Membrane Targeting of Disheveled Can Bypass the Need for Arrow/LRP5

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The highly conserved Wnt signaling pathway regulates cell proliferation and differentiation in vertebrates and invertebrates. Upon binding of a Wnt ligand to a receptor of the Fz family, Disheveled (Dsh/Dvl) transduces the signal during canonical and non-canonical Wnt signaling. The specific details of how this process occurs have proven difficult to study, especially as Dsh appears to function as a switch between different branches of Wnt signaling. Here we focus on the membrane-proximal events that occur once Dsh is recruited to the membrane. We show that membrane-tethering of the Dsh protein is sufficient to induce canonical Wnt signaling activation even in the absence of the Wnt co-receptor Arrow/LRP5/6. We map the protein domains required for pathway activation in membrane tethered constructs finding that both the DEP and PDZ domains are dispensable for canonical signaling only in membrane-tethered Dsh, but not in untethered/normal Dsh. These data lead to a signal activation model, where Arrow is required to localize Dsh to the membrane during canonical Wnt signaling placing Dsh downstream of Arrow.

Wnt signaling consists of a series of evolutionarily conserved pathways taking part in many developmental processes1–3. The main or canonical signaling branch regulates cytoplasmic levels of Armadillo (Arm, β-catenin) affecting cell fate and proliferation1. Non-canonical pathways are involved in a variety of cellular polarity processes from convergence & extension in vertebrate gastrulation to ommatidial rotation in the Drosophila eye4–7. In canonical signaling, Disheveled (Dsh in Drosophila, Dvl in vertebrates) functions to relay the Wnt message from the Wnt receptors, Frizzled (Fz) and Arrow (Arr in Drosophila, LRP-5/6 in vertebrates), nucleating the membrane-proximal activation complex (signalosome)8. Dsh, as the most downstream shared component between canonical and non-canonical signaling likely also determines which pathway is activated9, 10.

The canonical Wnt signaling pathway is activated by Wnt binding to its receptors, Fz and Arr. This tri-partite complex transmits the extracellular signal to the intracellular components11 by recruiting Dsh to the membrane and forming the membrane-proximal activation complex consisting of Arr, Axin, and the kinases CK1 and GSK38. Once this complex forms, the cytoplasmic destruction complex, consisting of APC, Axin, CK1 and GSK3 is disrupted allowing Arm to escape phosphorylation and ubiquitin mediated degradation by the proteasome. As the destruction complex ceases to do its work, levels of Arm increase, and Arm enters the nucleus where, along with the transcription factor TCF, it activates transcription of target genes12–16.

When Arr was originally discovered as a co-receptor for Fz, it seemed relatively obvious that it should function upstream of Dsh as most intracellular signaling components function downstream of transmembrane ligand receptors. This turned out, however, not to be the case as Arr was shown to function downstream of Dsh17–22. This discovery led to the current model of activation complex assembly, where the Fz receptor recruits Dsh to the membrane forming a binding site for other pathway components, and bringing the cytoplasmic, C-terminal portion of Arr into close proximity with GSK3 and CK1 leading to Arr phosphorylation. Phosphorylated Arr/Lrp in turn becomes a binding site for Axin, the limiting factor for the assembly of the destruction complex.

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functionally taking the destruction complex apart and preventing Arm degradation\textsuperscript{18,23–27}. Phosphorylated Arr/Lrp also directly inhibits GSK3 by providing pseudo-substrates for GSK3 to bind\textsuperscript{28,29}.

The Dsh protein contains highly conserved DIX, PDZ and DEP domains (Sup. Figure 1). The PDZ and DIX domains are thought to be involved in canonical signaling where the PDZ domain interacts with the intracellular domain of Fz\textsuperscript{30}, and the DIX domain binds tightly to the DAX (also called DIX) domain of Axin\textsuperscript{31}. The DEP domain was originally thought to be specific to non-canonical signaling as the original planar cell polarity defect causing mutation dsh\textsuperscript{1} contained a point mutation in the DEP domain that was thought to prevent Dsh protein from localizing to the membrane\textsuperscript{32}. Structurally the DEP domain stabilizes Dsh’s interaction with the membrane by binding to charged phospholipids at the plasma membrane\textsuperscript{33,34}, and binding directly to Fz\textsuperscript{35,36}. Additionally, the DEP domain was recently shown to function in canonical signaling by nucleating signalosome assembly\textsuperscript{36,37}. Taken together, the function of the DEP and PDZ domains is to localize Dsh to the membrane through Fz. The DIX and DEP domains then can function to nucleate signalosomes\textsuperscript{35,36–38}.

