The Drosophila Cadherin Fat Regulates Tissue Size and Planar Cell Polarity through Different Domains

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Abstract

The Drosophila Cadherin Fat (Ft) has been identified as a crucial regulator of tissue size and Planar Cell Polarity (PCP). However, the precise mechanism by which Ft regulates these processes remains unclear. In order to advance our understanding of the action of Ft, we have sought to identify the crucial Ft effector domains. Here we report that a small region of the Ft cytoplasmic domain (H2 region) is both necessary and sufficient, when membrane localized, to support viability and prevent tissue overgrowth. Interestingly, the H2 region is dispensable for regulating PCP signaling, whereas the mutant Ft lacking the H2 region is fully capable of directing PCP. This result suggests that Ft’s roles in PCP signaling and tissue size control are separable, and each can be carried out independently. Surprisingly, the crucial regions of Ft identified in our structure-function study do not overlap with the previously reported interaction regions with Atrophin, Dco, or Lowfat.

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Introduction

The Drosophila Fat (Ft), a large 34 cadherin domain-containing transmembrane adhesion molecule, plays a key role in the establishment of epithelial planar cell polarity (PCP) [1,2,3]. PCP is a form of polarity along an axis within the plane of an epithelium. It is manifested in many Drosophila tissues, including the uniform orientation of hairs and bristles on the adult body and wing, and the precise alignment of ommatidia in the eye. The role of Ft has been proposed to provide a directional signal for aligning PCP [1,2]. It has been intensively investigated how Ft and another group of conserved signaling molecules, core PCP proteins, are working together to mediate the PCP signaling [4,5,6,7,8,9]. Although considerable insight has been gained into the core PCP pathway, how Ft tranduces downstream PCP signal remains unclear.

In addition to regulating PCP, Ft has also been identified as a negative regulator of tissue growth. Animals lacking Ft function exhibit massive overgrowth of all imaginal discs, the larval epithelial tissues which ultimately form much of the adult body. Mutant animals subsequently die as pupae [10,11]. In addition to the overall increase in tissue size seen in animals lacking Ft function, studies of \textit{ft} mutant cells in mosaic animals have shown that cells lacking Ft function proliferate much more extensively than their wild-type neighbors [12]. Each of these observations suggests that cells lacking Ft are unable to respond to signals that normally limit their growth and proliferation.

Significant advancements have been made to elucidate the downstream pathway of Ft in past few years [13,14,15,16]. Several genetic studies have demonstrated that the Hpo/Wts signaling pathway is an important target of Ft in tissue growth regulation [17,18,19,20,21]. The core of the Hpo/Wts signal transduction system is a protein kinase cascade consisting of the serine/threonine kinases Hippo (Hpo) [22,23,24,25,26] and Warts (Wts) [27,28], and their associated scaffolding proteins Salvador (Sav) [29,30] and Mats (Mob as tumor suppressor) [31]. Upon activation of the kinases, the transcriptional activator Yorkie (Yki) is phosphorylated and inactivated, thus leading to reduced expression of Yki targets [32,33,34]. These targets include proliferation promoting factors such as Cyclin E and anti-apoptotic factors such as DIAP1 [25,30]. As a result, activation of the Hpo/Wts pathway leads to reduced cellular proliferation and increased cell death [35,36,37]. In contrast, inactivation of the Hpo/Wts protein kinase cascade in clones leads to elevated expression of Yki targets and overproliferation in a manner similar to clones of cells lacking Ft function. The similarity of clonal phenotypes as well as the induction of Yki targets in \textit{ft} mutant cells indicates that Ft regulates Hpo/Wts signaling. Genetic studies also placed Dachs, an unconventional myosin, Approximated, a palmitoyltransferase, and Expanded, a tumor suppressor, between Ft and downstream Hpo/Wts regulation and/or PCP signaling [17,18,20,38,39]. However, due to lack of biochemical link and complex nature of Ft signaling, the precise mode of action of Ft is not understood.

Several binding partners of Ft cytoplasmic domain have been identified and proposed to carry out some of Ft downstream activities. Atrophin/Grunge, a transcriptional repressor, which
binds to the C-terminal fragment of Ft, has been proposed to mediate downstream PCP signaling [40]. A conserved cytoplasmic protein, Lowfat and its human homologs LIX1 and LIX1-like, bind to the cytoplasmic domain of Ft and modulate Ft signaling by influencing the sub-apical membrane localization of Ft [41]. The Casein Kinase Discs overgrown (Dco) can bind to and phosphorylate the cytoplasmic domain of Ft, and control Ft signaling in both growth and PCP [42,43]. However, it has not been examined whether these interaction regions in Ft are crucial for regulating in PCP and tissue growth in vivo.

In order to further investigate the downstream signaling pathways regulated by Ft, we sought to identify regions of Ft that are essential for Ft function. Here, we identify a crucial region (H2 region) in the Ft cytoplasmic domain that, when membrane localized, is both necessary and sufficient to support viability and limit tissue size in animals lacking wild-type Ft function. Interestingly, this region is dispensable for PCP, indicating that Ft uses different regions to regulate PCP and tissue size. Moreover, the crucial regions of Ft identified in our structure-function study do not overlap with the previously reported interaction regions with Atrophin, Dco, or Lowfat.

