A complex of distal appendage–associated kinases linked to human disease regulates ciliary trafficking and stability

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Cilia biogenesis is a complex, multistep process involving the coordination of multiple cellular trafficking pathways. Despite the importance of ciliogenesis in mediating the cellular response to cues from the microenvironment, we have only a limited understanding of the regulation of cilia assembly. We previously identified Tau tubulin kinase 2 (TTBK2) as a key regulator of ciliogenesis. Here, using CRISPR kinome and bioinert identification screening, we identify the CK2 catalytic subunit CSNK2A1 as an important modulator of TTBK2 function in cilia trafficking. Superresolution microscopy reveals that CSNK2A1 is a centrosomal protein concentrated at the mother centriole and associated with the distal appendages. Csnk2a1 mutant cilia are longer than those of control cells, showing instability at the tip associated with ciliary actin cytoskeleton changes. These cilia also abnormally accumulate key cilia assembly and SHH-related proteins. De novo mutations of Csnk2a1 were recently linked to the human genetic disorder Okur-Chung neurodevelopmental syndrome (OCNDS). Consistent with the role of CSNK2A1 in cilia stability, we find that expression of OCNDS-associated Csnk2a1 variants in wild-type cells causes ciliary structural defects. Our findings provide insights into mechanisms involved in ciliary length regulation, trafficking, and stability that in turn shed light on the significance of cilia instability in human disease.

Results

CSNK2A1 Is a Modulator of TTBK2 at the Mother Centriole. To identify additional components of the TTBK2-dependent pathways that regulate ciliary assembly and/or stability, we undertook two unbiased screens: A BioID proximity-labeling screen to find proteins that physically interact with TTBK2 and might, therefore, be effectors of TTBK2 function in cilia regulation. CK2 is a widely expressed serine-threonine kinase involved in many cellular processes, including cell-cycle progression, circadian-clock regulation, and WNT signaling (16–20). Mutations in Csnk2a1 are associated with Okur-Chung neurodevelopmental syndrome (OCNDS), a dominantly inherited disorder characterized by dysmorphic facial features and neurological impairments (21–26). We show that CSNK2A1 acts in opposition to TTBK2 in establishing and maintaining ciliary structure, with knockout of Csnk2a1 partially rescuing ciliary phenotypes of Tbk2 hypomorphic mutant cells. We also demonstrate that CSNK2A1 mainly localizes to the mother centriole and is required for cilia structure and stability. Csnk2a1-knockout cells have long cilia that exhibit a variety of defects in trafficking and actin dynamics. Additionally, overexpression of Csnk2a1 variants associated with OCNDS causes ciliary structural abnormalities. Thus, we reveal a previously undescribed role for CSNK2A1 in cilia homeostasis, define a kinase network that regulates the stability of PC, and identify links between cilia integrity and human disease.

Significance

Primary cilia (PC) are sensory organelles essential for the development and maintenance of adult tissues. Accordingly, dysfunction of PC causes human disorders called ciliopathies. Hence, a thorough understanding of the molecular regulation of PC is critical. Our findings highlight CSNK2A1 as a modulator of cilia trafficking and stability, tightly related to TTBK2 function. Enriched at the centrosome, CSNK2A1 prevents abnormal accumulation of key ciliary proteins, instability at the tip, and aberrant activation of the Sonic Hedgehog pathway. Furthermore, we establish that Csnk2a1 mutations associated with Okur-Chung neurodevelopmental disorder (OCNDS) alter cilia morphology. Thus, we report a potential linkage between CSNK2A1 ciliary function and OCNDS.

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or upstream regulators of this crucial kinase (Fig. 1A); and a CRISPR loss-of-function screen to identify proteins that functionally interact with TTBK2 (Fig. 1B). In our proximity-labeling screen, we tagged TTBK2 at the N terminus with the biotin ligase BirA, mutated to be constitutively active (27). We verified that BirA*-tagged TTBK2 is active and able to perform its normal cellular function mediating cilium assembly (SI Appendix, Fig. S1A). We then inducibly expressed TTBK2-BirA* in HEK-293T cells, labeled cells with biotin, and carried out cell streptavidin affinity purification and mass spectrometry (SI Appendix, Fig. S1B and C). Using this approach, we identified 370 proteins that were enriched by twofold or more in cells expressing TTBK2-BirA* compared with the GFP-BirA* negative control (Fig. 1A and Dataset S1). Of these, 351 were statistically significant at a false discovery rate of 0.01.

In parallel, we performed a CRISPR loss-of-function kinome screen to identify additional kinases that cooperate with TTBK2 in cilium assembly. We employed a recently described Hedgehog (HH) pathway-responsive reporter, in which 8x GLI response elements drive the expression of blasticidin resistance genes (28). We expressed the reporter and a mouse-kinome small guide RNA (sgRNA) library [no. 75316, Addgene; (29)] in wild-type (WT) mouse embryonic fibroblasts (MEFs), as well as MEFs derived from hypomorphic Tbk2 mutant embryos [Tbk2<sup<noglubasa></sup> transheterozygous embryos (9) (SI Appendix, Fig. S2A)]. We defined a hit in our screen as any sgRNAs that became enriched within the population following blasticidin selection in the Tbk2<sup<noglubasa></sup> cells but not in the WT MEFs. In other words, hits were sgRNAs that specifically improve the responsiveness of cells to HH pathway activation when TTBK2 activity is impaired. This approach would predominantly identify negative regulators of TTBK2 function either in cilium initiation or stability or both (Fig. 1B, SI Appendix, Fig. S2B, and Dataset S2).

