LRP6 exerts non-canonical effects on Wnt signaling during neural tube closure

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Low-density lipoprotein receptor related protein 6 (Lrp6) mutational effects on neurulation were examined using gain (Crooked tail, Lrp6Cd) and loss (Lrp62) of function mouse lines. Two features often associated with canonical Wnt signaling, dorsal–ventral patterning and proliferation, were no different from wild-type (WT) in the Lrp6Cd/Cd neural tube. Lrp6−/− embryos showed reduced proliferation and subtle patterning changes in the neural folds. Cell polarity defects in both Lrp6Cd/Cd and Lrp6−/− cranial folds were indicated by cell shape, centrosome displacement and failure of F-actin and GTP-RhoA accumulation at the apical surface. Mouse embryonic fibroblasts (MEFs) derived from Lrp6Cd/Cd or Lrp62/2 embryos exhibited elevated and decreased RhoA basal activity levels, respectively. While ligand-independent activation of canonical Wnt signaling, bypassing Lrp-Frizzled receptors, did not activate RhoA, non-canonical Wnt5a stimulation of RhoA activity was impaired in Lrp6−/− MEFs. RhoA inhibition exacerbated NTDs in cultured Lrp6 knockout embryos compared with WT littermates. In contrast, a ROCK inhibitor rescued Lrp6Cd/Cd embryos from NTDs. Lrp6 co-immunoprecipitated with Disheveled-associated activator of morphogenesis 1 (DAAM1), a formin promoting GEF activity in Wnt signaling. Biochemical and cell biological data revealed intracellular accumulation of Lrp6Cd protein where interaction with DAAM1 could account for observed elevated RhoA activity. Conversely, null mutation that eliminates Lrp6 interaction with DAAM1 led to lower basal RhoA activity in Lrp62/2 embryos. These results indicate that Lrp6 mediates not only canonical Wnt signaling, but can also modulate non-canonical pathways involving RhoA-dependent mechanisms to impact neurulation, possibly through intracellular complexes with DAAM1.

INTRODUCTION

Low-density lipoprotein receptor related protein 6 (Lrp6), together with Frizzled (Fzd), is an essential component of cell surface receptors for the canonical Wnt/β-catenin signaling pathway (1,2). Wnts are a family of secreted molecules that regulate numerous developmental events through several distinct signal transduction pathways, which are broadly categorized as canonical or non-canonical (3–5). The canonical pathway signals via Wnt ligand binding to Lrp5/6-Fzd1-10 heterodimers to set off a downstream cascade that ultimately results in blocking the ubiquitination and destruction of β-catenin, resulting in cytosolic accumulation and translocation of β-catenin to the nucleus where it activates TCF/Lef-dependent target gene transcription (6).

Non-canonical Wnt signaling is characterized by Wnt ligand actions transduced through Fzd and/or other receptors, such as Ryk and Ror, that do not result in β-catenin stabilization (reviewed in 5,7,8). Non-canonical effectors have been broadly grouped into either modulators of cytoskeleton remodeling (including JUN N-terminal kinase and the small GTPases, principally RhoA), or Ca2+-dependent enzymes (e.g. Ca2+/calmodulin-dependent protein kinase and protein kinase C) (5,9). Non-canonical signaling that modulates the cytoskeleton is ascribed to the planar cell polarity (PCP) pathway, because genetic mutations of the relevant genes in Drosophila,
Xenopus, Zebrafish and mice result in a loss of cell polarity within the plane of an epithelial sheet (10–14). In mice, the Wnt/PCP phenotype has been closely associated with a severe NTD, craniorachischisis, in which the neural tube becomes largely or completely open (15–17). These morphogenetic alterations require exquisitely regulated changes in cell shape via cytoskeletal reorganization (18–20). Neurulation in vertebrates requires proper regulation of RhoA signaling (21), which is disrupted in Wnt/PCP mutant mice such as Loop tail (22). Thus, there is an established link between non-canonical Wnt signaling, RhoA regulation, cytoskeletal organization and NTDs.

Historically, Lrp6 has been considered necessary for canonical Wnt signaling only, but studies also implicated Lrp6 in events associated with Wnt/PCP signaling, such as Xenopus gastrulation (23), axis elongation in Xenopus and neural tube closure in a mouse (24). However, the mechanisms by which Lrp6 achieves these functions during neurulation remain an open question. Two NTD-prone mouse lines, Lrp6
cd and Lrp6
cd, provide examples of a gain of function and a loss of function mutation in Lrp6, respectively (25,26). Crooked tail (Cd) mice carry a single amino acid substitution (G494D) in the extracellular domain of Lrp6 (25), and homozygous Lrp6
cd/cd mice display a range of phenotypes, including embryonic lethality, exencephaly (failure of cranial neural tube closure) and runted pups with severe lumbosacral and tail deformities (crooked tails). Exencephaly in Cd/Cd embryos can be prevented by prenatal folate supplementation, making it an important model of human NTDs (27). In contrast, Lrp6 null (Lrp6
cd) gene-trap mice display an embryonic phenotype primarily characterized by body axis truncation, limb defects, eye and palate defects (28). Craniofacial abnormalities, including exencephaly and/or spina bifida (failure of caudal neural tube closure) (26). Unlike the rescue from NTD in Lrp6
cd/cd embryos, prenatal folate supplementation exacerbates NTD in Lrp6
cd/cd conceptsues and leads to earlier embryonic lethality (28).

The Lrp6
cd mutation results in a protein that is resistant to Dkk1 antagonism of cytosolic β-catenin stabilization in response to Wnt3a, indicating that Lrp6
cd is a gain of function mutation (25). This lack of inhibitory function would interfere with the temporal regulation of canonical Wnt signaling and could result in net hyperactivity. However, NTD associated with Wnt pathway genes have so far been ascribed to effects on Wnt/PCP signaling (reviewed in 29). The discovery that mutations in Lrp6 are associated with NTDs suggests either that the canonical pathway also regulates neurulation, or that Lrp6 has a role outside the canonical pathway to affect cranial neurulation in mice. Here, we attempt to resolve these possibilities through comparison of Wnt pathway effects in the neural folds of Lrp6
cd and Lrp6
cd embryos. We reason that Wnt pathway functions during neurulation that are altered by both loss and gain of function Lrp6 alleles are likely to reflect Lrp6-dependent cascades that are critically important for neural tube closure.