While we are preparing for the manuscript, a similar structure-function study was published based on analysis in the wing [44]. In comparison, the conclusion from this report is largely in agreement with our results with a few exceptions. Both studies have mapped to the same region in the cytoplasmic domain of Ft that is required to support viability and tissue size regulation, but dispensable for PCP. Surprisingly, in our observation, expression of the H2 region does not have discernible ability to activate Hpo/Wts signaling in the eye. In addition, the extracellular domain of Ft is essential for PCP signaling in the eye, but dispensable in the wing, consistent with previous observations of tissue-specific difference for regulating Ft PCP signaling [3,45]. Together, our study in the eye and the recent report in the wing support the idea that Ft regulates PCP and tissue growth through separated functional regions.

**Materials and Methods**

**Fly Stocks**

The following stocks were used: ft/+ [10] and ft(2) [11], E(fly)mob10.5 (R4 cell fate marker) [46], ft-lacZ [47]. The Ga4 drivers used were TubP-Gal4 [48], Omb3-Gal4 and Vgl-Gal4 [49], and Flip-out Gal4 (ActP>TOP, y+>Gal4) [50]. FLP/FRT [51] was used to generate mutant clones in imaginal discs. All rescue experiments were performed in ft null mutant combination ft(-/-) ft(2)/ft(2)N. UAS-FtΔECD [52].

**Expression Constructs**

The full-length Ft (Ft-FL) construct was generated as previously described [3]. Ft-FL, FtΔcyt, FL-H2, FL-H345 were placed into the pUASP vector, while all the intracellular mutants were in the pUAST vector [53,54]. Each mutant construct has a V5 tag at the C-terminus. The V5 tag does not interfere with function as full-length Ft with or without the V5 tag provides the same degree of rescue in lethality, disc size, DIAP1 expression and PCP assays (data not shown). Each intracellular domain mutant construct has a myristylation sequence at N-terminus. The myristylation sequence used was: MGKCCSKRQ [55]. The mutated version (Myr-mut) of Ft-intra construct contains a (GA) point mutation in myristylation sequence, which inactivates the myristylation signal. FtΔECD is described in [52].

Amino Acids (a.a.) numbering is based on the protein sequence from Swiss-Prot: P33450. Sequences of Ft mutants are: Ft-FL: 1–5147a.a., FtΔcyt: 1–4624a.a., FL-H2: full length with deletion of 4719–4900a.a., FL-H345: full length with deletion of 4979–5147a.a., Ft-intra: 4612–5147a.a., FL-intra: 4612–5147a.a., MH2345: 4719–4900a.a., MH1234: 4719–4900a.a., MH45: 4967–5147a.a., MH345: 4897–5147a.a., MH2345: 4719–5147a.a., MH2: 4719–4900a.a., MH3: 4719–4900a.a., MH4: 4967–5147a.a., MH1: 4719–4900a.a., MFT: 4897–5147a.a., MAMH: intra-cellular domain sequence with deletion of 4719–4900a.a. Complete sequences of those constructs are available on request.

**Sections and Immunohistochemistry**

Tangential sectioning of adult eyes was performed as previously described, except that the osmium steps were omitted [56]. Third instar eye discs were dissected in cold PBS and fixed in 4% formaldehyde for 45 min on ice. Antibodies used were as follows: Mouse anti-DIAP1 (1:200; a gift from B. Hay), Rat anti-CyC-E (1:400; a gift from H. Richardson), Mouse anti-β-gal (1:1000; Promega), Rabbit anti-GFP (1:1000; Molecular Probes), Chicken anti-GFP (1:1000; Molecular Probes), and Rabbit anti-V5 (1:5000, Sigma), Cy3, Cy5, and FITC-conjugated secondary antibodies (Jackson Lab) were used at 1:200 dilution. Fluorescein phalloidin was used at 1:100 dilution. Confocal pictures were taken with a Biorad MRC 1024 microscope.

**Immunoblotting**

Third instar larval imaginal discs were lyzed and boiled in protein sample buffer. Samples were subjected to SDS-PAGE, transferred to Hybond-P membrane, and probed with Rabbit anti-V5 (1:10,000; Sigma), Rabbit anti-DIAP1 (1:5000; a gift from K. White) or Mouse anti-β-tubulin (1:10,000; DSHB). Secondary antibodies used were Rabbit and Mouse anti-HRP.

For comparisons of DIAP1 protein levels, 20–30 eye imaginal discs were collected for each particular genotype for each immunoblot analysis. Using NIH ImageJ software, DIAP1 levels were quantified and normalized to β-tubulin, then normalized to the value in wt animals. Quantification result represents the mean of 3 independent experiments. Error bars show ± S.D. To assess statistical significance, data was analyzed by one-way ANOVA with post-hoc Bonferroni correction. Significance was placed at p<0.05.

**Results**

**Membrane Localized Cytoplasmic Domain of Ft is Essential to Support Viability and Limit Tissue Size**

In order to understand how Ft transduces growth regulatory signals to downstream pathways, we sought to identify regions of Ft that are essential for regulating tissue size. A series of deletion mutants of Ft were constructed, expressed in vivo, and assayed for their ability to rescue both the pupal lethality and eye imaginal disc overgrowth phenotypes associated with ft mutant animals (ft(2)N/ft(2)N). For this analysis, ubiquitous expression of the deleted Ft proteins was driven using the UAS/Gal4 system and a ubiquitous Tub-Gal4 driver. For each deletion mutant transgene, 2–3 independent insertions were analyzed. Schematic structures of Ft mutants and assay results are summarized in Figure 1. As expected, expression of the full-length wild-type Ft protein (Ft-FL) fully rescued pupal lethality (Figure 2A).