CSNK2A1, the catalytic subunit of CK2, was identified as a hit in both of these screens (Fig. 1A and B). Because it was the only protein identified by both of our approaches, we prioritized CSNK2A1 for further analysis. To examine the relationship between CSNK2A1 and TTBK2, we verified that these two kinases associate during ciliogenesis. We coexpressed TTBK2-V5 with GFP or CSNK2A1-GFP in serum-starved HEK-293T cells. We detected an interaction between the two fusion proteins by co-immunoprecipitation, whereas no interaction was observed in the control pulldown (Fig. 2A). This suggests that these two kinases interact either directly, or as part of a complex with additional proteins.

TTBK2 localizes to the distal appendages of the mother centriole prior to the initiation of cilium assembly (8, 13, 30). To resolve the precise subcellular localization of CSNK2A1 and assess whether CSNK2A1 colocalizes with TTBK2 in that compartment, we performed conventional and superresolution immunofluorescence microscopy using antibodies against endogenous CSNK2A1, as well as a GFP fusion construct. With both approaches, we found that CSNK2A1 is specifically enriched at the centrosome, consistent with previous findings (31) (Fig. 2B and SI Appendix, Fig. S3). We then used three-dimensional stimulated emission depletion (STED) microscopy to examine endogenous CSNK2A1 localization relative to the centriolar core component Centrin-2 and the ciliary membrane protein ARL13B. This analysis revealed that CSNK2A1 predominantly localizes to the mother centriole at the vicinity of the axoneme (Fig. 2C). CSNK2A1 signal shows a distinctive localization pattern, wherein it surrounds the mother centriole in a barrel-like fashion (Fig. 2C). We distinguished two main loci of the CSNK2A1 signal. A symmetrical arc-like structure is found at the proximal end of the mother centriole. Fainter parallel linkers to centriolar sidewalls connect the proximal CSNK2A1 to a tighter distal locus (Fig. 2C and F). In addition, we detect a weaker signal of the kinase at the daughter centriole (Fig. 2 C, Upper).

To further characterize the localization of CSNK2A1 at the distal mother centriole, we analyzed its positioning relative to the distal appendage protein CEP164 together with γ-tubulin and acetylated α-tubulin using the structured illumination microscopy (SIM) (Fig. 2D). In a side view, we observed an identical pattern to that obtained from colabeling with Centrin-2. In a top-down

![Graph A](https://example.com/grapha.png)

**Fig. 1.** Identification of CSNK2A1 as a putative TTBK2-interacting protein. (A) Volcano plot of TTBK2-proximate proteins labeled by BioID. CEP164, TTBK2, and MAPRE1, known interactors of TTBK2, are shown in green. CSNK2A1, a candidate TTBK2-proximate protein, is highlighted in yellow. The x-axis denotes the log2 fold change of TTBK2-GFP control. In the y-axis, the significance displays the negative log10-transformed P value for each protein. The red dots show the proteins considered as hits, delimited by the selected thresholds ≥ 1.5 for the log2 fold change and statistical significance. (B) The CRISPR screen plot for Ttbk2<sup<noglubasa></sup> cells. The suppressor screen identifies potential negative modifiers of TTBK2, including CSNK2A1 (red). Each gene is plotted using the mean of medians values (defined as the two median ratios of sgRNAs counts, calculated as follows: +blasticidin(+SAG); −Blasticidin(+SAG); note that four sgRNAs represent each gene). Orange dots represent the nontargeting sgRNAs.

![Graph B](https://example.com/graphb.png)

**Fig. 2.** CSNK2A1 interacts with TTBK2 during ciliogenesis. (A) Immunoprecipitation of TTBK2-V5 with GFP-BirA*-tagged TTBK2 is active and able to perform its normal cellular function mediating cilium assembly (SI Appendix, Fig. S1A). We then inducibly expressed TTBK2-BirA* in HEK-293T cells, labeled cells with biotin, and carried out cell streptavidin affinity purification and mass spectrometry (SI Appendix, Fig. S1B and C). Using this approach, we identified 370 proteins that were enriched by twofold or more in cells expressing TTBK2-BirA* compared with the GFP-BirA* negative control (Fig. 1A and Dataset S1). Of these, 351 were statistically significant at a false discovery rate of 0.01.