RESULTS

NTD in Crooked tail is not explained by deregulation of the canonical Wnt pathway

Wnt1 and Wnt3a play a role in the proliferation of progenitor cells in the neural tube, where targeted gene knockout causes loss of nervous system segments and overexpression results in overgrowth (reviewed in 30). Gain and loss of function mutations in β-catenin also recapitulate these effects (31,32). In addition to cell proliferation, canonical Wnt signaling is essential for dorsal–ventral patterning of the neural tube (reviewed in 33–35). Patterning defects were previously identified in Lrp6
cd/neural tube, which displays expanded Pax3 expression at E9.5 (26). We evaluated canonical Wnt signaling activity in the neural tube of Lrp6 mutants using a TCF/LEF transcriptional reporter mouse line and also compared cell proliferation and dorsal–ventral patterning in the neural tube of mutant and wild-type (WT) siblings.

Transgenic animals carrying a LacZ cassette expressed from a TCF/LEF-responsive promoter (BatGal (36)) were crossed with the Cd line to assess canonical Wnt/β-catenin-dependent gene transcription. Embryos recovered immediately before or after cranial neurulation displayed β-galactosidase reporter activity indicating that β-catenin-dependent gene transcription in Lrp6
cd/cd::BatGal embryos was reduced in crania and unchanged in somites at E9.5, regardless of whether cranial folds were open or closed (Fig. 1A). At E8.5, during completion of neural tube closure, reporter activity was similar in both WT::BatGal and Lrp6
cd/cd::BatGal embryos. In addition, LacZ expression detected in tissue sections by immunostaining was similar to whole-mount enzymatic detection with X-Gal (Supplementary Material, Fig. S1). Canonical Wnt signaling in the presence of mutant Lrp6 was further examined in mouse embryonic fibroblast (MEF) cells derived from the Lrp6
cd/cd and Lrp6
cd embryos (Fig. 1B–D). Cultures were incubated with vehicle or recombinant Wnt3a and collected protein lysates were subjected to western blot analysis, probed with antibodies for total β-catenin and unphosphorylated (activated) β-catenin (37). Compared with littermate controls, basal levels of active β-catenin were lower in mutant MEFs and were induced to a lesser extent by Wnt3a (basal unphosphorylated β-catenin: 33.96 Lrp6
cd/cd versus 100 WT; 52.19 Lrp6
cd/cd versus 100 WT; stimulated: 84.64 Lrp6
cd/cd versus 165.51 WT; 84.04 Lrp6
cd/cd versus 373.97 WT, n = 3, P < 0.05) (Fig. 1B and C). Similar to the reporter mouse embryos, the difference between mutant and WT activated levels was substantially smaller for Lrp6
cd/cd than for Lrp6
cd MEFs (stimulated values: 84.64 Lrp6
cd/cd versus 165.51 WT compared with 84.04 Lrp6
cd/cd versus 373.97 WT, or 2-fold reduction versus 4.5-fold reduction in active WNT signaling P < 0.05). Another readout of canonical signaling, Wnt-dependent transcription of Axin2 mRNA, was compared using quantitative RT-PCR (Fig. 1D). Transcriptional activity paralleled the changes observed for activated β-catenin [basal Axin2 mRNA: 21.32 Lrp6
cd/cd versus 100 WT; 9.97 Lrp6
cd/cd versus 100 WT; stimulated: 65.58 Lrp6
cd/cd versus 319.41 WT; 31.81 Lrp6
cd/cd versus 815.91 WT, or 4.8-fold (Lrp6
cd/cd) versus 25.5-fold (Lrp6
cd/cd) reduction in active Wnt signaling n = 3, P < 0.05). Together, these data indicate that Lrp6
cd is certainly not a hyperactive allele and may be hypomorphic with respect to canonical Wnt signaling through β-catenin in the neural tube.

We next evaluated markers for the dorsals (Pax3), middle (Pax6) and ventral (Nkx2.2) portions of the neural tube of Lrp6 mutant embryos. Compared with WT siblings (Fig. 2A, E, I), the Lrp6
cd embryos showed an expansion of the Pax3 and Pax6 protein expression domains (Fig. 2C and G) [Lrp6
cd versus
WT: 60.4 ± 0.32% versus 57.2 ± 1.01% μm/μm Pax3 n(4) = 3.19; 61.6 ± 3.75% versus 52.0 ± 1.54% μm/μm Pax6 n(5) = 2.69, P < 0.05, reported as percentages in order to control for size variation among embryos]. However, the extent of patterning gene expression was no different in Lrp6Cd/Cd embryos (Fig. 2B, F, J), suggesting that altered dorsal–ventral patterning is not the primary cause of NTD in Lrp6Cd/Cd embryos.

Phospho-histone3 (PH3), an M-phase marker, was used to label dividing neural tube cells at the midbrain/hindbrain junction. In contrast to the reduced divisions in Lrp6−/− embryos [Fig. 2B, D and (28)], (9.92 ± 0.89 Lrp6−/− versus 11.88 ± 0.27 WT, PH3cells/200 μm, n(18) = 2.88, P < 0.05), no difference was observed in proliferation between WT and Lrp6Cd/Cd neural folds (Fig. 3A, B, D).

Apoptosis is also essential to neural tube closure, as disruptions of nearly a dozen genes that are either anti- or pro-apoptotic have been found to cause NTDs in mice (reviewed in 29). Cell death among genotypes was compared using LysoTracker, a reagent that allows whole-mount visualization of cells with activated lysosomes, a hallmark of apoptosis. No differences in the pattern of cell death in the neural folds were observed between WT, Lrp6Cd/Cd and Lrp6−/− (Fig. 3E–J). Collectively, the above results show that defects in canonical Wnt signaling-dependent functions of proliferation, dorsal–ventral patterning and apoptosis are not sufficient to explain the exencephaly in Lrp6Cd/Cd mutant embryos.

Apical-basal cell polarity is disrupted in crooked tail and Lrp6−/− neural folds

Cranial neurulation requires formation of a dorsal–lateral hinge point (DLHP) in the neural folds (38). Therefore, the DLHP was