Examination of late 3rd instar eye discs from the rescued animals further showed that eye imaginal discs of rescued animals were normal in size compared to overgrown discs of mutant animals (Figure 2B&E). We began our next analysis with the generation of Ft proteins in which either the entire intracellular domain was deleted and replaced by a V5 epitope tag (FtΔcyt) or in which the extracellular...
and transmembrane domains were replaced by a membrane targeting Src myristylation motif (Ft-intra) (Figure 1). Consistent with a previously published report [52], we found that ubiquitous expression of Ft-intra was sufficient to rescue both the lethality and disc size overgrowth phenotypes of ft mutant animals (Figure 1F).

We further found that expression of either Ft-FL or Ft-intra under Eyeless-Gal4 control could restore eye imaginal disc size to normal, but not that of other tissues (Figure S1). This indicates that the ability of Ft to regulate tissue size is a tissue-autonomous function.

In contrast to the ability of Ft-FL and Ft-intra to regulate tissue size, expression of FtD cyt did not support the viability of ft mutant animals. Examination of mutant larvae expressing FtD cyt showed that eye discs appeared to have severe patterning defects and thus could not easily be evaluated for their overgrowth phenotype (Figure 2G). To assay the ability of FtD cyt to regulate disc size, the size of wing discs from ft mutant animals expressing FtD cyt was examined. While expression of the Ft-FL or Ft-intra in ft mutant animals restored their wing discs to normal size, the wing discs of ft mutant animals expressing FtD cyt remained overgrown (Figures 2I–N). Taken together, these results indicate that the intracellular domain of Ft contains essential regions necessary for supporting viability and limiting tissue size.

The ability of membrane targeted Ft-intra to rescue the ft mutant phenotype raised the question of whether localization to the plasma membrane is required for Ft function in these rescue experiments. Figure 1. Summary of Ft constructs and their ability to rescue ft mutant phenotype. Schematic structure of Ft mutant constructs is on the left with their short names. (See Materials and Methods for details.) Summary of rescue results is on the right. Measurement and quantification of each rescue assay can be found in indicated figures. In the lethality rescue assay, effective rescue (+) was determined by at least 25% recovery of the expected number of ftG2rv/ftl(2)fd; TubP-Gal4/UAS-transgene adults, whereas no rescue (−) was determined for those transgenes which did not produce any adult animal (eclosion). In the eye disc size rescue assay, effective rescue (+) was determined by >20% increase of size compared to wt disc. In the PCP rescue assay, effective rescue (+) was determined by >85% correct orientation of ommatidia; (−) indicates disorganized polarity that the equator was unable to be recognized, and <75% correct orientation can be determined. In the clonal DIAP1 and clone size assays, effective rescue (+) was determined by >60% of clones with successful rescue of clonal phenotype and DIAP1 expression level. ‘DN’ indicates dominant negative effect of FL-DH2. N.D.: not determined.

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Structure and Function Study of Ft
assays. To address this issue, a mutant form (Myr-mut) of Ft-intra was generated with a point mutation in the myristylation sequence that blocks myristylation and thus prevents membrane localization. This mutation abolishes Ft function in both the lethality and imaginal disc size rescue assays (Figures 2H, N), indicating that membrane localization is crucial for Ft function.

A Small Region of the Intracellular Domain can Provide Ft Function for Regulating Viability and Tissue Size

To identify crucial regions within the intracellular domain of Ft, a series of deletion mutations that removed portions of Ft-intra were constructed and assayed for function. To guide this process, the cytoplasmic sequence of Drosophila Ft was compared to its homologs from other species. Sequence alignment reveals several highly conserved blocks of amino acids, referred to as conserved blocks 1–5 (Figure 3A). A series of mutant constructs were then generated in which the regions containing the conserved blocks were successively deleted from the N- or C-terminus of the Ft-intra protein (Figure 3B). Since membrane localization is required for growth control, each mutant protein was directed to the plasma membrane by a myristylation sequence at its N-terminus. In addition, the presence of a V5 epitope tag at the C-terminus of each protein allowed us to demonstrate that the various mutant proteins were expressed at similar levels (Figure 3C). Truncated proteins were then tested for their ability to rescue the lethality and disc overgrowth phenotype. The ability of the different truncated proteins to rescue / mutant animals is summarized in Figures 3B, and examples in 3D–F.

This functional analysis led to the identification of a small region (H2) that is crucial for supporting viability and limiting tissue size (Figure 3B). Importantly, each of the truncated proteins lacking H2 was unable to rescue / mutant phenotypes. In contrast, every protein containing the H2 region retained rescuing ability. The striking correlation between the presence of H2 and rescuing ability suggested that the H2 region might be both necessary and sufficient for effective Ft function. In order to test these possibilities directly, two additional mutant constructs were generated. In one, only the H2 region was deleted from Ft-intra (MΔH2). This protein failed to provide Ft function. In the second, only the H2 region of Ft was present in the expressed protein (MH2). This protein provided effective rescuing function. Taken together, these results indicate that the H2 region is both necessary and sufficient for Ft function in these assays. Consistent with the importance of the H2 region, its deletion from the full-length Ft protein (FL-DH2) effectively abolishes rescuing ability (Figure 3F).

