The Ciliary Protein Nephrocystin-4 Translocates the Canonical Wnt Regulator Jade-1 to the Nucleus to Negatively Regulate β-Catenin Signaling*§

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Background: Deregulation of Wnt signaling contributes to the development of cystic kidney diseases such as nephronophthisis (NPH).

Results: The NPH protein NPHP4 stabilizes Jade-1, a negative Wnt regulator, and translocates Jade-1 to the nucleus.

Conclusion: NPHP4 and Jade-1 additively decrease canonical Wnt signaling.

Significance: Loss of NPHP4-mediated Wnt repression via Jade-1 may contribute to cystogenesis in NPH.

Nephronophthisis (NPH) is an autosomal-recessive cystic kidney disease and represents the most common genetic cause for end-stage renal disease in children and adolescents. It can be caused by the mutation of genes encoding for the nephrocystin proteins (NPHPs). All NPHPs localize to primary cilia, classifying this disease as a “ciliopathy.” The primary cilium is a critical regulator of several cell signaling pathways. Cystogenesis in the kidney is thought to involve overactivation of canonical Wnt signaling, which is negatively regulated by the primary cilium and several NPH proteins, although the mechanism remains unclear. Jade-1 has recently been identified as a novel ubiquitin ligase targeting the canonical Wnt downstream effector β-catenin for proteasomal degradation. Here, we identify Jade-1 as a novel component of the NPHP protein complex. Jade-1 colocalizes with NPHP1 at the transition zone of primary cilia and interacts with NPHP4. Furthermore, NPHP4 stabilizes protein levels of Jade-1 and promotes the translocation of Jade-1 to the nucleus. Finally, NPHP4 and Jade-1 additively inhibit canonical Wnt signaling, and this genetic interaction is conserved in zebrafish. The stabilization and nuclear translocation of Jade-1 by NPHP4 enhances the ability of Jade-1 to negatively regulate canonical Wnt signaling. Loss of this repressor function in nephronophthisis might be an important factor promoting Wnt activation and contributing to cyst formation.

Nephronophthisis (NPH) is the most common cause of cystic kidney disease in children, with no causative treatment currently available. The most common juvenile form of the disease manifests with urinary concentration defects in early childhood and leads to end-stage renal failure by an average of 13 years (1, 2). The disease is characterized histologically by a hallmark triad of tubular basement membrane disruption, interstitial fibrosis, and corticomedullary cyst development (1–3). Pathogenesis involves the loss of function of one or more NPHP genes, leading to the disease classification of a “ciliopathy” because all nephrocystin proteins (NPHP) studied to date localize to primary cilia or centrosomes (1, 4).

Primary cilia are assembled from the mother centriole in virtually all postmitotic eukaryotic cell types and have been associated with an increasing number of cell signaling pathways (5). An important example is the inhibition of canonical Wnt signaling by the primary cilium and basal body, which have been shown to promote proteosomal β-catenin turnover (6, 7) and sequester Wnt signaling components (8). Canonical Wnt signaling is required for cellular proliferation and differentiation during kidney development and during self-renewal of several adult tissues (9, 10) and has been implicated in disease processes such as tumor development when deregulated (10). When Wnt signaling is initiated, cytosolic β-catenin escapes degradation and is able to translocate to the nucleus, where it binds T cell factor/lymphoid enhancer factor-1 (TCF/LEF1) transcription factors and initiates Wnt-dependent gene transcription. In the absence of Wnt stimulation, cytosolic β-catenin is constitutively phosphorylated by the GSK3β-Axin2-APC destruction complex, subsequently ubiquitylated, and targeted for proteosomal degradation (10).