Despite Dsh being discovered more than 20 years ago as a Wnt pathway component\textsuperscript{39}, the mechanism of how it relays signal to specific cellular responses is still poorly understood\textsuperscript{40}. Here we investigate the role of plasma membrane localization of Dsh protein. We utilize the Drosophila embryo to express an allelic series of Dsh proteins in both dsh and arr null genetic backgrounds. We use stringent developmental rescue and molecular assays to establish the functionality of Dsh alleles. We find that Arr is required for canonical pathway activation only if Dsh is not membrane localized. We find that both PDZ and DEP domains are dispensable for signaling when Dsh is membrane localized, but not when it is cytoplasmic.

**Results**

**Expression of membrane-tethered Dsh activates signaling.** Although most studies have reported that Dsh is a cytoplasmic protein, there have been some reported instances where it was found in the nucleus\textsuperscript{41}. In order to test the sufficiency of Dsh protein expression at the membrane, we attached a Src derived myristoylation (Myr) sequence to the N-terminus of Dsh. This sequence was originally used to tether Arm protein to the membrane, and we have found it highly effective for membrane localization of GSK3, Axin and APC\textsuperscript{16,42–44}. We proceeded to express tethered and un-tethered Dsh versions in embryos to test their effect on patterning. Normal Drosophila embryos show a repeating pattern of naked cuticle and denticle covered cuticle a result of segment polarity patterning\textsuperscript{45}. When Wnt signaling is turned off, all ventral epidermal cells produce denticles. The opposite is true when Wnt is turned on ectopically and most cells do not make denticles causing the ‘naked’ phenotype. When we expressed Dsh and Myr-Dsh in embryos, in both cases we saw a strong Wnt activation as visualized by a naked phenotype (Compare a control embryo (ArmGal4) in Fig. 1A with Fig. 1B,C respectively). Both Dsh isoforms were tagged with HA, so we could examine the localization in cells with anti-HA staining (Fig. 1B’C’). Additionally, we used phospho-tyrosine staining to visualize cell outlines and denticle precursors (Fig. 1A’–C’). We observed a concentration of Myr-Dsh at the membrane, and a more diffuse, intracellular localization for Dsh with some punctate structures observed at higher expression conditions (Fig. 1B’).

Loss of Dsh leads to a strong segment polarity phenotype\textsuperscript{46}. To establish the functionality of Dsh and Myr-Dsh, it was necessary to express these in a dsh null background. Therefore, we made maternal and zygotic dsh null embryos (dsh M/Z) completely lacking Dsh activity by crossing females with dsh germ line clones\textsuperscript{47} (i.e. the only laid eggs are homozygous mutant) with males providing re-expressed UAS-Dsh and UAS-Myr-Dsh (note that paternally rescued embryos were excluded from analysis by being y\textsuperscript{+}; see Materials and Methods). As expected, expression of Dsh rescues the canonical Wnt signaling defect of dsh mutants and causes some ectopic activation leading to a mild naked phenotype (Compare Fig. 2A and B). Similarly, Myr-Dsh rescues the loss of signaling as shown by the naked cuticle patches in these embryos (Fig. 2C). We visualized this both at the cuticle level, and used pTyr/HA staining to show loss of denticle precursors in Dsh expressing cells (Fig. 2B’C’ and 2B’C’ respectively). Taken together, these results show that at least for canonical Wnt activation, membrane localized Dsh is active and asks the question whether Dsh membrane localization is sufficient to bypass the requirement of Wnt co-receptors.

**Membrane tethered Dsh activates signaling in arr mutants.** The Wnt co-receptor Arr binds Wnts along with Fz\textsuperscript{18,19,21}. In most signaling pathways, this should place it epistatically upstream of an intracellular component such as Dsh as is the case for Fz\textsuperscript{30}. However, this is not the case for Arr and Dsh, as simple overexpression of Dsh in an arr mutant does not activate the pathway\textsuperscript{47}. We repeated this experiment in order to compare the function of Myr-Dsh and Dsh, and as expected we find that expression of Dsh in maternally and zygotically mutant arr embryos does not activate signaling (Compare Fig. 3A to Fig. 3B). In contrast to normal Dsh, expression of membrane-tethered Myr-Dsh strongly activates signaling in arr (M/Z) embryos (Fig. 3C). Consistently, staining embryos for pTyr and HA-Dsh reveals denticle covered embryos in arr (M/Z; ArmGal4–UAS-Dsh despite the presence of HA-Dsh (Fig. 3B’B’). However, we observed a loss of denticles in arr (M/Z) expressing Myr-Dsh (Fig. 3C’C’). From these experiments, we conclude that signal activation through Dsh is indeed downstream of Arr, and that for the activation to occur, the requirement for Arr can be bypassed by localizing Dsh to the membrane.