Regions Crucial for PCP Signaling in the Eye

In addition to regulating viability and tissue size, Ft also plays a key role in the establishment of epithelial planar cell polarity (PCP) [1,2,3]. In the eye, planar cell polarity is apparent in the
Figure 3. The H2 region is necessary and sufficient for viability and tissue size regulation. (A) Sequence alignment of cytoplasmic domain of Drosophila Ft (538 a.a.) and human Ft4 (452 a.a.) reveals highly conserved regions (conserved block 1–5) shown in blue boxes. The starting and ending positions of the deletion mutants are indicated by red bars. (B) Schematic structure of Ft intracellular domain deletion mutants is on the left. Myristylation sequence is indicated in orange. Conserved homology regions are indicated in blue. V5 tag is indicated in pink. Regions deleted are shown with black lines. Quantification of the lethality rescue and disc size rescue on the right. (See Fig. 2AB for method details) (C) Expression levels of intracellular domain deletion mutants are revealed by an immunoblot of imaginal disc extracts from ft mutant animals expressing various transgenes. Blots were probed with anti-V5 and anti-
\[\beta\]-tubulin. (D–F) Deletion of the H2 region in either the intracellular domain (E) or the full-length Ft protein (F) effectively abolished rescuing ability, whereas the H2 region, when membrane localized (D), is necessary and sufficient to rescue both
orientation of the ommatidial units that comprise the compound eye. In wild type, all of the ommatidia in the dorsal half of the eye are oriented in the same direction while those in the ventral region uniformly adopt the opposite chiral form (Figure 4A, A'). The establishment of this ordered pattern of polarity is dependent on Ft, whose absence leads to a randomized pattern of dorsal and ventral type ommatidia throughout the eye (Figure 4B, B').

The importance of the intracellular domain (Ft-intra) for regulating viability and tissue size naturally raised a question as to whether Ft-intra is also necessary and sufficient for planar cell polarity signaling. To test this hypothesis, we carried out an eye PCP rescue experiment with Ft-intra. We have previously shown that ubiquitous Tub-Gal4 driven expression of Ft can fully re-establish the wild-type pattern of ommatidia in ft mutant animals (error rate of <1%) [3]. This assay allowed us to examine whether Ft-intra could provide PCP function in ft mutant animals.

We found that uniform expression of Ft-intra cannot restore the overall pattern of polarization (Figure 4E, E'). A different version of the intracellular domain (FtΔECD) [52], which has the endogenous transmembrane and cytoplasmic domains, also fails to rescue the pattern of polarization (Figure 4F, F'). These findings indicate that in the eye, the Ft intracellular domain alone is not sufficient to provide all the function of Ft in PCP, suggesting that the extracellular domain is essential for receiving and/or directing the downstream PCP signaling. This is in marked contrast to the observation in the wing, where the Ft transmembrane and cytoplasmic domains together are sufficient to mediate PCP signaling [44,52].

Ft's Role in PCP is Independent from Viability and Tissue Size Regulation

Although Ft-intra is not sufficient to rescue the PCP defect, it may still contain key regions necessary in PCP signaling. This is particularly interesting for the H2 region, given its key role for rescuing viability and tissue size. To test this hypothesis, and to extend it to the other regions of Ft, we tested two full length mutants, FL-DH2 and FL-DH345, for their ability in PCP signaling. Since FL-DH2 expression does not rescue ft mutant animals to adulthood, our analysis was conducted in 3rd instar eye discs. We found that the overall pattern of PCP is rescued when either FL-AH2 or FL-DH345 is expressed (Figure 4D, D', G, G'). In each case, only a low rate of polarity errors was observed. This finding indicates that neither the H2 region nor the H345 region is essential for directing PCP in the eye. However, the somewhat elevated occurrence of polarity mistakes may suggest that the strength or quality of information is slightly reduced when the structure of the cytoplasmic domain is impaired. These results suggest that Ft regulates PCP through an effector region that is distinct from that used during growth control, and more importantly, that Ft's roles in PCP signaling and tissue size control are separable and can be carried out independently. Consistent with this proposal, while MH2 sufficiently rescued ft animals to adulthood without restoring the normal pattern of planar cell polarity, expression of MH2 along with FL-AH2 can produce rescued adult animals with normal ommatidial organization (Figure 4H, H').

The Function of Ft to Regulate Tissue Size and Viability is Separable from its Ability to Regulate Clone Size and Hpo/Wts Signaling

The requirement of Ft function in growth control can be seen in two different settings: the overall tissue size increase in the ft mutant animal and the clonal overgrowth in the mosaic tissue. Although expression of MH2 is sufficient to rescue tissue size and support viability in ft mutant animal, it surprisingly failed to rescue the clonal phenotype of ft mutant cells in the mosaic tissue (Figure 5E). In a more quantitative comparison, expression of the wild-type Ft (Ft-FL) restored the size and shape of ft clones to normal, ft mutant clones expressing MH2 are still large in size and round in shape, not significantly different from those lacking Ft function (Figure S2). These results suggest that the essential function of Ft for regulating viability and tissue size is separable from its regulation of clone size.