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Fig. 2. CSNK2A1 is a centrosomal protein that opposes TTBK2 function. (A) TTBK2-V5 was overexpressed with GFP or CSNK2A1-GFP in HEK293T cells. GFP pulldowns were performed on total cell lysates, and Western blots were probed with antibodies to V5 and GFP tags. (B) Endogenous (Upper) and GFP-tagged (Lower, stably expressed, both magenta) CSNK2A1, costained with the centrosomal protein γ-tubulin (red) and ciliary marker ARL13B (green) in WT MEFs. (Scale bars, 1 μm.) (C) 3D STED superresolution images show the endogenous CSNK2A1 (red) surrounding the centriole core (Centrin-2 green, Upper). Lower shows CSNK2A1 signal at the vicinity of the cilium. (Scale bars, 500 nm.) (D) SIM acquisitions show that CSNK2A1 exhibits a complex basal body-associated organization. (Upper) A side view of CSNK2A1 (green) at the base of the cilium labeled with acetylated alpha-tubulin (magenta) and partially overlapping with CEP164 (red). (Lower) CSNK2A1 tightly surrounds the pericentriolar material, labeled by γ-tubulin (magenta), partially overlapping with CEP164 (Upper and Lower). (Scale bars, 500 nm.) (E) Representative two-dimensional STED images from WT MEFs, exhibiting the top view of CSNK2A1 (green), in a ring-shaped structure and colocalizing with TTBK2 (red). (Scale bar, 250 nm.) (F) Cartoon illustrating the subcellular organization of CSNK2A1 at the basal body. (G) Cilia defects in Ttbk2null/gt cells are partially rescued by Csnk2a1 knockout. Representative immunofluorescence images from cells labeled with ARL13B (Green) and γ-tubulin (red). DNA was stained with DAPI. (Scale bar, 2 μm.) Graphs show the mean percentage of ciliated cells (left graph: WT control gRNA (gRNA) n = 105; Ttbk2null/gt control gRNA n = 251; and Ttbk2null/gt Csnk2a1 gRNA n = 282 cells) and cilia length (right graph: WT control gRNA n = 105; Ttbk2null/gt control gRNA n = 48; and Ttbk2null/gt Csnk2a1 gRNA n = 58 cells). Error bars denote the SEM. Statistical comparison was performed by one-way ANOVA with Tukey’s multiple comparisons test. P values are shown on the graph.
view, we noted a ring of CSNK2A1 partially overlapping with that of CEP164 (Fig. 2 D and F). Top-view acquisition contained TTBK2 confirmed partial colocalization between CSNK2A1 and TTBK2 rings at the mother centriole (Fig. 2E) and indicates that CSNK2A1 is a centriolar distal appendage-associated protein (Fig. 2 D–F). These results show that CSNK2A1 and TTBK2 distal rings exhibit similar positioning at the distal appendages, with evidence of a more complex organization for CSNK2A1 (Fig. 2F).

We identified CSNK2A1 from a CRISPR screen for genes that modify the function of TTBK2, specifically as a gene that, when knocked out, increases the ability of hypomorphic \( \text{Tbbk}\text{mut} \) cells to activate a HH signaling response (Fig. 1B). \( \text{Tbbk}\text{mut} \) cells form cilia at a reduced frequency, and these cilia have defects in IFT protein localization as well as structural abnormalities (9). Therefore, we tested whether the loss of CSNK2A1 function is sufficient to rescue or partially rescue cilia defects seen in \( \text{Tbbk}\text{mut} \) cells. To this end, we tested five different sgRNAs against \( \text{Csnk2a1} \), from which we chose the most efficient sgRNA and selected individual clones to validate \( \text{Csnk2a1} \) knockout by Western blot and sequencing (SI Appendix, Fig. S4A). Simultaneously, we isolated a control clonal cell line, generated by a nontargeting sgRNA. Consistent with the identification of \( \text{Csnk2a1} \) in our CRISPR screen, sgRNA-mediated knockout of \( \text{Csnk2a1} \) in \( \text{Tbbk}\text{mut} \) cells partially rescued defects in cilia frequency as well as cilia length (Fig. 2G).

Loss of CSNK2A1 Perturbs Ciliary Trafficking. To further examine the requirements for \( \text{Csnk2a1} \) in cilia assembly and function, we generated \( \text{Csnk2a1} \)-knockout cell lines in WT MEFs using the CRISPR-Cas9 system. Two clonal cell lines were isolated with distinct frameshift mutations within the \( \text{Csnk2a1} \) gene (SI Appendix, Fig. S4B). The frequency of cilia formation was not affected in the absence of \( \text{Csnk2a1} \). However, \( \text{Csnk2a1} \)-depleted cells formed longer cilia with a wide range of lengths, suggesting possible ciliary instability (Fig. 3A).

Cilia formation is a multistep process with TTBK2 acting upstream of IFT complexes to trigger the final steps required for the conversion of the mother centriole into a basal body (8). Since CSNK2A1 partially opposes TTBK2 function, we evaluated IFT recruitment and localization of \( \text{Csnk2a1} \)-knockout cilia. In contrast with \( \text{Ttbk} \) mutant cells, components of the IFT-B complex, IFT81 and IFT88, are seen at abnormally high levels within \( \text{Csnk2a1} \)-knockout cilia, especially within the more distal axoneme (Fig. 3 B and D and SI Appendix, Fig. S4 C and D). We also assessed the localization and intensity of IFT140, a component of the IFT-A complex, which is implicated in retrograde transport as well as in the trafficking of specialized ciliary membrane proteins and HH pathway components (32, 33). In WT cells, IFT140 accumulates primarily at the ciliary base. In \( \text{Csnk2a1} \)-knockout cilia, we do not observe substantial changes in localization pattern; however, there is an increase in IFT140 intensity at the ciliary base and the axoneme (Fig. 3 C and D).