Consistent with the role of primary cilia in repressing canonical Wnt signaling, it has been demonstrated that the ciliary von Hippel-Lindau protein (pVHL) stabilizes Jade-1 (PHF17 isoform 1S), which, in turn, can act as an E3-ubiquitin ligase to target cytosolic as well as nuclear β-catenin for proteosomal degradation (11). Thus, Jade-1 might be able to “fine-tune”

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§ The abbreviations used are: NPH, nephronophthisis; NPHP, nephrocystin protein; TCF/LEF1, T-cell factor/lymphoid enhancer factor 1; pVHL, von Hippel-Lindau protein; qPCR, quantitative PCR; RPE-1, retinal pigmented epithelial-1; IP, immunoprecipitation; Dsh, Dishevelled; PFA, paraformaldehyde.
β-catenin levels. The loss of pVHL, as is the case in VHL disease and in the majority of renal cell carcinomas, therefore, would lead to derepressed canonical Wnt signaling via decreased Jade-1 stabilization. Loss of pVHL also disrupts ciliogenesis and leads to cystic kidneys (12–14). Taking into consideration that mouse mutants with overactive canonical Wnt signaling develop cystic kidneys (15, 16), that the protein products of the PKD genes mutated in autosomal dominant polycystic kidney disease negatively regulate Wnt signaling (17, 18), and that histological samples from patients with NPHP show abnormal Wnt up-regulation (19), an emerging model suggests that cystogenesis may in part be due to the overactivation of canonical Wnt signaling, with cilia playing a direct role in Wnt regulation (20, 21).

In line with this model, several of the cilia–associated nephrocystin proteins have also been identified as negative modulators of the canonical Wnt signaling pathway (22–24). However, reports are inconsistent regarding the contribution of NPH proteins to structural integrity of primary cilia (25–27). Furthermore, although some mouse models with a nephronophthisis phenotype present with slight changes of cilia number or length (22, 28), others retain morphologically normal cilia (29, 30). Therefore, it is likely that NPH proteins are involved with the regulation of cilia–mediated cell signaling rather than simply contributing to ciliary structure. Because Jade-1 has been identified as novel regulator of β-catenin levels (11), we questioned in this study whether Jade-1 could be a route through which the nephrocystin proteins influence Wnt signaling.

Nephrocystin-4 (NPHP4) has been shown to localize to the basal body (27, 31) and, furthermore, is thought to be involved in docking and organizing ciliary traffic at the transition zone (32, 33). Recently, NPHP4 has been reported to be a negative regulator of canonical Wnt signaling (34), acting in a similar manner to inversin (NPHP2) to regulate Dsh localization. The current study confirms that NPHP4 negatively influences canonical Wnt signaling and points to an additional mechanism of this regulation. We identify Jade-1 as a ciliary protein that interacts with NPHP4, which, in turn, facilitates Jade-1 nuclear translocation, resulting in decreased canonical Wnt signaling. We further demonstrate genetic interaction between NPHP4 and Jade-1 in zebrafish, consistent with a conserved role for NPHP4/Jade-1 in regulating canonical Wnt signaling.

**EXPERIMENTAL PROCEDURES**

*Plasmids, siRNA, and Antibodies—FLAG- or V5-tagged plasmids were generated by PCR from the fetal human kidney cDNA library (Stratagene) and inserted into a modified pcDNA6 vector (Invitrogen) using standard cloning techniques. All plasmids were verified by automated DNA sequencing. M50 Super 8 TOPFlash and M51 Super 8 FOPFlash were generated by the Moon laboratory and received from Addgene (plasmids 12456 and 12457). Renilla luciferase pGL4.74 was purchased from Promega (E6921). Control siRNA and NPH4 siRNA have been described previously (35). siRNAs purchased from Biomers (Konstanz, Germany) were directed against sequences as follows: NPHP4, 5′-AAGCGGAGCGAGATGTCTACA-3′ and scrambled/control, 5′-AAAT-GTACTGCGGTGGAGAC-3′. siRNAs purchased from Qia-

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*Cell Culture and Transfections—RPE-1 cells were cultured in DMEM-F12 medium (Sigma) supplemented with 10% FBS, 2.5 mM L-glutamate and 1.2 g/liter sodium bicarbonate. Cilia growth was stimulated by growing cells to 70–80% confluency and then serum-starving them for 24–48 h in the RPE1 medium as described above with no FBS supplement. HEK293T cells were cultured in DMEM (Sigma) supplemented with 10% FBS and seeded in 10-cm or 6-well dishes. For transfection experiments, cells were grown to 60% confluency and transiently transfected with plasmid DNA using the calcium-phosphate method as described previously (36, 37). For siRNA knockdown experiments, cells were grown to 80% confluency and transfected with Lipofectamine 2000 (Invitrogen) according to the instructions of the manufacturer. 24 h after transfection, cells in 6-well dishes were harvested in 1 ml of cold PBS, and the centrifuged pellet was boiled in Laemmli buffer at 95 °C for 10 min to obtain a whole cell lysate. Cells in 10-cm dishes were harvested with 6 ml of cold PBS, and 1 ml was immediately removed from the cell suspension to be used for “whole cell lysate” as described above. The remaining 5 ml were centrifuged, and the cell pellet used for coimmunoprecipitation.