Several reports have demonstrated that the Hpo/Wts signaling pathway is a target of Ft regulation in growth control [17,18,19,20]. This was most clearly shown in the mosaic animal in which clones of cells lacking Ft function have elevated Hpo/Wts signaling, resulting in the increase of clone size. The ability of Ft to regulate the Hpo/Wts pathway and the importance of the Hpo/ Wts pathway for controlling cell proliferation and cell death has led to the idea that the essential function of Ft in supporting viability and preventing tissue size overgrowth is to stimulate the Hpo/Wts pathway and thus limit Yki activity.

Because we have observed that expression of MH2 provides sufficient Ft function to regulate tissue size and support viability without rescuing ft clonal overgrowth, this raises a question of whether expression of MH2 is able to restore Ft-mediated activation of the Hpo/Wts pathway. We first examined this question at the whole tissue level. Since it is difficult to compare the precise level of DIAP1 expression between eye discs by immunofluorescence, we used western blot to compare DIAP1 protein levels with β-tubulin as an internal control. While we observed a significant increase of DIAP1 in both ft mutant discs and MH2 rescued eye discs compared to wt discs, we could not detect significant difference of DIAP1 between ft mutant and MH2 rescued eye discs (Figure S2G, H).

We then tested this question at the clonal level by examining whether ubiquitous expression of MH2 prevents the elevated expression of Yki transcriptional targets, such as DIAP1, fj-lacZ, and CyeC that would otherwise occur in clones of ft mutant cells (Figure 5). While ubiquitous expression of either Ft-FL or Ft-intra restored DIAP1 expression levels in ft clones to normal (Figures 5B, C), expression of MH2 appeared to have no effect on elevated DIAP1 expression in ft clones (Figure 5E). MH2 similarly failed to affect elevated CyeC and fj-lacZ (Figure S3, S4). These results are consistent with the failure of MH2 to regulate clonal phenotype of ft mutant cells, suggesting that MH2 lacks the ability to restore Ft regulation of Hpo/Wts signaling.

Since the ability to detect MH2 effect on the Hpo/Wts pathway may be limited by the need to make comparisons between discs, the possibility remained that MH2 might still regulate the Hpo/Wts pathway, albeit much more weakly than Ft-FL. In order to rigorously examine this possibility, we compared ft mutant clones that expressed MH2 to ones from the same eye disc that did not express MH2. For this experiment, ft mutant clones were generated in eye imaginal
Figure 4. The H2 region is not required for Ft in regulating PCP signaling. Schematic diagrams (A–H) of equatorial sections of adult eyes (A’, C’, D’, E’, F’, H’), or confocal images of third instar eye imaginal discs (B’, G’). (B’, G’) Ommatidial orientation was determined by examining the expression of E(spl)/tm005 (R4 marker) in each ommatidium. This marker consists of a portion of the enhancer region of the E(spl) gene, a transcriptional target of N activation, fused to a β-galactosidase reporter. This marker is strongly expressed in R4 cell (green) [46]. Elav staining (red) labels all photoreceptor cells in each ommatidial cluster. All images are oriented anterior to the left and dorsal up. The equator is indicated by the black line. Green arrows indicate ommatidia with dorsal chirality and orientation, whereas red arrows designate ventral type ommatidia. Yellow dots represent either incorrectly constructed ommatidia, ommatidia exhibiting severe misrotation or ommatidia with undetermined polarity. Relevant changes from wild type with regard to Ft function or expression are indicated at the bottom of each panel. Genotypes are listed below along with the fidelity rate (the percentage of ommatidia that are correctly constructed and rotated). Ubiquitous expression of Ft-FL, FL-H2 or FL-ΔH2Δ5 can restore the wild-type pattern of ommatidia in ft mutant animals. FL-H2Δ2 and FL-ΔH2Δ45 have a low rate of errors. (A, A’) w1118 (100%) (B, B’) w1118; ft–/– (Disorganized, n = 430) (C, C’) w1118; ft–/–; Tub-Gal4/UAS-FL-FL (99%), n = 788) (D, D’) w1118; ft–/–; Tub-Gal4/UAS-FL-FL-H345 (91.1%, n = 529) (E, E’) w1118; ft–/–; Tub-Gal4/UAS-FL-FT-intra (Disorganized, n = 1592) (F, F’) w1118; ft–/–; Tub-Gal4/UAS-FL-FT-ECOD (Disorganized, n = 907) (G, G’) w1118; ft–/–; Tub-Gal4/UAS-FL-FT-H2 (93.5%, n = 660) (H, H’) w1118; ft–/–; Tub-Gal4/UAS-FL-FT-H2 & UAS-MH2 (91.5%, n = 471).

discs in which the flip-out GAL4 system had been used was independently to generate independent clones of cells expressing MH2 under actin5C promoter control [50]. The resulting discs contained independent patches of ft mutant cells and MH2 expressing cells, which sometimes overlapped. As shown in Figure 6B and B’, the elevated DIAP1 level of ft mutant clones that expressed MH2 was quantitatively indistinguishable to that from ft mutant clones without expressing MH2. As expected, ft mutant clones that expressed Ft-FL effectively rescued DIAP1 expression level in this assay (Figure 6A). Together, these results indicate that the MH2 protein has little, if any, ability to activate the Hpo/Wts signaling pathway.