We postulated that the changes we observed in IFT localization may point to defects in intracellular transport. To examine the bidirectional IFT, we monitored IFT88-GFP within the cillum by live imaging and measured antero- and retrograde dynamics (Fig. 3E). Mutant cilia had a greater number of IFT88 tracks moving toward the ciliary tip, with increased velocity. Retrograde measurements did not show a significant change compared to control cilia (Fig. 3E). Altogether, these results suggest that CSNK2A1 regulates IFT levels in PC.

Transfuding of HH Pathway Components and Signaling Are Increased in \( \text{Csnk2a1} \)-Knockout Cilia. Given our observation of abnormally ciliary trafficking in \( \text{Csnk2a1} \)-knockout cells, we assessed whether the ciliary trafficking of key HH pathway components, including SMO, GLI2, and KIF7, is affected by the loss of \( \text{Csnk2a1} \). Upon activation of HH signaling, the transmembrane protein SMO becomes enriched throughout the ciliary membrane (34). GLI2, a transcriptional activator of the pathway, becomes enriched at the distal tip of cilia (35), as does the atypical kinesin KIF7, which regulates the structure of the ciliary tip compartment and participates in HH signaling (36, 37). For each of these proteins, we observed that their accumulation upon treatment with SMO agonist (SAG) was increased in \( \text{Csnk2a1} \)-knockout cilia (Fig. 4 A–C).

Increased accumulation or trafficking of HH pathway components into the cillum is often associated with changes in the transcriptional output of the pathway (38, 39). To test whether this is the case for the \( \text{Csnk2a1} \)-knockout cells, we assessed the expression of two direct transcriptional targets of the HH pathway, \( \text{Pch} \) and \( \text{Gli1} \), using qPCR. Consistent with the increased accumulation of HH pathway components within the \( \text{Csnk2a1} \)-knockout cilia, we observed higher expression levels of both \( \text{Pch} \) and \( \text{Gli1} \) upon SAG treatment in \( \text{Csnk2a1} \)-knockout cell lines relative to WT (Fig. 4D and SI Appendix, Fig. S5A). In addition, we also noted that levels of \( \text{Pch} \) are higher in the \( \text{Csnk2a1} \)-knockout cells relative to WT even in the absence of SAG, suggesting that the basal levels of HH might be higher in the absence of \( \text{Csnk2a1} \) (Fig. 4D). Altogether, these results suggest that CSNK2A1 is a negative regulator of Sonic Hedgehog (SHH) pathway.

CSNK2A1 Is Required for Cilia Stability. Because \( \text{Csnk2a1} \)-knockout cilia are elongated and exhibit a variety of trafficking defects, we further examined the stability of these cilia. \( \text{Csnk2a1} \)-depleted cells display a higher percentage of cilia with a tip gap that is positive for ARL13B but negative for acetylated α-tubulin (Fig. 5A). This gap, in addition to the accumulation of IFT81, GLI2, and KIF7 in mutant cilia, points to abnormalities in the ciliary tip compartment. To test whether defects at the ciliary tip were associated with decreased ciliary stability, we incubated the cells with nocodazole (Noco) to induce microtubule depolymerization and to challenge the axonomial integrity of the ciliary. We then calculated the ciliary length difference between the Noco-treated and the Noco-untreated conditions and noted that mutant cilia are less stable than control cilia (Fig. 5B). Taken together, these data indicate that \( \text{Csnk2a1} \)-mutant cilia have an abnormal ciliary tip compartment and are unstable.

The ciliary tip is also a site of ectocytosis, a process by which vesicles are shed from the cillum. This process is linked to signaling from PC, and regulation of ectocytosis is important for controlling the ciliary length and for maintaining stability (40–44). To test whether ciliary scission and ectocytosis are affected in the absence of CSNK2A1, we purified extracellular vesicles (EVs) from supernatants collected from serum-starved control and \( \text{Csnk2a1} \)-knockout cells. Nonciliated cells can shed EVs (45), however, under our conditions, EVs are enriched for ciliary proteins, suggesting shedding through tip scission (41). We noted that among the supernatants, we observed higher levels of FLOTILLIN, a marker of EVs, as well as IFT88 and ARL13B in the \( \text{Csnk2a1} \)-knockout cells relative to control cells WT cells (Fig. 5C). \( \text{Ttbk2} \) null cells, which do not form PC (8), have FLOTILLIN levels in the supernatant at levels comparable to WT cells; however, we observed a dramatic reduction of ciliary-associated proteins recovered in this fraction. Additionally, ciliary tip markers GLI2 and IFT81 were also found specifically enriched in supernatants from \( \text{Csnk2a1} \)-knockout cells (SI Appendix, Fig. S5B). Taken together, these data suggest that \( \text{Csnk2a1} \)-mutant cilia exhibit an increased amount of ciliary scission and/or EV formation.