**Dsh domains at the membrane.** In order to analyze this result further, we determined the domain requirement of Dsh for activation of Wnt signaling by taking a structure function approach. Based on results with the dsh\textsuperscript{1} allele causing PCP specific defects, the DEP domain was sometimes considered more important for non-canonical Wnt signaling than for canonical signaling, although overexpression and in vitro experiments suggested a role for the DEP domain also in canonical signaling\textsuperscript{35,36,49,50}. We therefore first verified that, compared to wild-type Dsh\textsuperscript{31,32}, Dsh lacking the DEP domain or the DEP domain and C-terminus expressed under control of its endogenous promoter cannot rescue the dsh\textsuperscript{1} null allele. Indeed, five independent transgenic insertions
lacking the DEP or DEP-C-terminal domain each fail to rescue viability (canonical signaling; Fig. 4A). Similarly, the three lines tested also are unable to rescue the PCP defects of the \(d_{sh}^{1}\) allele in the eye and wing (Fig. 4B–F).

We then investigated the effect of individual domains of Dsh in the membrane proximal activation complex. We made membrane tethered constructs where individual domains were deleted and expressed them in wild type embryos, \(d_{sh}\) (M/Z) and \(arr\) (M/Z) mutant embryos (Sup. Figure 1). We find that expression of Myr-Dsh lacking either the PDZ or DEP domains can activate the pathway. Both proteins when expressed in otherwise wildtype embryos show a strong gain of function (GOF) phenotype leading to nearly naked patterning (Fig. 5B,C). In contrast, expression of the Myr-Dsh\(\Delta\)DIX leads to a strong loss of signaling phenotype with many ectopic denticles, thus likely acting as dominant negative (Fig. 5A). This was verified with HA/pTyr staining to show loss of denticle precursors in Dsh expressing cells (Fig. 5B',C').

We then proceeded to test the deletion constructs (Sup. Fig. 1) in \(d_{sh}\) (M/Z) loss of function mutants to assess their ability to rescue the loss of endogenous Dsh. As expected from the dominant negative effect displayed in a wild-type background, Myr-Dsh\(\Delta\)DIX failed to rescue signaling in embryos (Fig. 5D). Dsh lacking the DEP or PDZ domains rescued \(d_{sh}\) (M/Z) embryos when expressed as Myr-Dsh\(\Delta\)DEP and Myr-Dsh\(\Delta\)PDZ (Fig. 5E,F; quantified in Table 1). This finding confirmed the notion that the DEP and PDZ domains stabilize Dsh membrane localization. Nevertheless, although with respect to frequency of rescued embryos, rescue was efficient, the extent

Figure 1. Cuticle preparations of (A) Wild type embryo showing six rows of denticles per segment. (B) Embryo expressing Dsh resulting in expansion of naked cuticle causing loss of one or two rows of denticles. (C) wild type embryo expressing Myr-Dsh showing signaling activation phenotype (GOF). To increase expression levels, we doubled embryonic Gal4 by combining daGal4 and ArmGal4 into a 2XGal4 line. (A’-A”) Wild-type embryo stained for Arm and denticle precursors (pTyr). (B’-B”) Embryo overexpressing Dsh stained for ectopic tag HA and denticle precursors (pTyr). Similar staining in wild type embryo expressing Myr-Dsh (C’-C”).” panels show merged images.
of the rescue was weaker than with the full-length Myr-Dsh (Fig. 2, compare the amount of naked cuticle between conditions) suggesting that the DIX and PDZ domains may have some further function in signal transduction at the membrane.

Next, we expressed the membrane tethered deletion constructs in arr mutant embryos. Again, we find that in contrast to Myr-Dsh, Myr-DshΔDIX failed to rescue signaling in embryos lacking both maternal and zygotic Arr (Fig. 5G). Expression of Myr-DshΔDEP and Myr-DshΔPDZ showed rescue in arr (M/Z) embryos as revealed by distinct regions of naked cuticle in arr mutants (Fig. 5H,I; quantified in Table 1). Taken together, these results suggest that bringing Dsh to the membrane can bypass the requirement for Arr, but that the DIX domain is required to activate signaling even under those circumstances, while both the DEP and PDZ are dispensable, but may enhance signaling as the rescue was not as good as with full-length Myr-Dsh.

Dsh forms the signalosome at the membrane to activate signaling by inhibiting the action of the destruction complex40. Downstream, Arm protein levels increase and signaling is activated. We tested whether signaling could be activated downstream of both complexes when disrupted. We expressed Arm alleles that were activated either by deleting the N-terminus or by changing a specific phosphorylation site (ArmS56A16, 42, 53, 54), both of which block phosphorylation and degradation of Arm. Their expression in dsh (M/Z), arr (M/Z) or wildtype embryos rescued the dphthalmocyte phenotype to varying degrees (Fig. 3C-F). Similar rescue was observed in double mutants and we found that ArmS56A is a stronger activator than ArmΔN (Fig. 3G; quantified in Table 1). These results indicate a shared role for Arm activation in dsh and arr mutants.