Interestingly, the protein lacking H2 region (FL-ΔH2) not only failed to suppress elevated DIAP1 expression levels in ft mutant clones, also induced dominant negative effect on Hpo/Wts signaling, as evident by the elevated level of DIAP1 in wt cells overexpressing FL-ΔH2 (Figure 6C). This dominant negative effect was also apparent when clones of cells that overexpressed FL-ΔH2 were generated in otherwise wild-type animals (Figure 6E), indicating that FL-ΔH2 lacks the ability to provide the Hpo/Wts regulatory function, also dominantly inhibited the Hpo/Wts signaling ability of the endogenous wild-type Ft protein.

While the results described above demonstrate that expression of the H2 region alone is not sufficient to regulate Hpo/Wts signaling, they neither addressed whether the H2 region is required for this function nor identified other regions Ft that contribute to Hpo/Wts regulation. We investigated these issues by examining clonal size and DIAP1 expression in ft mutant clones generated in animals ubiquitously expressing various mutant forms of Ft proteins (Figure 5). Since most ft mutant constructs cannot rescue the clonal phenotype and elevated DIAP1 expression, our interpretation is mainly focused on those with successful rescue. First, the expression of FL-ΔH2Δ45 effectively rescues Yki target gene expression and the ft clonal overgrowth phenotype, suggesting that the H345 region is not essential for this process. Second, the expression of Ft-intra and MH2Δ345 effectively suppresses the enhanced expression of Yki targets in ft clones, suggesting the extracellular domain and the H1 region is not necessary. Third, the comparison between MH12 and FL-ΔH345, and the comparison between MH2345 and MH2 suggest that the presence of the extracellular domain and the H345 region may significantly enhance the signaling strength of the protein in regulating the Hpo/Wts pathway. Taken together, these results indicate that the H2 region, although not sufficient, is necessary for regulating the Hpo/Wts pathway, whereas H1, H345 and extracellular domain are not essential, but could potentially provide important supporting role for Hpo/Wts pathway.

Discussion

Separation of the Growth Control and PCP Function of Ft

As the involvement of Ft in both tissue growth and PCP raises the question as to whether Ft regulates the two processes through the same or different regulatory domains. Here we identified two mutant proteins, and each harbors complementary properties. The H2 region is essential for tissue growth, but is completely dispensable for PCP signaling. The FL-H2Δ2 protein lacks regulatory function for tissue growth but retains full function in PCP. This strongly indicates that roles of Ft in tissue growth regulation and PCP signaling are independent, and each can be carried out by a separate effector region.

There are several reported binding partners of Ft cytoplasmic domain, which have been reported to mediate some function of Ft [40,41,42,43,57]. The key interaction regions are indicated in Figure 1. However, these regions are not essential for Ft function in tissue size, PCP and Hpo/Wts activity. This is best shown by one mutant Ft protein, FL-ΔH345, which lacks the reported binding regions, but is still sufficient to rescue Ft functions in these processes. This result indicates that these binding partners are not likely the primary effectors, but modulators in mediating downstream signaling.

Different Requirement for Ft Function in PCP Signaling between the Eye and Wing

In the eye, we showed that the crucial region of Ft for PCP signaling includes the extracellular domain. In the absence of the extracellular domain, the eye displays severe PCP defects. This result is consistent with the model in previous in vivo and in vitro studies that Ft and Ds can form a heterodimer through their extracellular domains and this interaction is essential for the ordered PCP pattern [1,45]. Second, this result draws a clear distinction between the eye and wing. In the wing, the transmembrane and cytoplasmic domains of Ft are sufficient to rescue the PCP defect in the wing, suggesting this protein contains all the effector regions for PCP signaling [52]. Since the extracellular domain of Ft is required to interact with Ds to receive the upstream directional information in the eye, the dispensability of the extracellular domain during the polarization of the wing indicates that this directional information may not be essential in the wing. This is consistent with the result shown in a previous study that the Ds and Fj gradients are essential for the eye, but do not play a major role in directing PCP in the wing [3]. It is worth pointing out that the dispensability of Ds and Fj gradients is NOT equivalent to the removal of Ds function, as Ds has an essential function to stabilize and/or activate Ft on the plasma membrane in addition to its role of providing a directional cue. Together, these observations suggest a model in which Ft needs Ds to activate its PCP function; and
both the activation and its graded fashion across the tissue are required in the eye to provide a directional cue, whereas in the wing this activation is not necessary to be graded. Thus, this study supports the notion that the eye and wing have different requirements for Ft function, indicating different strategies used to direct PCP.

Ft’s Role in Tissue Size Regulation is Genetically Separable from its Function in Clone Size Regulation

Role of Ft in regulating growth can often be revealed in two different settings. First, animals lacking Ft function exhibit massive overgrowth of all imaginal discs. Second, in mosaic animals, cells lacking Ft function proliferate much more extensively than their wild-type neighbors, resulting in large clones of mutant cells. While each of these observations suggests that cells lacking Ft are
unable to respond to signals that normally limit their growth and proliferation, they may reflect different roles of Ft. The increase of clone size may result from mutant cells possessing a competitive advantage over their neighbors, such as an increased rate of proliferation or resistance to cell death. In contrast, the imaginal disc overgrowth in animals lacking Ft function may indicate that ft mutant cells are unable to sense or respond to the signals which normally restrain tissue growth and thus help specify tissue size. Although these two properties can often be affected by the same cellular events, they are not always correlated. In this study, we indeed observed the dissociation of these two aspects of Ft function. This is most clearly shown by MH2, which is sufficient to rescue tissue size overgrowth but lacking the ability to rescue clonal overgrowth defect.