To extend these findings and examine how the loss of CSNK2A1 might affect cilia stability through ciliary scission, we performed live imaging of cilia in MEFs derived from a transgenic mouse line expressing ARL13B-mCherry (46) and transduced with previously described lentiviruses coding for control or \( \text{Csnk2a1} \) sgRNAs. After selection, we validated by immunofluorescence a 60% depletion of CSNK2A1. \( \text{Csnk2a1} \)-depleted cilia frequently showed a thinning of the distal axoneme followed by breakage of the ciliary tip. Compared to control cells, mutant cilia showed an average increase of
CSN2A1 regulates cilia length and IFT trafficking. (A) Csnk2a1 knockout cells form longer cilia. Representative immunofluorescence images stained for ARL13B (green) and γ-tubulin (red), from control and Csnk2a1 KO-1 cells as indicated. Graphs show the mean percentage of ciliated cells and the average cilia length ± SEM. Statistical comparison was performed using the nonparametric Mann–Whitney U test. P values are displayed on the graph (ns, not significant). The total number of cells analyzed for both graphs is 69 cells per condition. (Scale bar, 2 μm.) (B) IFT81 accumulates at the basal body and cilia in Csnk2a1 KO-1 cells. Representative immunofluorescence images labeled with ARL13B (green), Csnk2a1 gRNA (red), and γ-tubulin (magenta). Graphs show the mean intensity of IFT81 at the basal body (left graph) and the axoneme (right graph) in control and Csnk2a1 KO-1 cells. Statistical comparison was performed using the nonparametric Mann–Whitney U test. P values are shown on the graph (ns, not significant). Number of cilia quantified is as follows: anterograde transport, control gRNA = 29 cells; right graph, control gRNA = 35 cells. (C) IFT88 accumulates at the basal body and cilia in Csnk2a1 KO-1 cells. Representative immunofluorescence images stained with ARL13B (green), γ-tubulin (red), and IFT88-GFP (magenta). Graphs show the mean intensity of IFT88-GFP at the basal body (left graph) and the axoneme (right graph) in control and Csnk2a1 KO-1 cells. Statistical comparison was performed using the nonparametric Mann–Whitney U test. P values are shown on the graph (ns, not significant). Left graph, control gRNA = 28 and Csnk2a1 gRNA n = 29 cells; right graph, control gRNA n = 30 and Csnk2a1 gRNA n = 35 cells. (D) The anterograde trafficking of IFT88-GFP, but not the retrograde, is increased in Csnk2a1 KO-1 cells. Representative kymographs of IFT88-GFP for anterograde (red) and retrograde (green) trafficking are shown from the base to the ciliary tip for a duration of 30 s. Graphs exhibit the mean number of IFT88-GFP events per cilium and their velocities. Error bars denote the SEM. Statistical comparison was performed using the nonparametric Mann–Whitney U test. P values are shown on the graph (ns, not significant). Number of cilia quantified is as follows: anterograde transport, control gRNA n = 18 and Csnk2a1 gRNA n = 15 cilia; retrograde transport, control gRNA n = 15 and Csnk2a1 gRNA n = 13 cilia.
Control gRNA

Csnk2a1 gRNA

+ SAG

A

B

C

D

-5-fold in the frequency of tip breakage/scission (Fig. S5 and Movies S1 and S2).

CSNK2A1 Regulates Actin Modulators and Ciliary Actin Dynamics. Previously, actin and actin-binding proteins have been linked to ciliary scission (41, 42). We therefore examined whether the ciliary breaks observed in the Csnk2a1-knockout cells were driven by changes in the actin cytoskeleton within the PC. Live imaging of cells expressing SSTR3-GFP to label cilia and Lifeact-mCherry (F-actin) showed that F-actin was transiently recruited to the breakage site in Csnk2a1-knockout cilia, prior to the excision event (Fig. S4 and Movies S3 and S4). Similarly, induction of cilia disassembly by serum readdition led to a distal enrichment of F-actin in Csnk2a1-knockout cilia (SI Appendix, Fig. S5C). These results indicate that CSNK2A1 is required for the integrity of the ciliary tip and that the protein depletion causes spontaneous ciliary tip excision, likely driven by changes in actin cytoskeleton dynamics.