**Figure 2.** Cuticle preparations of (A) dsh (M/Z) mutant showing a lawn of denticles without activation of signaling. (B) dsh (M/Z) mutant embryo expressing Dsh resulting in expansion of naked cuticle. (C) dsh (M/Z) mutant embryo expressing Myr-Dsh showing signaling activation phenotype. (A'-A') dsh (M/Z) mutant embryo stained for Arm and denticle precursors (pTyr). (B'-B') dsh (M/Z) mutant embryo expressing Dsh stained for ectopic tag HA and denticle precursors (pTyr). Similar staining in dsh (M/Z) mutant embryo expressing Myr-Dsh (C'-C''). panels show merged images.
embryos led to loss of denticles or an activation of Wnt signaling (Fig. 5J–L). These results suggest that the downstream pathway is unaffected by the loss of the signalosome (see also discussion).

Role of membrane-tethered Dsh in canonical signaling. Previous work has led to a model where the DIX domain of Dsh is required for bringing Axin to the membrane, taking it away from the destruction complex55. Dsh transgenes lacking the DIX domain act as dominant negatives (Fig. 5A 17). In order to test our Dsh model further, we used a TopFlash assay, where the luciferase gene is attached to multimerized TCF binding sites to analyze functionality of Dsh variants in cell culture. We used untethered overexpressed Dsh in S2R+ cells as the baseline for Wnt signaling activation and compared it to the various membrane-tethered Dsh constructs. We found that full length Myr-Dsh could activate the TopFlash promoter to a higher level than untethered Dsh (Fig. 6A; see Fig. 6B for expression levels). Deletion of the DIX domain in Myr-DshΔDIX did not activate the reporter. In contrast, Myr-DshΔDEP activated TopFlash reporter to a similar extent to full-length Myr-Dsh whereas Myr-DshΔPDZ showed a somewhat lower extent (Fig. 6A), thus correlating with the in vivo results.

We next looked at endogenous target genes downstream of Wnt signaling. Wnt signaling activates and maintains its own activity by activating wg and en transcription64. Using qRT-PCR, we therefore quantified en and wg transcript levels in embryos upon overexpression of the various Dsh mutant transgenes relative to wildtype Dsh (normalized to the housekeeping gene Rpl32). We compared the various membrane-tethered Dsh constructs expressed in otherwise wild-type embryos (Fig. 7A; transgene expression levels are shown in 7B), and found that full length Myr-Dsh activated to a similar level to untethered Dsh. Consistent with the DN effect in vivo (Fig. 5A) the deletion Myr-DshΔDIX lowered the overall abundance of wg and en. Myr-DshΔPDZ showed an insignificant reduction in levels of wg and en, whereas Myr-DshΔDEP showed strong activation (Fig. 7A). We looked at Arm protein levels in the various conditions, and these correlated with the levels of Dsh activity with Myr-Dsh and Myr-DshΔDEP showing increased Arm protein (Fig. 7C). Taken together, these results support the overall activity levels of Dsh shown in the in vivo rescue and epistasis assays.
Membrane localized Dsh is protected from degradation. As we looked at the levels of HA-tagged Dsh in embryos (Fig. 7B), we noticed consistently that the levels of cytoplasmic protein (Fig. 7C) were much lower than the expressed myristoylated forms (in spite of comparable activities). As these lines were made by phiC31 integration into identical sites, we expected similar levels of protein, but this was clearly not the case. We looked at the mRNA expression levels by qPCR, and observed that all Dsh forms were expressed at similar levels at the transcript level (see HA levels in Fig. 7B), but the protein levels were much lower for the un-tethered version of Dsh (Fig. 7C). This result shows that membrane localization may protect Dsh from degradation in the cytoplasm in vivo.56.

Discussion
The identification of Dsh as an activator of the Wnt pathway and its placement in the pathway upstream of GSK3 and Arm led to a simple genetic description of the Wnt pathway. Yet years later, the molecular function of Dsh is still the subject of debate. Scaffold proteins such as Axin and Dsh perform complex roles in signaling by bringing several proteins into close proximity in different cellular compartments. Our work in this paper focuses on the role of membrane localization of Dsh, its relationship with the Wnt co-receptor Arr, and the domains utilized in the signaling process. We show that, genetically, the role of Arr is to localize Dsh to the membrane in response to Wnt, as Arr's role in signaling is bypassed when Dsh is targeted to the membrane.