Figure 1. Lrp6Cd-driven TCF/Lef transcription in situ and in vitro. (A) WT and Lrp6Cd/Cd embryos carrying the TCF/Lef-reporter transgene, BatGal, were collected at E8.5 and E9.5 and stained with X-gal to visualize canonical Wnt signaling-dependent gene transcription in whole-mount embryos. At E8.5, there is no phenotypic difference between mutant and WT sibling. Two E9.5 Lrp6Cd/Cd embryos are shown, one with open cranial folds and one that successfully completed cranial neurulation despite being homozygous for the mutation (far right panel), and therefore is phenotypically comparable with the WT sibling. (B) Western blot of Wnt3a responses measuring unphosphorylated (active) β-catenin (red channel) normalized to total β-catenin (green channel) in MEFs derived from the Lrp6Cd and Lrp6−/− mouse lines. (C) Quantification of basal and recombinant Wnt3a (rW3a) stimulated levels of activated β-catenin, in mutant compared with littermate-derived WT control MEFs. (D) Quantitative RT-PCR of Axin2 mRNA in control versus rW3a treated MEF cells. *P < 0.05, n = 3.
examined using the mutant lines intercrossed with mice that ubiquitously express myristoylated Venus fluorescent protein (39) to allow visualization of cell shape in vivo. Lrp6Cd/Cd::myr-Venus or Lrp62/2::myrVenus embryos often fail to form a DLHP (Fig. 4B and C). DLHP formation requires cells to lengthen in the apical–basal axis and narrow along the apical surface of cells in the medial–lateral axis (apical constriction), thus creating a bend in the tissue (40,41). Both Lrp6Cd/Cd and Lrp62/2 embryos showed a significant increase in the circularity of cells along the apical surface (Fig. 4D), consistent with a cell shape defect that could impair DLHP formation (WT 0.57 ± 0.003 versus Lrp6Cd/Cd 0.72 ± 0.018 t(4) = 4.83, P < 0.01 or WT 0.61 ± 0.018 versus Lrp62/2 0.70 ± 0.005 t(4) = 6.72 P < 0.01).

Despite the prior association of Lrp6 with canonical signaling, the defects in cell shape observed in Lrp6Cd/Cd and Lrp62/2 embryos showed a significant increase in the circularity of cells along the apical surface (Fig. 4D), consistent with a cell shape defect that could impair DLHP formation (WT 0.57 ± 0.003 versus Lrp6Cd/Cd 0.72 ± 0.018 t(4) = 4.83, P < 0.01 or WT 0.61 ± 0.018 versus Lrp62/2 0.70 ± 0.005 t(4) = 6.72 P < 0.01).

Figure 2. Dorsal–ventral patterning of the neural folds is normal in Lrp6Cd/Cd and modestly altered in Lrp62/2. (A–D) The linear extent of Pax3 immunolabeling is slightly expanded ventrally in Lrp62/2, but no different in Lrp6Cd/Cd compared to WT in E9.5 neural folds. (E–H) Pax6 immunolabeling is expanded dorsally in Lrp62/2, but remains unchanged in Lrp6Cd/Cd. (I–L) Localization of the ventral floorplate marker, Nkx2.2, was the same in all genotypes. *P < 0.05.
F-Actin and RhoA are deregulated in Lrp6 mutants in vivo and in vitro

During neurulation, F-actin accumulates at the apical surface (Fig. 5A, reviewed in (45)). Histological sections from Lrp6Cd/Cd and Lrp6−/− embryos showed a failure of apical F-actin accumulation in the neural folds (Fig. 5B and C). A GST-tagged Rhotekin-binding domain (RBD) peptide was applied to tissue to bind active RhoA in vivo. Compared with WT, active RhoA was generally increased in Lrp6Cd/Cd and diminished in Lrp6−/− neural folds with no accumulation at the apical surface (Fig. 5D, E, F).

RhoA activity was next measured in vitro using RBD fused to agarose beads to pull down GTP-bound RhoA (Fig. 5G and H). Lrp6Cd/Cd MEFs contained elevated GTP-RhoA levels compared with WT MEFs, whereas there was significantly less active RhoA in Lrp6−/− MEFs than WT controls (257 ± 56.5% of WT in Lrp6Cd/Cd, t(5) = 2.76, P < 0.05; 62 ± 10.8% of WT in Lrp6−/−, t(7) = −3.50, P < 0.05). Stimulation of mutant MEFs with recombinant Wnt3a, sufficient to increase GTP-RhoA 2-fold in WT cells, was ineffective in either Lrp6−/− or Lrp6Cd/Cd cells. RhoA is a known downstream target of the Wnt/PCP pathway (47,48). Myosin light chain (MLC) is a prominent target downstream of RhoA and its effector kinase, ROCK. Phospho-MLC2 is required for vertebrate neurulation (21) and is a key element of PCP signaling in the embryonic node (49) and for apical constriction during neurulation. Levels of p-MLC2 were therefore examined in NIH3T3 cells transfected with pLrp6–GFP or pLrp6Cd (Fig. 5I and J) and in lysates of WT, Lrp6Cd/Cd and Lrp6−/− MEFs (Fig. 5K and L). In both settings, p-MLC2 levels were elevated in Lrp6-expressing cells, and the phosphoprotein was reduced in Lrp6 null cells, providing strong evidence for a role of Lrp6 in RhoA-dependent, non-canonical Wnt signaling.

To further test the participation of Lrp6 in non-canonical Wnt signaling, we examined RhoA activity levels in the presence of recombinant Wnt5a, a ligand considered to act exclusively in the non-canonical Wnt pathway, likely through the Ror tyrosine kinase receptor family (8). Again, basal levels of active RhoA were elevated in Lrp6Cd/Cd and reduced in Lrp6−/− MEFs (Fig. 6A). Interestingly, the Wnt5a ligand failed to further stimulate RhoA activity in MEF cultures from either mutant line (Fig. 6A). Moreover, when the GSK3β inhibitor CHIR99021 was used to activate canonical Wnt signaling downstream of the Lrp-Fzd receptors (50), there was no activation of basal RhoA activity (Fig. 6B, Supplementary Material, Fig. S2).
Thus, whether directly or indirectly, Lrp6 has an impact on the ability of Wnt5a to regulate non-canonical signaling, while the canonical pathway by itself is insufficient to activate RhoA. Proper RhoA regulation is required for neural tube closure in Lrp6Cd/Cd and Lrp6−/− embryos

We next sought to determine whether the deregulation of RhoA observed in the neural folds is an essential mechanism underlying the NTDs in both Lrp6Cd/Cd and Lrp6−/− mutants. Embryos (n = 156 Lrp6−/− line; n = 272 Lrp6Cd line) were collected at E8.5 and were subjected to ex vivo roller culture in media containing a ROCK inhibitor to antagonize the major downstream effector of RhoA during cranial neurulation. Previous reports demonstrated a differential effect of ROCK inhibitor on convergent extension and neurulation in other mutant mice with defective PCP signaling (22). In the present study, heterozygous Lrp6+/− embryos were more susceptible to NTD in the presence of ROCK inhibitor compared with their WT littermates (Fig. 6C). In contrast, NTD in Lrp6Cd/Cd embryos, in which basal RhoA activity is elevated, were rescued by increasing doses of ROCK inhibitor (Fig. 6D). The 33% rate of neural tube closure observed for Lrp6Cd/Cd embryos in the control culture condition is consistent with previous reports of neurulation rates in Cd/Cd