Several studies have reported that CSNK2A1 regulates actin cytoskeleton through actin modulators such as PLEKHO1 (pleckstrin homology domain containing O1), cortactin, and capping proteins (47-50). To investigate how CSNK2A1 present at the distal appendages regulates the ciliary tip stability, we examined the localization of PLEKHO1 and CAPZB (F-actin-capping protein subunit beta) in control and Csnk2a1-knockout cilia. PLEKHO1 is a bona fide interactor of CK2 (49), and both proteins are reported to inhibit the activity of actin-capping proteins at the barbed ends, thereby promoting polymerization of branched actin (48). In control cilia, there was no detectable signal of PLEKHO1 and patched transcripts in Csnk2a1 KO-2 cells. Graphs were produced from three replicates per condition from two independent experiments. Statistical comparison was performed by two-way analysis ANOVA with Tukey’s multiple comparisons test. P values are shown on the graph (ns, not significant).
CSNK2A1 is essential for tip stability. (A) A tip gap is a tip positive for ARL13B and negative for acetylated alpha-tubulin (illustrated by “a”). (Scale bars, 1 μm.) Graph shows the mean percentage of cilia that display tip gaps ± SEM (47 control cilia and 49 mutant cilia). (B) Under Noco treatment, mutant cilia are less stable than control cilia. Cells were serum starved for 48 h and then treated with 10 μM Noco or dimethyl sulfoxide (DMSO) as a control for 35 min. Cilia length was collected and data were displayed as a mean difference of cilia length [+Noco]−[−Noco] ± SEM. Measurements were produced from 74 control cilia and 70 Csnk2a1 mutant cilia. (C) Csnk2a1-knockout cells shed more ciliary EVs. Western blots from total cell lysates (TCL, left blots) and EVs purifications (right blots) are probed with antibodies to FLOTILLIN-1, ARL13B, and IFT88. For each probe, the fold change value was calculated as follows: ratio of [EVs_Area under the curve]/[TCL_Area under the curve]. The fold change was normalized to the control value. Dashed line marks where the Western blot images were cut to remove experimental conditions that were not relevant to the current study. (D) Csnk2a1-knockout cilia are prone to tip breakage. Nonclonal control and Csnk2a1-depleted cells were generated in transgenic ARL13B-mCherry MEFs. Cilia were imaged every 15 min for 75 min. Breakage events were quantified and normalized to the total number of imaged cilia. The graph represents the mean percentage of tip breakage ± SEM for control (n = 51 cilia) and Csnk2a1-depleted cells (n = 63 cilia). (Scale bars, 3 μm.)

Fig. 5. CSNK2A1 is essential for tip stability. (A) A tip gap is a tip positive for ARL13B and negative for acetylated alpha-tubulin (illustrated by “a”). (Scale bars, 1 μm.) Graph shows the mean percentage of cilia that display tip gaps ± SEM (47 control cilia and 49 mutant cilia). (B) Under Noco treatment, mutant cilia are less stable than control cilia. Cells were serum starved for 48 h and then treated with 10 μM Noco or dimethyl sulfoxide (DMSO) as a control for 35 min. Cilia length was collected and data were displayed as a mean difference of cilia length [+Noco]−[−Noco] ± SEM. Measurements were produced from 74 control cilia and 70 Csnk2a1 mutant cilia. (C) Csnk2a1-knockout cells shed more ciliary EVs. Western blots from total cell lysates (TCL, left blots) and EVs purifications (right blots) are probed with antibodies to FLOTILLIN-1, ARL13B, and IFT88. For each probe, the fold change value was calculated as follows: ratio of [EVs_Area under the curve]/[TCL_Area under the curve]. The fold change was normalized to the control value. Dashed line marks where the Western blot images were cut to remove experimental conditions that were not relevant to the current study. (D) Csnk2a1-knockout cilia are prone to tip breakage. Nonclonal control and Csnk2a1-depleted cells were generated in transgenic ARL13B-mCherry MEFs. Cilia were imaged every 15 min for 75 min. Breakage events were quantified and normalized to the total number of imaged cilia. The graph represents the mean percentage of tip breakage ± SEM for control (n = 51 cilia) and Csnk2a1-depleted cells (n = 63 cilia). (Scale bars, 3 μm.)

OCNDS-Associated Csnk2a1 Mutations Cause Defects in Cilia Morphology. Dominant mutations in Csnk2a1 are associated with the human genetic disorder OCNDS (21–26). Because there is some phenotypic overlap between OCNDS and ciliopathies, and because we have shown that Csnk2a1 is required for normal ciliary trafficking and structure, we tested whether the OCNDS mutations are associated with ciliary defects. We stably expressed mouse CSNK2A1WT as well as three different OCNDS-associated variants: R80H, D156H, and R191Q (Fig. 7A) (22, 24, 52) in WT MEFs. GFP tagged-CSNK2A1WT localizes to the centrosome, similar to the endogenous protein (Fig. 2B). Most Csnk2a1 variants were still recruited to the centrosome, except for CSNK2A1R80H, which exhibited impaired centrosomal recruitment (Fig. 7B). We assessed cilia frequency and found no significant difference between these cell lines (Fig. 7C). Because the primary phenotypes of Csnk2a1-knockout cilia are related to cilia stability, we examined cilia morphology and consistently found between the three OCNDS variants a higher frequency of cilia with morphological abnormalities compared to WT CSNK2A1 (Fig. 7D). About 70% of these abnormalities include a bifurcated ciliary tip and/or membranous branch-like protrusions from the distal axoneme. An additional 30% of these abnormal cilia showed a collapsed morphology, where ARL13B-positive cilia exhibited nonelongated

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bulgy and hollow-curled structures (Fig. 7D). Our results indicate that the OCNDS-associated mutations may alter the morphology and structure of PC. Our work, therefore, establishes a link between Csnk2a1 and cilia regulation and raises the possibility that OCNDS might be linked, at least partially, to disrupted cilia structure or function.