*Reagents—Okadaic acid sodium salt was purchased from Calbiochem and resuspended in water. Cells were incubated with 1 μM okadaic acid in DMEM for 45 min. MG132 was purchased from Calbiochem and used at a concentration of 1 μM in DMEM for 4 h. Control cells were incubated with an equal amount of the solvent dimethyl sulfoxide.

*Coimmunoprecipitation—Harvested 293T cell pellets were resuspended in 1 ml of lysis buffer (1% Triton X-100, 20 mM Tris-HCl (pH 7.5), 50 mM NaCl, 50 mM NaF, 15 mM Na4P2O7, 2 mM Na3VO4, and protease inhibitors). After 15 min of incubation on ice, cells were centrifuged (20,000 g, 15 min, 4 °C), and the supernatant was ultracentrifuged (100,000 × g, 30 min, 4 °C). 50 μl of the supernatant was kept as the “IP lysate” and boiled with Laemmli buffer, and the remaining supernatant was incubated at 4 °C with anti-FLAG–agarose beads (M2, Sigma) for 1 h. The beads were washed extensively with lysis buffer, boiled in Laemmli buffer, and resolved by 10% SDS-PAGE.

*Subcellular Fractionation—Harvested 293T cells pellets were resuspended in 100 μl of hypotonic buffer containing 10 mM Hepes, 1.5 mM MgCl2, 10 mM KCl, and protease inhibitors. Cells were incubated on ice for 10 min, subsequently disrupted
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14 times using a 21-gauge needle, and then centrifuged at 4 °C at 100 × g for 30 min. The supernatant was then ultracentrifuged for 1 h at 100,000 × g to yield the cytosolic fraction, whereas the nuclear fraction was obtained by washing the pellet with 1 ml of cold PBS and centrifuged at 4 °C at 9200 × g for 10 min and boiled immediately or first resuspended in a high-salt buffer, sonicated, and repelled. Fractions were boiled in Laemmli buffer and resolved by 10% SDS-PAGE.

Luciferase Assay—HEK293T cells were seeded into 96-well plates and transfected at 80% confluency using Lipofectamine 2000 (Invitrogen) according to the instructions of the manufacturer with a total amount of 140 ng of DNA per well (50 ng TOPFlash or FOPFlash firefly luciferase; 25 ng constitutively active luciferase pGL4.74; 5 ng FLAG-β-catenin; 50 ng V5/Jade1S or empty vector; 10 ng V5/NPHP4 or empty vector). In all cases, experimental plasmids were added to a prepared master mix containing FLAG-tagged β-catenin (F.β-catenin) and the reporter plasmids. Non-stimulated cells received only the reported plasmids balanced with empty vector. For knockdown experiments, 20 nM of siRNA was incubated in Lipofectamine together with plasmids, as indicated in figure legends. After 24 h, cells were lysed with passive lysis 5× buffer (Promega). Firefly and Renilla luciferase activities were analyzed using a dual-luciferase reporter assay system (Promega) in a luminometer (Mithras LB940, Berthold) and normalized for Renilla luciferase activity to control for transfection efficiency. Each transfection was performed in triplicate, and each experiment was repeated at least three times. Data were normalized by setting the “β-catenin plus empty vector” treatment group to 100%, and significance was assessed for all β-catenin stimulated treatment groups using a one-way analysis of variance with Tukey’s post-hoc test. Error bars represent S.E. To assess the effect because of Jade-1 siRNA, data were normalized to set β-catenin stimulation to 100% separately for each Jade-1 siRNA condition. The amount that NPHP4 reduced the signal was then analyzed by normalizing so that the reduction by NPHP4 in the negative siRNA condition was set to 100%.