Figure 4. Defects in cell shape and polarity in the cranial neural folds of both Lrp6Cd/Cd and Lrp6−/− embryos. (A–C) Sections through the cranial folds of E9 embryos carrying the myr-Venus reporter immunolabeled for GFP reveal cell shape and show that compared with WT there is a failure of cell elongation throughout the Lrp6−/− neural folds, particularly at the DLHP. (A′–C′) Higher magnification images (×63) in which individual cells have been traced (yellow) illustrate the changes in cell shape. (D) The average shape of cells along the apical surface is measured as circularity, where 1.0 reflects an exactly round measurement and numbers closer to zero reflect an increasing deviation from a perfect circle. Cells are significantly more rounded in both Lrp6Cd/Cd and Lrp6−/− neural folds (n = 3 animals per genotype, 100–150 cells per animal, *P < 0.01). (E–H) Cell polarity is disrupted in Lrp6Cd/Cd and Lrp6−/− neural folds and Cdc42 is deregulated in mutant MEFs. (E–G) Merged immunofluorescence images show cell shape (myr-Venus GFP, green) and centrosomes (γ-tubulin, red). In contrast to WT sections, centrosomes fail to accumulate γ-tubulin or position themselves at the apical surface in Lrp6Cd/Cd and Lrp6−/− neural folds. (H) Western blots from PAK-bead pulldowns of the GTP-Cdc42 from MEF cells show a decrease in the active form compared with total cdc42 levels. n = 4, *P < 0.05.
mice (27), while at 5 μM ROCK inhibitor 82% of Lrp6Cd/Cd embryos closed their neural tubes, a significant rescue (z = 2.13, P < 0.05). Furthermore, the rescued Lrp6Cd/Cd embryos show a restoration of centrosomes and their positioning at the apical pole of neural fold cells (Fig. 6E). Thus ROCK inhibition interferes with neurulation in Lrp6+/− through a mechanism that further decreases the already reduced levels of GTP-RhoA. In contrast, the Lrp6Cd/Cd mutation, associated with elevated RhoA activity, renders embryos resistant to deleterious effects of ROCK inhibitor, which instead rescues Lrp6Cd/Cd mutants from NTD and restores centrosomes to the apical pole of the neural tube.

Improper intracellular regulation of Lrp6Cd

We previously showed that the Cd point mutation impairs the interaction of Lrp6Cd with its known chaperone protein MESD, reducing levels of biotinylated Lrp6 at the cell surface (25). Subsequent studies identified Lrp6 as the target of glycosylation that changes the electrophoretic mobility of the protein so that it appears as a doublet on western blots at around 220 and 240 kDa, well above its predicted molecular weight of 185 kDa (51,52). Here, the Cd mutation is shown to produce a 220 kDa immature form of the protein, as well as a novel fragment of ~150 kDa (Fig. 7A). The latter fragment is consistent

Figure 5. Defects in cytoskeletal RhoA activity regulation in Lrp6 mutants. (A–C) Both Lrp6Cd/Cd and Lrp6−/− embryos fail to accumulate F-actin at the apical surface of the neural folds. (D–F) GTP-RhoA is elevated throughout the neural folds in Lrp6Cd/Cd. In contrast, RhoA activity in Lrp6−/− is generally reduced and GTP-RhoA fails to accumulate at the apical surface. (G, H) MEF cells isolated from Cd/Cd embryos show elevated basal levels of GTP-RhoA that cannot be further stimulated by recombinant Wnt3a (rW3a), whereas Lrp6−/− MEFs show diminished basal levels that are unresponsive to Wnt stimulation. n = 6 *P < 0.05. (I) NIH 3T3 cells transfected with pLrp6-GFP or pLrp6Cd-GFP plasmids immunostained for GFP and p-MLC2, phosphorylated on threonine residue 19. In contrast to cells expressing Lrp6−GFP (white *), those with Lrp6Cd−GFP (white arrowhead) showed visibly more p-MLC2 labeling than untransfected cells, quantified in the bar graph (J; n = 12 cells each condition *P < 0.02). (K, L) Western blots of MEF cell lysates from mutants and their WT siblings showed p-MLC2 was elevated to 175% (Lrp6Cd/Cd versus WT) and reduced to 70% (Lrp6−/− versus WT) of controls (loading normalized either to ribosomal protein S3, or to total MLC2 *P < 0.01, n = 6 each genotype).
with improper cleavage of Lrp6 occurring at the site of the Cd mutation, which would be detectable with the antibody directed against the C-terminal portion of Lrp6. Treatment of WT and Cd/Cd MEF lysates with the deglycosylating agent PNGase F caused a shift in Lrp6 protein to an ~200 kDa species (Fig. 7B, Lane 3) consistent with the removal of the post-translational glycosylation of Lrp6 reported previously (51,52). Treatment of cells with Tunicamycin to prevent glycosylation in Figure 6. Lrp6 genotype and non-canonical stimulation of RhoA signaling in NTDs. (A) RhoA activity in MEF cells derived from Lrp6<sup>Cd/Cd</sup> or Lrp6<sup>+/−</sup> and their WT littermate embryos. Cultures were stimulated for 10 min in recombinant Wnt5a. (B) WT MEFs. Basal RhoA activity levels were examined after exposing cells for 4 h to the GSK3β inhibitor, CHIR99021 (10 μM). Canonical Wnt activation independent of Lrp-Fzd does not elevate RhoA activity levels. (C and D) Whole embryo cultures of animals harvested at E8.5 and maintained ex vivo for 24 h before phenotypic scoring and genotyping. Graphs show the effects of ROCK inhibitor, Y-27632, on neural tube closure in the Lrp6<sup>−/−</sup> (C) and Lrp6<sup>Cd/Cd</sup> (D) mouse lines. (C) Heterozygous, Lrp6<sup>+/−</sup> embryos failed neurulation more frequently at lower doses of Y-27632 when compared with WT littermates, producing a leftward shift in their dose response curve. (D) The 5 μM dose of Y-27632 rescued Lrp6<sup>Cd/Cd</sup> embryos from NTD, decreasing the incidence of NTD compared with those cultured with no inhibitor. *P < 0.05. (E) Whole embryos cultured in the presence of 5 μM Y-27632. In contrast to a WT with closed cranial folds, the open cranial folds in a WT embryo show displaced γ-tubulin-labeled centrosomes, while the closed cranial folds of the Lrp6<sup>Cd/Cd</sup> embryo show more centrosomes are aligned at the apical surface.
Figure 7. Lrp6\textsuperscript{Cd} is improperly processed within the cell. (A) Western analysis of Lrp6 in MEFs shows that Lrp6\textsuperscript{Cd} fails to form the typical doublet seen in WT and instead produces only the lower isoform and a novel 150 kDa fragment. (B) Treatment of MEF cells with deglycosylating agents shifts Lrp6 bands to smaller apparent molecular weights. Lanes 1 and 2: lysates from untreated cultures prepared with or without glycosylation denaturing buffer; lane 3: lysates treated with PNGase; lane 4: lysates from cells cultured in the presence of Tunicamycin, an inhibitor of protein glycosylation. (C) Lrp6 and DAAM1 are reciprocally co-immunoprecipitated (IP) from cells overexpressing Lrp6 or Lrp6\textsuperscript{Cd} protein. Left panels are immunolabeled for DAAM1, right panels for Lrp6. (D) Transfections of NIH3T3 cells with LRP6–GFP fusion protein—with equivalent transfer efficiencies for the WT and mutant plasmids confirmed by equal total pixels immunolabeled GFP/cell in the transfected cultures—show intracellular GFP fluorescence (green) and substantial overlap (yellow) with WGA-labeled Golgi apparatus (red in upper panel). Dual labeling with anti-pan-cadherin antibody (red in lower panel) shows the whole cytoplasm and edge of the plasma membrane. When pLrp6–GFP is cotransfected with pMESD, more of the fluorescently labeled receptor gets out of the Golgi (decreased GFP/WGA overlap) and more reaches the plasma membrane (green fluorescence at the cell periphery, arrowheads). In contrast, cells expressing Lrp6\textsuperscript{Cd–GFP} display substantial overlap with labeled Golgi, even in the presence of exogenous MESD, with almost no overlap with the plasma membrane. (Blue = DAPI stained nuclei). (E) Quantified overlap of GFP-fluorescence with WGA-labeled Golgi apparatus in cells expressing either Lrp6–GFP or Lrp6\textsuperscript{Cd–GFP} with or without added MESD. *P < 0.0015 compared with WT. n = 12–18 cells per condition. (F) HEK293 cells (live, unfixed) show less mutant Lrp6\textsuperscript{Cd–GFP} (green) colocalization (yellow) with the plasma membrane (red, arrowheads) compared with Lrp6–GFP, when cotransfected with myristoylated-red fluorescent protein (pMyrRFP) and pMESD. (G) Transcriptional reporter activity in HEK293 cells transfected with pLrp6\textsuperscript{WT} or pLrp6\textsuperscript{Cd} with or without pMESD. MESD enhances transcriptional activity of Lrp6\textsuperscript{WT}, but has no effect on Lrp6\textsuperscript{Cd}-dependent transcription.