Our structure/function studies in vivo suggest that the DIX domain is absolutely required for signal activation, and further, its loss can cause a dominant negative effect. The DEP and PDZ domains are dispensable for canonical signaling only if Dsh is membrane tethered, but their absence decreases the effectiveness of membrane tethered Dsh in activating canonical signaling in the absence of endogenous Dsh and Arr, as shown phenotypically in our rescue assays of maternal-zygotic null alleles. We show that the PDZ domain is dispensable for Dsh function in canonical signaling as expression of Myr-DshΔPDZ rescues dsh (M/Z) embryos and activates strongly in otherwise wildtype embryos showing that the interaction of PDZ with Fz isn't crucial if Dsh is at the membrane. But the PDZ domain does contribute to signaling as its absence weakens the activation in all our assays.

Although our results do not directly explain how the destruction complex is inactivated, they do point to a model of how the membrane-proximal activation complex or signalosome functions. Under normal signaling conditions, Dsh recruitment to the membrane is followed by GSK3/CK1 phosphorylation of sites on the cytoplasmic tail of Arr forming a binding site for Axin effectively disrupting the degradation complex. These sites work in conjunction with the Axin DAX/Dsh DIX interaction to form Wnt signal activating signalosomes.18, 23, 25–27, 31, 57, 58.

Our results suggest that localizing Dsh to the membrane is sufficient to remove Axin from the destruction complex, thereby blocking Arm degradation, especially as the membrane localized Dsh is protected from degradation. These findings do not necessarily distinguish between the several models for destruction complex inactivation, but in the absence of Arr, pseudo-substrate sites for inhibition of GSK3 cannot be formed at the membrane.
Figure 5. Cuticle of (A) Myr-DshΔDIX expressed in 'wild type' embryos, producing more ectopic denticles due to signaling inactivation. (B,C) Cuticles of Myr-DshΔPDZ and Myr-DshΔDEP expressing embryos, showing a naked phenotype (GOF). Staining for ectopic tag HA and denticle precursors (pTyr) in wild type embryos expressing Myr-DshΔDIX (A’-A”), Myr-DshΔPDZ (B’-B”) and Myr-DshΔDEP (C’-C”). D–F) Myr-DshΔPDZ (E) and Myr-DshΔDEP (F) expression in dsh (M/Z) mutant embryos rescued Wnt signaling whereas expressing Myr-DshΔDIX (D) in dsh (M/Z) mutant embryos did not. (G) Arrow null embryo expressing Myr-DshΔDIX showing no activation of signaling and hence the wingless phenotype. (H–I) Arrow null mutant suppressed by expression of Myr-DshΔPDZ and Myr-DshΔDEP suggesting requirement for a membrane recruitment. Cuticle of arr (M/Z) (J), dsh (M/Z) (K) and wildtype (L) embryos expressing activated Arm alleles either lacking the N-terminus (ArmΔN) or a specific ArmS56A substitution (ArmSA) both of which block phosphorylation and degradation of Arm. Expression in these backgrounds led to loss of denticles, reflecting constitutive activation of Wg signaling.
recombination is induced using the FLP/FRT method in ovaries 47, 66, 67. Oregon R was used as the wild-type strain.

dshV26 (or 3) and ovoD1 use the G13 FRT. The following crosses were conducted:

| Crosses and expression of UAS constructs. |
|-------------------------------------------|
| 1. arr2 FRTG13/ovoD1 FRTG13; da-Gal4/+ females x arr2/CyO-GFP; UAS-Dsh-3XHA |
| 2. arr2 FRTG13/ovoD1 FRTG13; da-Gal4/+ females x arr2/CyO-GFP; UAS-Myr-Dsh-3XHA |
| 3. arr2 FRTG13/ovoD1 FRTG13; da-Gal4/+ females x arr2/CyO-GFP; UAS-Myr-DshΔDIX-3XHA |
| 4. arr2 FRTG13/ovoD1 FRTG13; da-Gal4/+ females x arr2/CyO-GFP; UAS-Myr-DshΔPDZ-3XHA |
| 5. arr2 FRTG13/ovoD1 FRTG13; da-Gal4/+ females x arr2/CyO-GFP; UAS-Myr-DshΔDEP-3XHA |
| 6. y, dshΔPDZ-3XHA FRT18E/ovoD1 FRT18E; arm-Gal4/+ females x UAS-Dsh-3XHA |
| 7. y, dshΔPDZ-3XHA FRT18E/ovoD1 FRT18E; arm-Gal4/+ females x UAS-Myr-Dsh-3XHA |
| 8. y, dshΔPDZ-3XHA FRT18E/ovoD1 FRT18E; arm-Gal4/+ females x UAS-Myr-DshΔDIX-3XHA |
| 9. y, dshΔPDZ-3XHA FRT18E/ovoD1 FRT18E; arm-Gal4/+ females x UAS-Myr-DshΔPDZ-3XHA |
| 10. y, dshΔPDZ-3XHA FRT18E/ovoD1 FRT18E; arm-Gal4/+ females x UAS-Myr-DshΔDEP-3XHA |
| 11. y, dshΔPDZ-3XHA FRT18E/ovoD1 FRT18E; arm-Gal4/+ females x UAS-ArmΔN2,2XHA |
| 12. arr2 FRTG13/ovoD1 FRTG13; da-Gal4/+ females x arr2/CyO-GFP; UAS-Myr-ArmΔN-2XHA |