qPCR—HEK 293T cells transiently transfected for 24 h with the plasmids described in the figure legend were washed with PBS, and RNA was extracted using TRIzol (Invitrogen). After DNase treatment (Invitrogen), reverse transcription was performed using a high-capacity cDNA kit (Applied Biosystems). Jade-1 mRNA was assessed by SYBR Green qPCR using the following primers (Jade-1 forward primer, 5’-TGAAGCCACCCCCGTACGGGA-3’ and Jade-1 reverse primer, 5’-GCGGTTGTACCAGCCCTTTGCTGC-3’), using HPRT1 as an endogenous control (HPRT1 forward primer, 5’-TCCAGCTGGCAATACACATTGGCA-3’ and HPRT1 reverse primer, 5’-GTTCCCTTTGACACGGCAAGC-3’). All qPCR experiments were performed on a real-time PCR system (7900HT, Applied Biosystems). Data were analyzed by Student’s t test, and error bars depict S.E.

Immunofluorescence—Cells grown on glass coverslips were fixed with 4% paraformaldehyde (PFA) or ice-cold methanol, as indicated in figure legends, washed with Dulbecco’s PBS, permeabilized with 0.05% Triton-X100, and blocked with 5% normal donkey serum. Primary antibodies were incubated sequentially for 1 h each with anti-Jade1 used first, except when combined with endogenous NPHP4 staining, in which case anti-NPHP4 was used first. Secondary antibodies were incubated for 30 min. Coverslips were washed with DPBS between each step and extensively before mounting onto glass slides using Prolong Gold antifade reagent with DAPI (Invitrogen) mounting medium. Cells were visualized using an Axiovert 200 microscope (objective EC Plan Neofluar ×40/1.3 Oil) equipped with an AxioCam MRm and apotome system (Carl Zeiss Micro-Imaging, Jena, Germany). Images were captured using Axiovision 4.8 (Carl Zeiss MicroImaging).

Zebrafish—Morpholinos were obtained from Gene Tools (Philomath, OR) with sequences as follows: ATG-targeting NPHP4 (5’-GCGGTTTCCACCTCACTACGAG-3’), ATG-targeting Jade-1 (5’-TGGGGACACGGCTCCTCACTCC-3’), and 5’UTR-targeting Jade-1 (5’-GTGGGCTCCTCAAGTGATCCGGAAT-3’) and were diluted in Danieau’s buffer (58). Wild-type zebrafish embryos were injected at the 1- to 2-cell stage with 1.5 nl of the morpholino solution at 20 μM (low dose) or 100 μM (high dose). Embryos were kept in E3 buffer (5 mM NaCl, 0.17 mM KCl, 0.33 mM CaCl2, 0.33 mM MgSO4, 0.00001% methylene blue) at 28 °C and after 72 h were analyzed for body curvature. Only embryos displaying a distinct curvature of the tail angled more than 45° from the head-neck axis were scored as positive. Over 150 fish were counted for each data point. Data for low-dose comparisons were collected from three independent experiments per condition, and average phenotype frequencies and standard errors were calculated (Excel software). Significance was assessed on the basis of percent WT fish using a one-way analysis of variance with Tukey’s post-hoc test. Error bars represent S.E. Cystic morphants used for sectioning were PFA-fixed and embedded in paraffin. Sections were stained using H&E.