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vivo resulted in a single band in both WT and Lrp6\textsuperscript{Cd/Cd} cells, each migrating at the predicted molecular weight of Lrp6 of 185 kDa (Fig. 7B, lane 4). Absence of the putative cleavage product in Tunicamycin treated Lrp6\textsuperscript{Cd/Cd} MEFs suggests that degradation of Lrp6\textsuperscript{Cd} requires that the protein is glycosylated and that the aberrant cleavage occurs during or after post-translational processing.

A previous report provided genetic evidence for a role of Lrp6 in the regulation of small GTPases (24). However, the cellular consequences during neurulation and molecular mechanism(s) underlying its effects have not been characterized. Disheveled-associated activator of morphogenesis 1 (DAAM1) is a requisite formin protein linking PCP signaling with changes in actin skeleton organization. Changes in these features in the Lrp6 mutants are associated with altered RhoA activity levels. Moreover, RhoA activation in the presence of non-canonical Wnt pathway ligand Wnt5a is impaired in the absence of Lrp6. The data support a role for Lrp6 in the regulation of cell functions beyond β-catenin-dependent signaling.

DISCUSSION

This study examined the potential mechanisms underlying NTDs in two different mouse Lrp6 mutants. Numerous essential processes for neural tube closure that are regulated by Wnt signaling were evaluated, including β-catenin-TCF/LEF-dependent gene transcription, cell proliferation and apoptosis, dorsal–ventral patterning, establishment of apical–basal cell polarity and organization of the cytoskeleton. Changes in these features in the Lrp6 mutants are associated with altered RhoA activity levels.

Looking for evidence of canonical Wnt effects on neurulation, we found impaired proliferation and expansion of Pax3 and Pax6 expression domains in Lrp6\textsuperscript{−/−} embryos, but these were unchanged in the Lrp6\textsuperscript{Cd/Cd} embryonic neural tube. Neither mutation visibly increased apoptosis during advanced stages of neurulation.

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Looking for evidence of canonical Wnt effects on neurulation, we found impaired proliferation and expansion of Pax3 and Pax6 expression domains in Lrp6\textsuperscript{−/−} embryos, but these were unchanged in the Lrp6\textsuperscript{Cd/Cd} embryonic neural tube. Neither mutation visibly increased apoptosis during advanced stages of neurulation.

Nevertheless, the LacZ reporter mouse indicated that canonical Wnt/TCF/LEF-dependent transcription was not elevated in Lrp6\textsuperscript{Cd/Cd} embryos. This ran counter to the expected outcome given the elevated levels of cytosolic, total β-catenin in Lrp6\textsuperscript{Cd/Cd} MEF cells, increased nuclear β-catenin in Lrp6\textsuperscript{Cd/Cd} somites and lack of antagonism by Dkk1 in MEFs and transfected cells previously observed (25). Clearly, the Lrp6\textsuperscript{Cd} mutation alters the temporal aspect of Wnt signaling as, once activated, Dkk1 does not antagonize the canonical pathway (25). Attenuated elevations of activated β-catenin and Axin-2 mRNA levels in response to Wnt3a in mutant compared with WT cells, together with the inability of cotransfected MESD to enhance Lrp6\textsuperscript{Cd}-dependent transcriptional reporter activity, or to increase transit of Lrp6\textsuperscript{Cd} out of the Golgi, all suggest that Lrp6\textsuperscript{Cd} transcriptional responses in the neural tube may be blunted. Nevertheless, the present study shows that Lrp6\textsuperscript{Cd} is a gain of function mutation as it is associated with elevated RhoA activity, while Lrp6\textsuperscript{−/−} embryos display reduced RhoA activity levels and pharmacological inhibition of ROCK, the major effector downstream of RhoA, can rescue NTD and cell polarity defects in Lrp6\textsuperscript{Cd/Cd} embryos. Our in vivo data lend strong support to a growing literature demonstrating context-dependent outcomes of Wnt signaling (reviewed in 5).