Table 1. Quantification of embryonic phenotypes. Column (ArmGal4) represents the phenotype of a simple Gal4 driver to UAS construct cross where all embryos are expected to overexpress Dsh. Column (arr) represents embryos where all maternal arr is removed, paternally rescued embryos are removed by GFP selection and 50% of embryos are expected to express Dsh. Therefore, ‘50% naked cuticle’ corresponds to full suppression. Column (dsh) represents embryos where all maternal endogenous Dsh is removed, paternally rescued embryos are ignored based on y +, and 50% of embryos are expected to express UAS-Dsh. Therefore, ‘50% naked cuticle’ corresponds to full rescue. The naked phenotype is defined as fewer denticles than in wild type, but not necessarily that all denticles are absent.

suggesting that this may not be the only way that GSK3 can be inhibited, and that the most likely mechanism of activation is the titration of Axin away from the cytoplasm.

We find that membrane localized Dsh accumulates to higher levels than normal Dsh through a post-translational mechanism. Previous studies have suggested that Dsh can be degraded through proteasomal degradation 60, 61, but another recent finding suggests that the basolateral complex protein Discs Large protects Dsh from degradation 60. This adds an interesting dimension to Dsh regulation as we have previously observed interactions between Wnt pathway components and apicobasal machinery 54, 61–63. We attributed these effects to non-canonical signaling, but it could have effects on canonical signaling as well 55, 64, 65.

It was found that Dsh in vertebrate cell culture and Xenopus embryos shuttles between the cytoplasm and nucleus and that the ability to enter the nucleus is important for Dsh function specifically in canonical Wnt signaling 61. It was suggested that nuclear Dsh might affect degradation of β-Catenin in the nucleus or indirectly in the cytoplasm. Our results showing that stabilized Arm is constitutively active in a M/Z dsh null background argues that any nuclear function of Dsh acts upstream of Arm and is not an additional, Arm-independent nuclear function of Dsh, a scenario that previously had not been excluded.

Taken together, we suggest that the membrane proximal activation complex brings together several proteins and enzymes – Fz, Arr, GSK3, CK1, Axin and Dsh. Formation of the complex leads to phosphorylation of Arr by GSK3 and CK1, creating binding sites for Arr brought to the membrane by the DIX domain of Dsh. Dsh is likely brought to the membrane through Fz binding to the PDZ and DEP domains and DEP binding to charged phospholipids. In our system, we can bypass the creation of Axin binding sites on Arr by directly tethering Dsh to the membrane.

Materials and Methods

Crosses and expression of UAS constructs. Maternally mutant eggs were generated by the dominant female sterile technique where balanced mutants are crossed to the dominant female sterile mutation ovoD1 and recombination is induced using the FLP/FRT method in ovaries 63, 66, 67. Oregon R was used as the wild-type strain. Please see Flybase for details on mutants used (flybase.bio.indiana.edu). Mutants used: dshΔPDZ, arrΔDIX-3XHA and arrΔN2. For mis-expression experiments, the ArmGal4 2nd chromosome and daGal4 3rd chromosome drivers were used. All X-chromosome mutants use FRT 101 except for dshΔPDZ-3XHA that has FRT 18E and second chromosome arrΔN-2XHA use the G13 FRT. The following crosses were conducted:

1. arr2 FRTG13/ovoD1 FRTG13; da-Gal4/+ females x arr2/CyO-GFP; UAS-Dsh-3XHA
2. arr2 FRTG13/ovoD1 FRTG13; da-Gal4/+ females x arr2/CyO-GFP; UAS-Myr-Dsh-3XHA
3. arr2 FRTG13/ovoD1 FRTG13; da-Gal4/+ females x arr2/CyO-GFP; UAS-Myr-DshΔDIX-3XHA
4. arr2 FRTG13/ovoD1 FRTG13; da-Gal4/+ females x arr2/CyO-GFP; UAS-Myr-DshΔPDZ-3XHA
5. arr2 FRTG13/ovoD1 FRTG13; da-Gal4/+ females x arr2/CyO-GFP; UAS-Myr-DshΔDEP-3XHA
6. y, dshΔPDZ-3XHA FRT18E/ovoD1 FRT18E; arm-Gal4/+ females x UAS-Dsh-3XHA
7. y, dshΔPDZ-3XHA FRT18E/ovoD1 FRT18E; arm-Gal4/+ females x UAS-Myr-Dsh-3XHA
8. y, dshΔPDZ-3XHA FRT18E/ovoD1 FRT18E; arm-Gal4/+ females x UAS-Myr-DshΔDIX-3XHA
9. y, dshΔPDZ-3XHA FRT18E/ovoD1 FRT18E; arm-Gal4/+ females x UAS-Myr-DshΔPDZ-3XHA
10. y, dshΔPDZ-3XHA FRT18E/ovoD1 FRT18E; arm-Gal4/+ females x UAS-Myr-DshΔDEP-3XHA
11. y, dshΔPDZ-3XHA FRT18E/ovoD1 FRT18E; arm-Gal4/+ females x UAS-ArmΔN2,2XHA
12. arr2 FRTG13/ovoD1 FRTG13; da-Gal4/+ females x arr2/CyO-GFP; UAS-Myr-ArmΔN-2XHA