One downstream pathway of Fat is to stimulate the activity of the Hpo/Wts signaling pathway, thus leading to inhibition of the transcriptional activator Yki and reduced expression of its targets [14,36]. These targets include important positive regulators of growth and negative regulators of apoptosis such as CycE and DIAP1. The ability of Ft to regulate the Hpo/Wts pathway and

Figure 6. The H2 region has no detectable activity in the Hpo/Wts pathway. ft mutant clones, marked by the absence of GFP expression, were generated in third-instar eye imaginal discs, in which Ft mutant transgenes (blue): Ft-FL (A) and MH2 (B) were expressed after flip-out of ActP>STOP, y+>Gal4. Arrowheads point to ft mutant clones with expression of Ft transgenes. Arrows point to ft mutant clones alone. Eye imaginal discs were stained with GFP (green), DIAP1 (red), and V5 (blue). (A) Expression of Ft-FL suppressed elevated DIAP1 expression levels in ft clones. (B) Expression of MH2 failed to suppress elevated DIAP1 expression levels in ft clones. (B') DIAP1 expression level in the ft clone was not significantly different from that of ft clone expressing MH2 in the same disc. DIAP1 expression level in the center of the clones (indicated in the yellow boxes in 6B) were quantitatively measured using histogram function of Adobe Photoshop, and normalized. n=5. Two-tailed Student’s t-test. (C) Expression of FL-ΔH2 in ft clones did not suppress elevated DIAP1 expression levels. Cells in FL-ΔH2 expressing clones exhibited elevated levels of DIAP1 expression (E), whereas cells in Ft-FL (D) expressing clones did not have a dominant negative effect.

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the similar clonal overgrowth phenotypes of animals lacking either Ft or Hpo/Wts pathway activity has led to the idea that the essential function of Ft in supporting viability and preventing tissue overgrowth is to activate the Hpo/Wts pathway. Indeed, genetic interactions between ft and the Hpo/Wts pathway are consistent with this idea. For example, the imaginal disc overgrowth phenotype of ft mutant animals is partially suppressed and enhanced by heterozygosity for Yki and Wts mutations, respectively [20].

Given the strength of these arguments, our expectation was that the minimal region of Ft required for regulating tissue size and supporting viability would also be capable of regulating the expression of Yki targets. Surprisingly, our analysis demonstrates that the ability to regulate tissue size and viability does not correlate with the ability to repress Yki activity. This is most clearly seen in the ability of the H2 region, when membrane-tagged, to rescue viability and tissue size without having any detectable ability to regulate Yki target expression. Thus, these data argue that the essential function of Ft in viability and tissue size regulation is genetically separable from its role in Hpo/Wts pathway regulation. In agreement with our observation, a previous study has reported growth defect in ft mutant tissue that cannot be explained by loss of expanded (ex), an upstream component of the Hpo/Wts pathway, suggesting that there are other, as-yet-unidentified, mediators of ft [18].

Previous research has shown that regulation of tissue growth and proliferation by Ft is complex [17,18,19,20,36,58,59]. The complex nature of the signaling network may reflect multiple levels of regulation in tissue growth and proliferation control. It is possible that multiple mechanisms at different levels regulate distinct aspects of tissue growth. Indeed, our studies support such a model. During the control of tissue growth and proliferation, Hpo/Wts signaling may serve primarily as a local regulatory mechanism to coordinate cell proliferation and cell death among neighboring cells, whereas an unidentified pathway mediated by Ft may play a more crucial role in allowing the tissue as a whole to regulate its size. Although not favored by our data, the possibility cannot be excluded that dissociation of tissue size control from clone size regulation is due to quantitative difference in Hpo/Wts signaling. However, within the sensitivity range of all the in vivo assays used in this study, we could not detect any activity of MH2 in regulating the Hpo/Wts pathway. Therefore, our results are more consistent with the model in which an unknown pathway mediated by Ft regulates tissue size. To completely understand how Ft fulfills these roles and the underlying biochemical mechanisms will likely require the identification of key binding partners for the crucial H2 region.

While this manuscript was under preparation, a similar structure-function analysis was published based on the analysis of Ft function in the wing [44]. In agreement with our analysis, Matakatsu et al. have found a region (termed Hpo-active domain) required for viability, tissue size regulation and Hpo/Wts signaling activity, which largely overlaps with the H2 region. The study from Matakatsu et al. did not report any mutant construct that can clearly rescue viability but not activate the Hpo/Wts signaling pathway. However, we have observed such phenomenon that expression of the H2 region is not sufficient to activate Hpo/Wts signaling in our assays even though the H2 region is fully capable to support viability and limit tissue size. Several differences in the experimental set up between the two studies may contribute to the difference in observation. Besides the tissue difference between the wing and eye, the two studies used different version of the truncation mutants of Ft, different pathway readouts for Hpo/Wts signaling, different Gal4 drivers, and different tissue locations to examine the effect on growth and Hpo/Wts signaling. In the future, it will be interesting to test whether such phenomenon can be observed across different tissues. Comparing to the required regions for PCP in the wing, we have demonstrated that the extracellular domain of Ft is essential for PCP in the eye. This difference reflects different strategies to organize PCP between eye and wing [3,45,52]. Finally, both studies support the idea that the functional region for PCP is distinct from the region for tissue growth regulation.