The present data suggest that canonical Wnt signaling defects do not readily account for NTD in Lrp6\textsuperscript{Cd/Cd} and that Lrp6 also affects non-canonical functions that, in view of the rather modest patterning changes in the Lrp6\textsuperscript{−/−} embryos, may be more crucial to advancing neurulation.
were observed in the width of the floor plate in either Lrp6 mutants. We did not specifically examine convergence extending neurulation defects in mice closely associated with PCP defects in both mutants. Unlike mouse mutants closely associated with PCP defects, no differences were observed in the width of the floor plate in either Lrp6 mutant line. We did not specifically examine convergence extending neurulation defects in the neural tube of both the Lrp6Cd/Cd and Lrp6+/−/− embryos. The failure of cells to elongate in the neural folds could account for the improper DLHP formation in both cell. The mutant Lrp6Cd accumulates within the cell to alter levels of β-catenin and increase GTP-RhoA levels through complexation with DAAM1. In the Lrp6−/− cells, only Lrp5 remains available for signal transduction, and it is insufficient to activate either the canonical or non-canonical pathways to the level necessary for proper neural tube closure.

In contrast to the limited effects of the mutations on canonical signaling parameters in the neural tube, a number of apical–basal cell polarity deficits associated with non-canonical Wnt signaling were found in the neural tube of both the Lrp6Cd/Cd and Lrp6+/−/− embryos. The failure of cells to elongate in the neural folds could account for the improper DLHP formation in both mutants. Unlike mouse mutants closely associated with PCP defects, no differences were observed in the width of the floor plate in either Lrp6 mutant line. We did not specifically examine convergence extending neurulation defects in both Lrp6Cd/Cd and Lrp6+/−/−/− mice coincided with disruption of γ-tubulin localization in the neural folds and deficient Cdc42 activation in vitro in both cases, indicating defects in establishing apical–basal cell polarity, a prerequisite for PCP-driven events. This failed polarity was associated with disrupted network elements implicated in apical constriction (reviewed in(46,61), including altered localization of apical F-actin, and GTP-RhoA in the neural folds of both Lrp6 mutants.

GTP-RhoA levels were elevated in Lrp6Cd/Cd and decreased in Lrp6+/−/− MEFs and embryonic neural tube compared with WT. Disruption of RhoA signaling exacerbates NTDs in other mouse models associated with PCP defects (22). Here, ROCK inhibition prevented NTDs in cultured Lrp6Cd/Cd embryos while exacerbating neurulation defects in Lrp6+/−/− animals. Recent reports indicate that phosphorylation of ML2c, a target of ROCK, is controlled by the PCP pathway and that phosphorylation of ML2c is required for neural tube closure and planar polarization of the embryonic node (21,49). We further demonstrate that, compared with WT, Lrp6Cd expression in Lrp6Cd/Cd MEFs or transfected cells produces significant elevation of intracellular phospho-ML2c. Thus a connection between Lrp6 mutants, disregulated RhoA activity, altered phosphorylation of its downstream (ML2c) target through ROCK and, therefore, integrity of non-canonical Wnt signaling pathway elements is compelling. Pharmacological activation of canonical, β-catenin-dependent Wnt signaling using a compound that bypasses Lrp6/Fzd receptors did not increase RhoA activation, suggesting that changes in RhoA activity in the Lrp6 mutants are not directly related to changes in transcription. Importantly, the degree of RhoA activation in response to Wnt5a, a Wnt ligand acting independently of Lrp6 and instead dependent upon other receptors, activates RhoA to the same levels as Wnt3a, which has mixed canonical and non-canonical signaling potential (63,64). Moreover, Wnt5a treatment does not further increase the already elevated RhoA activity in Lrp6Cd/Cd MEFs, while RhoA activation by Wnt5a is impaired in Lrp6−/− MEFs. These results strongly implicate non-canonical Wnt pathway involvement in Lrp6 function with regard to neurulation.

The point mutation in Lrp6Cd changes an amino acid in the second extracellular β-propeller domain of Lrp6 (25), a region implicated in ligand binding (65). Although Dkk1 still binds Lrp6Cd protein, the mutation renders Dkk1 unable to antagonize β-catenin stabilization in Lrp6Cd/Cd MEFs (25). The mutation also reduces cell surface localization most likely because Lrp6Cd cannot bind its chaperone protein, MESD (Fig. 7) (25,58). Data presented here expand upon these findings to show that Lrp6Cd fails to produce the mature Lrp6 species and forms a truncated C-terminal fragment, which appears to result from proteolysis dependent upon the Cd mutation. The immature and truncated Lrp6Cd products were susceptible to the effects of PNgase and Tunicamycin, suggesting that the altered species are generated due to a defect in Lrp6 post-translational glycosylation. Importantly, unlike WT fusion protein, Lrp6Cd−GFP trafficking was much less efficient, regardless of whether MESD was co-expressed. We propose a model in which Lrp6Cd abnormally accumulates intracellularly because it cannot be properly processed and inserted into the membrane (Fig. 8).

Assessing previous characterization of Lrp6 function in both the canonical and non-canonical Wnt pathways in light of our model, several possible interpretations of our data are evident. Recent work reevaluating the canonical Wnt signaling pathway implicated the Lrp6 cytoplasmic tail as essential for efficient...
Dvl signaling and stabilization of β-catenin in the cytosol (66). Intracellular accumulation of C-terminal Lrp6Cd fragments in Lrp6CD−/− embryos could facilitate this interaction and contribute to cytosolic β-catenin levels observed in Lrp6CD−/− embryos (25). Increased cytoplasmic Lrp6Cd may also result in abnormal Dvl activation during frog neurulation (23). These new data refine our understanding of the role of Lrp6 Cd in neural tube closure primarily as a modulator of a RhoA-dependent, non-canonical signaling pathway (53,54). Here, immunoprecipitation and western blotting showed that Lrp6 can interact with DAAM1, a formin required for Wnt-dependent cell migration (39). Lrp6Cd/Cd mice were originally generated by Dufort and colleagues (36). Myristoylated Venus GFP (myr-Venus) mice express fluorescent protein that localizes to the inner leaflet of the cellular plasma membrane (39).

Together, these results indicate that both Lrp6 loss and gain of function disrupt apical–basal cell polarity in the neural folds, and this is associated with suppressed Cdc42 activity and especially with suppressed (in Lrp6−/− embryos) or elevated (in Lrp6CD−/− embryos) RhoA activity. Furthermore, inhibition of RhoA activity (i.e. ROCK inhibition) exacerbates (Lrp6−/−) or rescues (Lrp6CD−/−) mutant embryos from NTD in a genotype-dependent manner. This suggests that Lrp6 functions in neural tube closure primarily as a modulator of a RhoA-dependent, non-canonical Wnt pathway. The present evidence in a mouse is further illuminated by recent studies by Sagher et al. (2013), which directly demonstrate that Lrp6 impacts convergent extension during frog neurulation (23). These new data refine our model of defective Lrp6Cd action, implicating Lrp6 interaction with non-canonical signaling through a DAAM1-dependent complex, further illuminate the role of Lrp6 in neural tube closure and underscore the importance of further study of Lrp6 point mutations toward understanding the context-dependent functions of this versatile receptor.