RESULTS

Endogenous Jade-1 Localizes to the Ciliary Base and Interacts with NPHP4—Primary cilia are critical regulators of canonical Wnt signaling (6–8). As Jade-1 is a recently identified Wnt regulator that interacts with the cilia-associated protein pVHL (11), we investigated whether Jade-1 might also localize to cilia or to the ciliary base. In previous studies, Jade-1 was found to be localized both in the cytosol and nucleus, excluding nucleoli (11), which is consistent with our data. In addition, in polarized ciliated retinal pigmented epithelial (RPE-1) cells, we found that Jade-1 primarily localized to the nucleus and perinuclear region, as well as being concentrated at the ciliary base and at times visualized within the cilium proper (Fig. 1A). Some cells did not have a concentration of Jade-1 in the nucleus, which suggests that Jade-1 is a dynamic protein, possibly shuttled in and out of the nucleus under yet to be defined conditions. To confirm the localization of Jade-1 at the centrosome, we costained using appropriate markers and a second Jade-1 antibody in both HEK 293T cells and RPE-1 cells. In both cell types, we again noted a predominantly nuclear staining pattern together with an accumulation at the centrosome, which colabeled with the centrosomal marker γ-tubulin in both dividing and interphase 293T cells (Fig. 1B), and was located at the basal body region at the base of the cilia in RPE-1 cells (Fig. 1C). When the microtubule network was dissociated after 30 min of
incubation on ice, Jade-1 could still be identified at the ciliary base (supplemental Fig. 1). This provides evidence that Jade-1 is stably retained at the centrosome, which suggests Jade-1 to be a bona fide centriole-associated protein (38). Further analysis revealed that endogenous Jade-1 localizes both to the basal body and to the transition zone, where it always colocalized with the transition zone marker NPHP1 (supplemental Fig. 2A) and was often also observed in the cillum (supplemental Fig. 2B).

These results demonstrated that the Wnt regulator Jade-1 localizes to the basal body of primary cilia where it is in close proximity to NPHP1, a central component of the NPH protein complex. We therefore investigated whether any of the NPH proteins that have also been described to localize to the ciliary base coprecipitated with Jade-1. Immunoprecipitation (IP) experiments demonstrated that NPHP4 specifically precipitated with Jade-1. V5-tagged Jade-1 (V5.Jade-1) was expressed together with FLAG-tagged NPHP4 (F.NPHP4) or a control protein (F.EPS1-225). V5.Jade-1 was coimmunoprecipitated by F.NPHP4 but not F.EPS1-225. Note that in the cytosolic IP lysates, Jade-1 levels were decreased by NPHP4, whereas in whole cell lysates taken from the same experimental culture dish prior to the IP procedure, Jade-1 levels were increased by NPHP4. Endogenous tubulin staining from whole cell lysates shows equal amounts. \( \beta \), immunoblot. \( \beta \), in the reverse experiment, 293T cells were transiently transfected with V5.NPHP4 and either F.Jade-1 or F.EPS1-225. Immunoprecipitation was performed using an anti-FLAG antibody. F.Jade-1 but not F.EPS1-225 coimmunoprecipitated V5.NPHP4. C, 293T cells were transiently transfected with F.NPHP4 or a control protein. Immunoprecipitation was performed using an anti-FLAG antibody. Endogenous Jade-1 was detected in the precipitates of F.NPHP4 but not in the control lane. The expression of Jade-1 was found to be equal in the lysates. HC, heavy chain. Blots are representative from three experiments.
endogenous NPHP4 demonstrated a perinuclear localization in RPE-1 cells (supplemental Fig. 3A, left panel), which is in accordance with previous reports (31). Cells with a perinuclear concentration of endogenous Jade-1 showed colocalization with NPHP4 in this region (supplemental Fig. 3A). NPHP4 is furthermore known to colocalize with NPHP1 in the transition zone of the primary cilium, and it could also be identified in the cilium together with endogenous Jade-1 (supplemental Fig. 3B).

**NPHP4 Stabilizes Jade-1**—To confirm that NPHP4 overexpression specifically increased the total protein level of Jade-1, a whole cell lysate using V5.GFP coexpressed with V5.Jade-1 as a transfection control was performed. This experiment demonstrated that F.NPHP4 increased the overall abundance of V5.Jade-1 (Fig. 3A) without affecting V5.GFP expression. F.NPHP4 did not alter the transcription rate of Jade-1, as assessed by qPCR analysis (Fig. 3B). Instead, NPHP4 appears to increase Jade-1 protein levels by blocking proteasomal degradation, as evidenced by a lack of additional Jade-1 protein accumulation in the presence of both NPHP4 and the proteasome inhibitor MG132 (Fig. 3C). Interestingly, NPHP4 predominantly enhanced the abundance of a smaller band of the Jade-1 protein.