**Discussion**

Our present work identifies the CK2 catalytic subunit CSNK2A1 as a negative regulator of TTBK2 and the SHH pathway. CSNK2A1 opposes TTBK2 function mediating cilia structure and trafficking. Our results reveal that Csnk2a1 and TTBK2 physically and functionally interact and colocalize at the distal appendages. In the absence of CSNK2A1, mutant cilia are longer and exhibit instability at the ciliary tip with significant changes in actin cytoskeleton dynamics within the cilium. These defects are accompanied by increased antherograde trafficking and accumulation of IFT and SHH-related proteins within the cilium (Fig. 8). In addition, OCNDS-associated mutations to Csnk2a1 cause ciliary abnormalities, suggesting this human disorder may be linked to ciliary dysfunction. CK2 plays a key role in numerous cellular processes, including signal transduction, cell-cycle progression, and apoptosis (53–57). The CK2 holoenzyme is generally comprised of two catalytic subunits (CSNK2A1 and/or CSNK2A2) and two regulatory CSNK2B subunits (15, 58, 59). It is not yet clear whether the role we have identified for CSNK2A1 in ciliary stability and trafficking is an independent function of that subunit or whether it represents a requirement for the CK2 holoenzyme. Several reports showed that CK2 and the catalytic subunit CSNK2A1 can localize differentially. For example, CK2 was found associated with the Golgi complex and the endoplasmic reticulum (60), whereas Csnk2a1 (61) and Csnk2a2, but not Csnk2b, were detected at the centrosome (31). CSNK2A1 can also autonomously interact and/or phosphorylate an additional set of proteins, including PP2A, CALMODULIN, and MDM2 (62–65). On the other hand, both Csnk2a1 and Csnk2b were identified as potential negative regulators of the SHH pathway in a cell-based screen (66). Thus, the role of the regulatory subunit warrants further study in the context of PC function.

Our imaging reveals a complex localization pattern for Csnk2a1 at the centrosome, in particular, the cage-like organization observed from side views. A comparable structure was observed with the secreted metalloprotease ADAMTS9 (67), which is required for ciliogenesis. The ADAMTS9 signal is similar but not identical to that of Csnk2a1, with a larger and less-organized structure surrounding the centrosome (67). Csnk2a1 also forms a ring structure at the distal appendages, which is a more common centriolar structure similar to...
and colocalizing with TTBK2 and CEP164 (Fig. 2 D and E). A fainter signal of Csnk2a1 was also detected at the daughter centriole. Altogether, Csnk2a1 barrel-like localization needs further examination to uncover its role in ciliary function.

Several studies reported that CK2 regulates actin cytoskeleton dynamics (47, 48, 68, 69). Our data showed that Csnk2a1 mediates the ciliary tip stability, likely by regulating the localization of actin modulators such as PLEKHO1 and CAPZB, respectively, at the cilium and the centrosome (Fig. 6 B and C). PLEKHO1 becomes enriched at the ciliary tip in Csnk2a1-knockout cilia (Fig. 6B). Expression of PLEKHO1 is reportedly sufficient to increase F-actin levels (48). Thus, the presence of PLEKHO1 at the distal tip of mutant cilia could trigger the formation of transient actin filaments surrounding the ciliary axoneme (72). The centrosome itself also appears to function as an actin-organizing center capable of promoting the assembly of actin filaments at the vicinity of the ciliary rootlet (73). Additionally, the kinase-dead mutation of Csnk2a1 displayed a dominant-negative effect on the ciliary localization of PLEKHO1 but not that of CAPZB, which indicates a possible implication of Csnk2a1 kinase activity on regulating PLEKHO1 through phosphorylation. Altogether, our findings suggest that Csnk2a1 controls the dynamics of actin modulators within the cilium-centrosome compartment in order to inhibit the aberrant formation of ciliary F-actin (Fig. 8).

In addition to knockout of Csnk2a1 causing ciliary defects, we show that stable expression of variants associated with the human disorder OCNDS is sufficient to perturb cilia morphology and structure. OCNDS-associated missense mutations occur throughout the kinase domain and are predicted to be deleterious to the kinase activity (9). Our data show that the overexpression of the OCNDS mutation located at the active site is not sufficient to affect either PLEKHO1 or CAPZB localization, contrary to the kinase-dead mutation. This suggests that the human mutation may not lead to complete loss of the kinase activity. Most of the Csnk2a1-mutated variants that we examined still localize to the centrosome (Fig. 7B). Thus, these may interfere and/or compete with the endogenous Csnk2a1 WT, affecting its overall activity and/or function. An additional possibility is that mutations to Csnk2a1 could alter its protein–protein interaction networks, leading to branched and collapsed ciliary phenotypes we observed.

In summary, our present work identifies a distal appendage-associated kinase module that regulates cilia stability and signaling. Csnk2a1 acts as a negative modifier of TTBK2 function and modulates the ciliary actin cytoskeleton, mediating PC trafficking and stability.
and stability. Thus, our work proposes a failsafe mechanism for ciliary tip stability mediated by CSNK2A1 by which it prevents aberrant F-actin polymerization within the cilium. Our findings also expand our knowledge of a potential implication of ciliary dysfunction in OCNDSDS. Further molecular dissection will be important to define the ciliary mechanisms/changes through which OCNDSDS...
mutations trigger clinical features and neurodevelopmental delay in patients. Examining this link in detail and in vivo is therefore of high interest to us moving forward.