MATERIALS AND METHODS

Animals

Mice were housed in climate-controlled Thoren units with a 12 h light-dark cycle. All procedures were in accordance with NIH Guidelines and were approved by the Institutional Animal Care and Use Committee at Weill Cornell Medical College. Cd mice were maintained on an A-strain background. Gene-trap reporter mice (BatGal) were originally generated by Dufort and colleagues (36). Myristoylated Venus GFP (myr-Venus) mice express fluorescent protein that localizes to the inner leaflet of the cellular plasma membrane (39). E9.5 embryos were labeled with LysoTracker (Invitrogen, L-75256), per manufacturer’s instructions, to evaluate cell death. Whole mount images were collected on a fluorescent stereomicroscope (Leica M165 FC) with a DFC310 FX camera. Immunohistochemistry

Heterozygous breeding pairs were placed together one evening and separated the following morning, which was designated E0.5. Embryos from intercrosses were harvested on E9.5 and yolk sacs were collected for genotyping. Embryos were fixed in 4% PFA overnight at 4°C prior to paraffin processing (Tissue Tek 2000, Miles Laboratories) or cryoembedding in OCT. Paraffin-embedded sections were sectioned coronally at 6 μm, as before (67). Primary antibodies included: anti-PH3 (Upstate Biotechnology, 16-189, 1:1000), anti-Pax3 (Developmental Studies Hybridoma Bank (DSHB), 1:1000), anti-Pax6 (DSHB, 1:1000), anti-Nkx2.2 (DSHB, 1:500), anti-GFP (Santa Cruz, sc-9996, 1:1000), anti-γ-tubulin (Santa Cruz, sc-10732, 1:1000), anti-F-Actin (DSHB, 1:5000), anti-phospho-myosin light chain 2 (p-MLC2), phosphorylated on threonine 19 (Cell Signaling 3675, 1:1000 on Western and 1:200 on ICC) and anti pan-cadherin (Abcam 16505, 1:2000). Immunolabeling was visualized using the species appropriate secondary antibody conjugated to a fluorophore [Alexa Fluor 488, Molecular Probes, A-11070 (rb) & A-11001 (ms), 1:500; Alexa Fluor 568, Molecular Probes, A-10042 (rb), 1:500; Alexa Fluor 488, Molecular Probes, A-21206, 1:500] or conjugated to HRP (Signet A-11070 (rb) & A-11001 (ms), 1:500; Alexa Fluor 568, Molecular Probes, A-10042 (rb), 1:500) or conjugated to HRP (Signet Streptavidin detection system, #2246) for detection with DAB. In fixed cells, Golgi apparatus was labeled with Wheat Germ Agglutinin AlexaFluor®594 (WGA594, Life Technologies W32466) at 10 mg/ml. Cryosections for in situ GTP-RhoA labeling were post-fixed in ice cold 4% PFA for 5 min at room temp, washed in H2O, quenched with H2O2 for 5 min, washed 3 × 5 min in PBS, blocked with SNIPER for 1 h at room temp and incubated in 10 μg/ml RBD-GST fusion protein (Cytoskeleton Inc., RT01) overnight at 4°C. After fusion protein incubation, sections were washed then fixed for 10 min with 4% PFA. Sections were incubated with primary rabbit anti-GST antibody (Calbiochem, PC53, 1:200) for 2 h at 37°C. Immunolabeling of fusion protein was visualized using a secondary antibody conjugated to a fluorophore (Alexa Fluor 488, Molecular Probes A-21206, 1:500).

Assessment of dorsal ventral patterning

Paraffin-embedded tissues were sectioned at 6 μm and every 10th section was taken for labeling with each of the antibodies. Sections extended from the midbrain to just below the level of the otic vesicle, or ~10–15 sections per marker per animal, times 4 embryos (40–50 sections per marker per genotype). The measures were simple ratios of traces in NIH Image J of the length of labeled neural fold divided by the total length of the fold. Measurements were taken bilaterally to avoid any bias due to asymmetrical embedding and values were averaged over the 10–15 sections for each embryo to avoid sampling error. Statistics were generated on the average ratio for each embryo examined (four embryos per each genotype).
Estimation of cell circularity

Apical basal polarity has been linked to cell shape changes in the neural folds of vertebrates (68). One measure of cell polarity is their circularity, which measures the degree to which the length and width of the cell approximates a circle (e.g. approaches a ratio of 1:1) (69). Using ImageJ software (NIH), the perimeter of cells at the apical surface nearest to the location of the DLHP was traced individually on sections of Lrp6<sup>Cd/Cd</sup>-::myrVenus and Lrp6<sup>+/−</sup>-::myrVenus embryos and their WT siblings. The circularity of cells was determined by the ImageJ software using the equation: circularity = 4π(area/perimeter<sup>2</sup>).

Plasmids and transfections

pLrp6 and pLrp6<sup>Cd</sup> plasmids were used in transfections as previously described (25). The DAAM1 coding region including a mycDKK tag at the C-terminus was cloned into the pCMV6 mammalian expression plasmid under the control of a CMV promoter (Origene, RC217675). Human pLrp6–GFP and mammalian expression plasmid under the control of a CMV promoter (Origene, RG218918). The Cd point mutation was introduced into the human pLrp6–GFP construct by site-directed mutagenesis (QuikChange, Stratagene) per manufacturer’s instructions. MsdM plasmid (pMsdM) was a gift of Dr B.C. Holder, SUNY Stony Brook. Co-localization of Lrp6–GFP with cell compartments was assessed using WGA594 to label the Golgi apparatus or cadherin to label plasma membrane. Co-transfections with myristoylated-RFP (Myr-RFP) plasmid (pCS2-myrm-Cherry, a kind gift from Dr Sean Megason, Harvard Medical School) assessed co-localization of the mutant and WT Lrp6–GFP fusion proteins with the plasma membrane. Plasmid transfers were carried out in NIH3T3 or HEK293 cells in 60 mm dishes and transfected with either single plasmids or in combination using Transfectin (BioRad) per the manufacturer. Transfection efficiencies were normalized either to total GFP/cell area or to Fop luciferase. Top/Fop luciferase reporter assays were carried out as previously described (70).