**NPHP4 Translocates Jade-1 to the Nucleus**—To address whether NPHP4 promotes the nuclear localization of Jade-1, subcellular fractionation was performed following transient transfection of V5.Jade-1 with either V5.NPHP4 or a control protein. These experiments confirmed that expression of V5.NPHP4 reduced the amount of V5.Jade-1 in the cytosol but increased the amount of V5.Jade-1 detected in the nucleus as compared with V5.Jade-1 coexpressed with a control protein (Fig. 4A). The NPHP4-induced nuclear shift of Jade-1 was further supported in immunofluorescence experiments. We expressed low levels of V5.Jade-1 together with F.NPHP4 and observed low levels of nuclear Jade-1 in cells transfected only with V5.Jade-1 but a greatly increased nuclear signal in cells cotransfected with F.NPHP4 (Fig. 4B). It is noteworthy that after cell fraction, overexpressed NPHP4 predominantly associated with the nuclear fraction, in stark contrast to the cytosolic control protein V5.EPS (Fig. 4A), whereas immunofluorescence data reveals that F.NPHP4 does not actually enter the nucleus (B, center panel).

In our experiments, V5.Jade-1 presents as a double band in whole cell lysates. In the absence of exogenous NPHP4, we generally see the smaller band of Jade-1 in the cytosolic cell frac-
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Jade-1 appears consistently as a double band in immunoblot analyses of whole cell lysates, and it is the smaller of these that appears to be specifically influenced by NPHP4. We demonstrated that NPHP4 and Jade-1 are involved in a common pathway negatively regulating canonical Wnt signaling.

DISCUSSION

Recent work has demonstrated NPHP4 to be, like many of the nephrocystin proteins, a negative regulator of canonical Wnt signaling (34). This work corroborates this finding and points to a novel mechanism downstream from that suggested previously. Our study identifies Jade-1 as a novel component of the NPH protein complex being regulated by NPHP4. Jade-1, which functions as a single-subunit E3-ubiquitin ligase targeting the downstream canonical Wnt activator β-catenin to the proteasome, modulates both cytosolic and nuclear β-catenin levels and was suggested to fine-tune canonical Wnt stimulation (11). Here, we show that NPHP4 not only stabilizes Jade-1 but shifts the localization of Jade-1 from the cytosol to the nucleus, enhancing the ability of Jade-1 to block β-catenin-induced Wnt gene activation.

We performed siRNA knockdown experiments to determine the interdependent roles of NPHP4 and Jade-1 in reducing Wnt reporter activity. Knocking down endogenous NPHP4 led to a heightened level of β-catenin containing an S33A mutant, which mirrors previously published independent results (11). Additionally, Jade-1 consistently led to heightened reporter activity induced by the β-catenin S33A mutant, whereas body curvature prevalence was weaker after Jade-1 knockdown (56.5% prevalence) using the same ATG-targeting morpholino dose. However, when the injection doses were individually titrated to yield almost no body curvature phenotype using either morpholino, the combination of NPHP4 and Jade-1 knockdown at this dosage yielded a body curvature phenotype in 92.0% of embryos, whereas body curvature prevalence was weaker after Jade-1 knockdown (31.5%, indicating that NPHP4 and Jade-1 are involved in the same functional pathway in vivo (Fig. 6B). Using the Jade-1 5′UTR-targeting morpholino, we furthermore demonstrated the presence of pronephric cysts (Fig. 6C). These data clearly demonstrated genetic interaction and support a model whereby NPHP4 interacts with Jade-1 to control Jade-1 stability, nuclear translocation, and canonical Wnt signaling.
data suggest that Jade-1 may need to be in a non-phosphorylated form to enter the nucleus, where it presents as a slightly larger band and thus is likely to be rephosphorylated by nuclear kinases. NPHP4 enhances the stability of the smaller size Jade-1 protein form, resulting in increased nuclear translocation. NPHP4 does not alter the transcriptional rate of Jade-1 but
rather interacts on a protein level, possibly blocking kinase access to phosphorylation sites. This may also prevent Jade-1 from being directed to the proteasome, as the protein stabilization by NPHP4 mimics that achieved by the proteasome inhibitor MG132.