Materials and Methods

Cell Culture and Cell Lines. WT, Tbk2flour[2], Tbk2null[2], and CRISPR-engineered mouse MEFs, and HEK293T cells were grown in high glucose Gibco Dulbecco’s modified Eagle medium (DMEM) with 10% fetal bovine serum (FBS) in a humidified 5% CO2 incubator at 37 °C. Cilia formation was induced by serum starving the cells using media with 0.5% FBS. WT cell lines stably expressing TTBK2-FLAG-EGFP, CSNK2A1D156H, CSNK2A1R191Q, and CSNK2A1K68M-FLAG-EGFP were maintained in growth media with 500 μg/mL of Geneticin.

 Constructs. We employed the Gateway destination vector pFLAP-Dest-EGFP-3xFLAG, where we introduced Csnk2a1WT in the C terminus of EGFP, separated by a linker of 67 amino acids. The site-directed mutagenization method was utilized to generate pFLAP-Dest-CSNK2A1R191Q-EGFP, CSNK2A1D156H-FLAG-EGFP, CSNK2A1K68M-FLAG-EGFP, and CSNK2A1WT-FLAG-EGFP. The following plasmids were also used in this study: pmCherry-pmCherry-Lifeact (Plasmid no. 54491, Addgene, Addgene), pEGFPN3-SSTR3 (Plasmid no. 35623, Addgene), pCAGGS-mCherry-ARL13B (Goetz laboratory), and pCAGGS-EGFP-IT88 (Goetz laboratory). BioID constructs for TTBK2 and GFP were generated using Gateway cloning into pDEST 5 (Addgene), pEGFPN3-SSTR3 (Plasmid no. 35623, Addgene), pCAGGS-mCherry-ARL13B (Goetz laboratory), and pCAGGS-EGFP-IT88 (Goetz laboratory). BioID constructs for TTBK2 and GFP were generated using Gateway cloning into pDEST 5 Bearing the full-length CK2 γ subunit (BioID).

 Live Imaging of Cilia. Cells were seeded in 8-well Nunc Lab-Tek II chambered cover glass and serum starved for 24 h. At all times, cells were kept in a humidified chamber at 37 °C. The live imaging was performed using a Zeiss Axios Observer-Z1 widefield microscope. The Zeiss Plan-Apochromat 63x/1.4 Oil objective with the AxioVision 506 monochrome camera recorded the fluorescent signal. The Zeiss definite focus was employed to maintain cilia on focus. Time-lapse series were generated using the Fiji software.

 IFT88 Transport Measurements. Cells were cotransfected with pCAGGS-mCherry-ARL13B and pCAGGS-EGFP-IT88 and serum starved for 24 h. Cilia were live imaged every 250 ms for 30 s to generate fast time-lapse series. These image sequences were converted into an 8-bit format. The unspecific background was removed with the correction of the unwanted cilia movement. IFT88 bidirectional transport along the cilium was displayed using kymographs that were generated using a Fiji macro toolset called KymographClean 2.0 (2017).

Ciliation Rate and Fluorescence Intensity Quantification. Measurements of cilia rate and fluorescence intensity were performed from at least five randomly selected fields of cells for each condition. The fluorescence intensity of centrosonal and ciliary markers was quantified using the Fiji software. The polygon tool was chosen to select the positive pixels of the immunostained protein. We subsequently measured the mean fluorescence intensity, which integrates the total fluorescence intensity by the selected surface area.

 RT-qPCR. Control and Csnk2a1 knockout cells were plated in a 6-well plate. Cells were serum starved for 48 h and treated or not with 200 nM of SAG. Total RNA was purified using the RNeasy kit (Qiagen). From each condition, a total of 600 ng RNA was reverse transcribed to complementary DNA (cDNA) using the Script Reverse Transcription Supermix for RT-qPCR (no. 1708840). Real-time quantitative PCR was performed following the PowerUp SYBR Green (ThermoFisher) manufacturer instructions. We used the following primers: GLI1 forward 5′-TTATGGACGGACCCAGAGA-3′, reverse 5′- GAGCAGTCGTTTGTCTTAA-3′; PTH1 forward 5′- TGACAAAGGCGCTAATCGTC-3′, reverse 5′- AGGCTACTGCTGGCCTCT-3′; and ACTIN forward 5′- TGCTCCTAGCACCAGTA-3′, reverse 5′- CCACGATCCACACAGA-3′.

Statistical Analysis. Data are reported as arithmetic means ± SEM. Statistical analyses were performed using the nonparametric Mann–Whitney U test with GraphPad Prism 8 software. For multiple comparisons, we executed a one- or two-way analysis ANOVA with Tukey’s multiple comparisons test. To analyze the intension profile data, we implemented multiple t tests (one per row), found in the grouped analyses tab in GraphPad Prism 8 software. P < 0.05 was used as the cutoff for statistical significance.

Additional materials and methods are described in detail in SI Appendix.

Data Availability. All study data are included in the article and/or supporting information.

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