GTPase assay, western blotting and immunoprecipitation

MEFs were prepared from E14.5 Lrp6<sup>+/+</sup>, Lrp6<sup>Cd/Cd</sup> and Lrp6<sup>+/−</sup> mouse embryos (71). Active GTP-bound forms of RhoA and Cdc42 were measured using a pull-down assay according to the manufacturer’s instructions (Cytoskeleton, PAK02). Before cell lysis, cultures were treated for 10 min with 50 ng/ml recombinant Wnt3a or 100 ng/ml recombinant Wnt5a. To activate canonical Wnt signaling independently of Lrp-Fzd binding, Lrp6<sup>+/+</sup> cultures were treated with a highly selective GSK3β inhibitor CHIR99021 (50) at 10 μM for 4 h—a dose and incubation time selected from our experimental time course at which we found increases in nuclear β-catenin and cyclin D1 protein (Supplementary Material, Fig. S2). Cells were then harvested for detection of GTP-RhoA or GTP-Cdc42. Rhotekin-PBD or WBD agarose-bound proteins (Cytoskeleton, RT02) were separated by SDS–polyacrylamide gel electrophoresis (SDS–PAGE) and transferred to nitrocellulose. Blots were blocked in 3% BSA and incubated in mouse anti-RhoA (Santa Cruz, sc-418, 1:500) or mouse anti-Cdc42 antibody (BD, 610929 1:500). Labeling was detected using HRP-conjugated anti-mouse secondary antibodies and chemiluminescent reagents (Pierce, pico-34078 or femto-34096). X-ray films were scanned on an optical densitometer and relative protein concentrations were determined using Quantity One software (Bio-Rad). Optical densities of bands corresponding to GTP-bound GTPases were normalized to the amount of total RhoA or Cdc42, measured in western blot lanes loaded with equal amounts of lysate total protein. Alternatively, blots assessing protein expression were probed using rabbit anti-dephosphorylated β-catenin (Cell Signaling 8814, 1:1,000) and mouse anti-β-catenin (BD 610153 1:4,000), detected using fluorescent antibodies and quantified on an Odyssey IR imager (LiCor) per manufacturer’s protocol.

NIH3T3 cells were transfected with pLrp6 or pLrp6<sup>Cd</sup>, with or without cotransfected pDAAM1-mycDKK constructs. After 36 h recovery, replated cells were lysed in 350 μl of RIPA buffer [150 mM NaCl, 1.0% IGEPA<sup>®</sup> CA-630, 0.5% sodium deoxycholate, 0.1% SDS and 50 mM Tris, pH 7.4; 1 mM EDTA with protease inhibitor cocktail (Sigma)]. Lysates were clarified at 10 000 rpm for 5 min, pre-cleared with 25 μl of Protein A/G beads (Santa Cruz, sc-2003) at 4°C for 30 min and then incubated with 5 μl of Lrp6 antibody (Santa Cruz, sc-25317) or mycDKK antibody (Origene, TA50011) at 4°C with shaking overnight. Lrp6 and mycDKK antibodies were precipitated using 25 μl of Protein A/G beads overnight. IP fractions and whole lysates were separated by SDS–PAGE and transferred to nitrocellulose as described above. Blots were blocked with either 1% milk or 3% BSA and probed with Lrp6 or mycDKK antibody.

q-PCR

Total RNA was extracted from 70 to 80% confluent cultures of MEF cells grown in 12-well plates with TRI reagent (Sigma T9424). Template cDNA was prepared using an iScript first-strand cDNA synthesis kit (Bio-Rad 170-8891). Real-time PCR reactions were set up with a GoTaq SYBR green qPCR master mix (Promega) and run on an Applied Biosystems 7500 Fast Real-Time PCR system with the following primers: Axin2-F' TGATCTTCTTCTGAGCTG; Axin2-R' ACAGCGAGTTATCCAGCGAC; Gapdh-F' TTGATGGCAACAATCTCCAC; Gapdh-R' CGTCCCGTAGACAAAAATGGT.

Lrp6 glycosylation studies

Cell Culture: WT (Lrp6<sup>+/+</sup>) and Cd/Cd (Lrp6<sup>Cd/Cd</sup>) MEF cells were maintained in DMEM (Invitrogen Corp) containing high glucose with 10% fetal bovine serum, 2 mM l-glutamine, and penicillin (1 U/ml)/streptomycin (1 μg/ml) (Invitrogen) as supplements. RIPA lysis: Cells were washed with cold PBS and lysed with RIPA buffer, on ice. Lysed cells were centrifuged at 12,000 g for 15 min at 4°C and supernatants were collected as cell lysates. Deglycosylation: PNGase F (New England BioLabs Inc.) removed carbohydrate residues from proteins according to the company’s protocol. Tunicamycin (Sigma-Aldrich) was prepared as a 5 mg/ml stock solution in DMSO. In order to prevent post-translational protein glycosylation, cell cultures were treated with Tunicamycin (5 μg/ml, Sigma-Aldrich) for 24 h, and lysates prepared as above. Lrp6 proteins were detected on
Embryo culture and inhibitor treatment

Embryos from timed pregnancies of either Lrp6Cd/+ or Lrp6+/− mating pairs were collected when recovered embryos displayed between 10 and 16 somites and primary neural tube closure was underway, but cranial neurulation was not yet complete. Embryos were prepared for whole embryo culture as detailed elsewhere (72). Embryos were incubated in the roller apparatus (BTC Engineering, Cambridge, UK) at 37 °C for 2 h prior to addition of a Rho kinase (ROCK) inhibitor, Y-27632 (Enzo Life Sciences, ALX-270-333) or vehicle control (1 µl/ml DMSO). Embryos were allowed to develop in culture for 24 h from harvest to a time at which cranial neurulation should be complete and WT embryos without inhibitor displayed an average of 23–26 somites. Yolk sacs were removed for genotyping and embryos were photographed on a stereomicroscope with CCD camera (Leica) and scored for somite number.

Statistical treatment of data

Continuous variable measurements, including the extent of immunohistochemical labeling (Fig. 2), number of PH3 positive cells per µm (Fig. 3), cell circularity (Fig. 4), optical density (Figs 4 and 5) and percent fluorescence overlap with Golgi (Fig. 7) were statistically compared using a two-tailed Student’s t-test. In order to avoid sampling error from single sections, an average value was generated for each individual embryo that was then used for statistical comparison. Multiple measurements were taken from a series of histological sections (minimum of four sections per immune- or fluorescent protein label, serially sampling the caudal midbrain to otic vesicle) (Figs 2–4). Error bars in all figures represent the standard error of the mean. The frequency of neural tube closure (Fig. 6) was statistically compared using a z-test calculated using Microsoft Excel software. The exact values for each graph are reported in the results section using the following format: t(n − 1) = X or z = X, P < 0.05, where n = sample size (e.g. embryos or cultures), X is the computed t or z value and the significance threshold is a P-value < 0.05.

SUPPLEMENTARY MATERIAL

Supplementary Material is available at HMG online.

Conflict of Interest statement. None declared.

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