| | ArmGal4 | dsh (M/Z) | arrow (M/Z) |
|---|---|---|---|
| UAS-Dsh | Naked 100% | naked cuticle (rescue or GOF) 50% | wg phen. 100% |
| N > 100 | N = 98 | N = 220 |
| UAS-MyrDsh | Naked 100% | Naked 48% | Naked 49% |
| N > 100 | N = 95 | N = 223 |
| UAS-Myr-DshΔDIX | wg phen 100% | wg phen 98% | wg phen. 100% |
| N > 100 | N = 101 | N = 288 |
| UAS-Myr-DshΔPDZ | Naked cuticle expansion 100% | Naked 48% | Naked 49% |
| N > 100 | N = 120 | N = 294 |
| UAS-Myr-DshΔDEP | Naked cuticle expansion 100% | Naked 48% N = 154 | Naked 47% N = 138 |
X chromosomes were marked with the yellow mutation and the CyO balancers were marked GFP to simplify analysis. For rescue of ds, paternally rescued embryos were excluded as they were either y+ (fathers all had wildtype y alleles). For rescue of ar, paternally rescued embryos were excluded by selecting against a GFP balancer (genotype of fathers was with arr2 over CyO-GFP). As mothers were heterozygous for the Gal4 source, maximal rescue is reflected by a drop of phenotype to 50% (only half of the embryos will express Gal4). For all crosses, more than 100 embryos were analyzed in multiple, separate experiments (n > 95).

Transgenes and GAL4 driver lines. Two ubiquitous drivers were used for expression of transgenes: the weaker armadillo-GAL4 and the stronger daughterless-GAL4. UAS constructs were made using Gateway recombination (Invitrogen). Myristoylated constructs were made by adding a sequence identical to the NH2 terminus of src (MGNKCCSKRQGTMAGNI) to the NH2 terminus of GSK-3 by PCR. This sequence has proven to be very effective for membrane targeting of Arm16, 42, 43, 53. The PCR products were then transferred by Gateway cloning (Invitrogen) into pUASg.attB with C-terminal 3XHA tag (A kind gift from J. Bischof and K. Basler, Zurich).

pCasp_dshDshΔDEP was made by amplifying the C-terminus of Dsh lacking the DEP domain with primers DshΔDEP_For_Xho (TAACCTCGAGGAGATCGTTAAGGCGATGACGAAGGAGCGCA ATCCCAATCTGTTG) and DshΔDEP_rev_Xba (TAGTTCTAGATCGGCGCCGCTTTAATACGTAA TTAAATACGGA) and cloned as XhoI/XbaI fragment into pCasp_dshDsh_silentKpnSac_EGFP. pCasp_dshDshΔDEP-CT was made by replacing the XhoI/XbaI fragment of pCasp_dshDsh_silentKpnSac_EGFP with annealed oligos DshΔDEP_CTerminalelements of DshΔDEP_CT_lower and upper (CTAGAGTCGCGCCGCGCTTTACTTCTGT CAGCCGGCTTAAGATGAGCACGAAGGCGGCGGACT). Transgenes were injected into either w1118(Casp constructs) or attP2(Strain #8622)P[CaryP]attP2 68A4 by Rainbow Transgenics or BestGene Inc. (California).

Figure 6. (A) Effects of Dsh deletion constructs on TOPflash reporter activity. S2R+ cells were co-transfected with TOPflash reporter plasmid, Renilla luciferase-Pol III Vector and the indicated Dsh constructs. Mock transfected S2R+ cells were treated as the baseline (control) for Wnt signaling activation. Results are representative of three independent experiments and the average of three replicates (mean ± SD). Statistical significance was tested using the Student’s t-test. **p < 0.01 relative to control. (B) Western blot comparing total Arm protein levels in cells transfected with the Flag tagged Dsh constructs indicated with * . α-tubulin was used as a loading control.
non-disjunction events). The relative rescue index was counted as fractions of these males normalized to the average rescue efficiency of wild-type Dsh (note that wild-type Dsh constructs contain two silent point mutations\(^{51}\)). To score activity of Dsh variants for non-canonical Wnt signaling, transgenic males were crossed to dsh\(^{1}\) females and PCP defects in eye sections and wings were assessed in male offspring as described in\(^{71,72}\).