Jade-1 has been described previously to localize both to the cytosol and the nucleus (11). Our findings corroborate this and further demonstrate that Jade-1 localizes to the ciliary compartment. At the ciliary base, Jade-1 colocalizes with at least two members of the NPH protein complex and was confirmed to interact with NPHP4 by coprecipitation experiments. The factors that influence the ciliary versus centrosomal localization of Jade-1 remain to be determined. That Jade-1 is a highly dynamic protein is further exemplified by its nuclear versus cytosolic localization under basal conditions. Although we most often saw Jade-1 concentrated in the nucleus, we also identified cells within the same experiment where Jade-1 was instead mainly localized in the perinuclear region. In these cells, there was a striking colocalization with endogenous NPHP4, which also localizes perinuclearly. In our biochemical analyses, NPHP4 fractioned predominantly with the nucleus, although it is apparent using immunofluorescence that NPHP4 is not located within the nucleus itself. This is highly suggestive of NPHP4 associating with the nuclear envelope in a prime position to influence nuclear trafficking. These data are of interest in light of the recent parallels drawn between nuclear and ciliary protein transport whereby proteins required for nuclear import and export are also part of the ciliary proteome (46), supporting the possibility of a “ciliary pore” complex in analogy to the nuclear pore complex (33, 47, 48). The presence of NPHP4 at both the cilium and the perinuclear compartment may reflect parallel functions at these regions and warrants further investigation.

Both Jade-1 and NPHP4 have been demonstrated individually to inhibit canonical Wnt signaling (11, 34). Our data demonstrate that these two proteins act in concert. The additive ability of Jade-1 and NPHP4 to further reduce Wnt reporter activity would not have been seen if the two proteins acted completely independently of one another. We further elucidated individual roles in siRNA knockdown experiments. NPHP4 knockdown led to an increase in β-catenin-induced Wnt gene activity, which could be reversed by the overexpression of Jade-1. This reversal did not block Wnt activation to the extent that Jade-1 cotransfected with scrambled siRNA did. This indicates that the function of Jade-1 was reduced by NPHP4 siRNA, likely because of a decreased ability of Jade-1 to translocate to the nucleus under these conditions. Conversely, the degree to which NPHP4 is able to reduce Wnt reporter activity was impaired in the presence of Jade siRNA. However, there was substantial variation in the degree of impairment. Thus, other players are likely involved. Taken together, it is apparent that Jade-1 requires NPHP4 to be fully effective, but NPHP4 might act through several different means. Indeed, the ability of NPHP4 to regulate canonical Wnt signaling by decreasing the protein level of the canonical Wnt activator Dishevelled (Dsh) was demonstrated recently (34). β-Catenin acts downstream from Dsh in the canonical Wnt pathway. Therefore, the effect reported in our study is not likely to involve Dsh. Although the ability of NPHP4 to block Dsh-induced Wnt reporter activity could reflect a downstream effect

FIGURE 6. Jade-1 and NPHP4 additively augment an abnormal body curvature phenotype in vivo. Zebrafish embryos were injected at the 1- to 2-cell stage with either high dose (HD) or low dose (LD) morpholinos against either Jade-1 or NPHP4 and assessed at 72 hours post fertilization for body curvature. Only embryos with a body curvature of the tail angled more than 45° from the head-neck axis were scored as positive. A, representative images of zebrafish embryos injected with high doses of NPHP4 or Jade-1 morpholinos. The abnormal body curvature phenotype has been previously associated with mutations in cystic kidney disease-causing genes, ciliary abnormalities, pronephric cysts, and defective Wnt signaling (see text). B, when NPHP4 and Jade-1 ATG-targeting morpholino injections were individually titrated to a low concentration yielding no obvious phenotype individually, the combination of these two morpholinos at this low concentration resulted in a dramatic increase in the occurrence of an abnormal body curvature (F(2,8) = 25.35; p < 0.001, Tukey’s post hoc describes difference between combination and each individual condition, ** = p < 0.01). C, injection of a translation-blocking Jade-1 morpholino resulted in the development of pronephric cysts (PT, pronephric tubule; PD, pronephric duct; G, glomerulum).
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It has been proposed that the genetic mutations in ciliary “cystoproteins” can lead to an imbalance between Wnt-PCP and canonical Wnt signaling, contributing to cyst development (6, 7, 20, 34, 53–55). Many components of Wnt signaling pathways localize to primary cilia (53), and phosphorylated 

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