, 2017; Zhu et al., 2020). Intriguingly, Whirlin was reported to interact with actin-free Espin, another actin crosslinking protein, at the ankle links and stereocilia tips in hair cells (Wang et al., 2012). This interaction accelerated the exchange between pools of actin-free Espin and actin-bound Espin and may indirectly weaken the cross-links of Espin with actin filaments (Wang et al., 2012). The two mechanisms used by Whirlin (Whirlin-Espin axis and Whirlin-Eps8-Myo15 axis) to regulate actin dynamics are obviously different. These data strongly suggest that Whirlin, a versatile organizer of actin dynamics, may coordinate and balance multiple signaling pathways to precisely regulate actin dynamics in different compartments of stereocilia and/or different stages of stereocilia development.

Actin-based protrusions, including stereocilia, filopodia, and microvilli, serve a huge variety of functions in eukaryotic cells (Sahin et al., 2005). When comparing the three types of actin protrusions, the most intriguing commonality is the presence of the condensed tip complex density at the distal tips of these protrusions (Sahin et al., 2005). In filopodia and microvilli, the dense matrix contains actin polymerization proteins (e.g., VASP, Formins), actin bundling proteins (e.g., Fascin, Espin, villin, Eps8), myosins, and other signaling proteins (Breitsprecher et al., 2008; Harker et al., 2019; Kerber and Cheney, 2011; Postema et al., 2018; Sahin et al., 2005; Schirenbeck et al., 2005). It will be interesting to test whether the tip density of filopodia and microvilli could also be formed by phase separation.

STAR★METHODS

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SUPPLEMENTAL INFORMATION

Supplemental Information can be found online at <https://doi.org/10.1016/j.celrep.2021.108770>.

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AUTHOR CONTRIBUTIONS

J.Z., L.L., and Y.S. designed the experiments; L.L., Y.S., and M.W. performed the experiments; L.L. and Y.S. carried out the X-ray data collection and structure determination; J.Z., L.L., Y.S., M.W., C.W., Q.L., and R.Z. analyzed the data; L.L. and J.Z. drafted the manuscript; all of the authors commented on the manuscript; and J.Z. coordinated the project.

DECLARATION OF INTERESTS

The authors declare no competing interests.

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