**Antibodies and Immunofluorescence.** Embryos were fixed with Heat-Methanol treatment\(^{73}\) or with heptane/4% formaldehyde in phosphate buffer (0.1 M NaPO4 pH 7.4)\(^{16}\). The antibodies used were: anti-Armadillo (mAb N2 7A1, Developmental Studies Hybridoma Bank developed under the auspices of the NICHD and maintained by The University of Iowa, Department of Biological Sciences, Iowa City, IA 52242), anti-HA (ratAb 3F10 and mouse 12CA5, Roche), rabbit anti-Armadillo\(^{74}\), phospho-tyrosine pY99 (Santa Cruz Biotechnology), anti-\(\beta\)-tubulin (E7, DSHB), and anti-FLAG (F9291, Sigma-Aldrich). Staining, detection and image processing as described in\(^{75}\).

**Western Blotting.** Embryos were selected for fertilization and developmental stage, lysed in RIPA buffer (Cell Signaling Technology) with protease inhibitor cocktail (Roche), the extracts were separated on 4–20% gradient SDS-PAGE gel (Biorad), and blotted as described in\(^{16}\).

**TOPflash assay.** TOPflash luciferase assays (TCF/LEF reporter assays) were performed to assess the effect of the Dsh deletion constructs on canonical Wnt-signalling. S2R+ cells were co-transfected with dTF12 TOPflash reporter (TCF Reporter Plasmid; A kind gift from R. DasGupta, Singapore)\(^{77}\), Renilla luciferase-Pol III Vector (Promega) and the respective Dsh constructs using lipofectamine 3000 (ThermoFisher Scientific) according to the manufacturer’s instructions. Cell lysates were prepared 48 h after transfection and luciferase activity was measured using the Dual-Luciferase Reporter Assay System (Promega) according to the manufacturer's instructions. The relative TOPflash luciferase activity was measured using the ratio of firefly/renilla luciferase activity and the data was presented as mean ± SD.

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**Figure 7. (A) Comparison of gene expression levels of wz & en in embryos expressing the indicated Dsh constructs.** Reduced wz and en expression levels were observed in Myr-Dsh\(\Delta\)DIX expressing embryos compared to control embryos and embryos expressing Dsh. *\(p < 0.05\), **\(p < 0.01\) relative to control. (B) Gene expression levels of HA Tag in embryos expressing the respective Dsh constructs. Significantly elevated expression of HA was detected in all transgenic lines compared to control. ***\(p < 0.01\). (C) Western blot comparing total Arm protein levels in embryos expressing the various HA tagged myristoylated Dsh constructs.
RNA Extraction, cDNA Synthesis and qPCR. Total RNA was extracted for each experimental condition from 50ul of Drosophila embryos (collected 14–16hrs after deposition) using RNeasy Mini Kit (Qiagen) as per the manufacturer's protocol. Total RNA concentration was measured using NanoDrop ND-2000 Spectrophotometer and the purity of the samples was determined by the OD ratios. A$_{260}$/A$_{280}$. One µg of total RNA was reverse transcribed in a 20 µl reaction volume using the QuantiTect reverse transcription kit (Qiagen) according to the manufacturer's protocol. Gene specific primer sequences were obtained from Fly Primer Bank (en forward primer, 5' TCCGTGATCGGTCGATGAGT-3'; en reverse primer, 5'-CGCCCGATCTATCCACATC-3'; wg forward primer, 5'-GACCCCGAGTCACCTCTAC-3'; wg reverse primer, 5'-CGGCGATTCTGAACCTGTTG-3'; HA forward primer, 5'-GGTCCCTGACTAGCGGGGCTA-3'; HA reverse primer, 5'-AGCGTAACTCGGAAAGTCAT-3'; Rpl32 forward primer, 5'-CCCAAGGGTATCGCAACAGAAG-3'; Rpl32 reverse primer, 5'-CGATCTCGCCGCGATACAC-3').

Quantitation of mRNA was performed using SYBR® Green Assay (Thermo Fisher Scientific) on the PikoReal™ Real-Time PCR System (Thermo Fisher Scientific) and a PCR product dissociation curve was generated to ensure specificity of amplification. Rpl32 was used as an endogenous control and relative quantitation was performed using relative quantification (2^{-ΔΔCt}). Results were generated from three technical replicates for each mRNA. The average relative expression ± standard deviation (SD) was determined and two sample t-test was carried out to determine statistical significance.

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**Author Contributions**

P.K., V.Y.M.L., A.G.M., J.S., A.J., N.S.T. designed and performed the experiments. N.S.T., P.K. and A.J. wrote the paper. All authors reviewed manuscript.

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