During the revision of this manuscript, another structure-function study of Ft was published based on the analysis in the wing and abdomen [60]. Using a complementary approach - small deletion or point mutations in ft genomic constructs rather than UAS-Gal4 system, Pan et al. also reached the same conclusion that effects of Ft on wing growth versus PCP can be separated at the level of Ft itself. Interestingly, one unique finding from Pan et al. indicates a 4 amino acids C-terminal motif that is essential for Ft-mediated PCP in certain regions of the wing and abdomen. Intriguingly, we found the mutant Ft without C-terminal region (FL-AH345) is still capable of directing PCP in the eye with a fidelity rate of 91.1%. In the future, it would be interesting to examine how much of the reduced fidelity rate of FL-AH345 in the eye is because of missing this 4 amino acids motif.

**Supporting Information**

**Figure S1** Ft regulates tissue size through a tissue-autonomous manner. Expression of either Ft-FL or Ft-intra under Eyeless-Gal4 control restore eye imaginal disc size to normal, but not that of other tissues, such as wing discs. (A–D) Third-instar eye imaginal discs stained with fluorescein phalloidin. (A’–D’) Third-instar wing imaginal discs. Genotypes are: (A, A’) wt, (B, B’) ft<sup>2</sup>/–, (C, C’) ft<sup>2</sup>/–; Ey-Gal4/UAS-Ft-FL, (D, D’) ft<sup>2</sup>/–; Ey-Gal4/UAS-Ft-intra (PDF)

**Figure S2** ft mutant clones expressing MH2 are still large in size and round in shape, similar to those lacking Ft function. (A–D) Examples of clone/twin-spot pairs showing increased size and round shape of ft mutant clone compared to its twin-spot. This clonal phenotype is fully rescued by the ubiquitous expression of Ft-FL, but not significantly affected by the expression of MH2. (A) wt clones (marked by loss of GFP, indicated by white arrow) are similar in size and shape to their twin spot (GFP/GFP, indicated by yellow arrow). (B) ft clones (marked by loss of GFP, indicated by white arrow) are larger than their twin spot (indicated by yellow arrow) in larval eye discs. (C) Ubiquitous expression of wild-type Ft (Ft-FL) with TubP-Gal4 restored the size and shape of ft clones to normal. (D) ft clones expressing MH2 retained the large size and round shape. (E) Quantification of the ratio of clone size/twin spot. Clone and twin spot areas were measured using the histogram function of Adobe photoshop. Black bar represents average ratio. Average ratio was 0.94 for wt, 6.56 for ft clone, 0.85 for ft clone+Ft-FL and 5.94 for ft clone+H2. (F) Quantification of the ratio of clone shape index/twin spot. Shape of each individual clone or twin spot was determined by clone shape index (A/L<sup>2</sup>) of roundness (L<sub>2</sub> circumference of the clone, A: area of the clone). The rational for using A/L<sup>2</sup> formula was as follows: for a given number of cells in a clone, the stronger the tendency of these cells to avoid mixing with cells located outside of the clone, the smaller the circumference (L) would be relative to the area (A) [61], and the larger the clone shape index (A/L<sup>2</sup>). The area (A) and the circumference (L) of a clone or twin spot were measured using the histogram function of Adobe photoshop. Black bar represents average ratio. Average ratio was 1.12 for wt, 2.89 for ft clone, 1.04
for ft clone+Fl-FL and 2.40 for ft clone+H2. (GH) Immunoblot analysis of DIAP1 expression in eye imaginal disc extracts from wt, ft mutant, and ft mutant animals expressing Ft-FL or MH2 transgenes. Blots were probed with anti-DIAP1 and anti-β-tubulin. DIAP1 expression levels in the ft mutant and MH2 rescued discs are significantly increased compared to the wt and Ft-FL rescued discs (p<0.05). No significant difference can be detected between the ft mutant and MH2 rescued discs. DIAP1 levels were quantified and normalized to the value in wt animals. An example of immunoblot is shown in (G). Quantification result in (H) represents the mean of three independent experiments and error bars show S.D. Statistical significance was analyzed by one-way ANOVA with post-hoc bonferroni correction. (PDF)

Figure S3 The H2 region is not sufficient to rescue elevated Cyclin E expression in ft clones. ftΔ249 mutant clones, marked by the absence of GFP expression (green channel), were generated in third-instar eye imaginal discs in which Ft mutant transgenes were expressed by TubP-Gal4. β-gal staining was shown in red. (A) ft mutant clones show autonomous increases in β-gal. Ubiquitous expression of Ft-FL (B), and Ft-intra (C) rescues the elevated β-gal levels in ft clones, whereas MH2 (D) and MAH2 (E) do not rescue (see result section). (PDF)

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Figure S4 The H2 region is not sufficient to rescue elevated β-gal expression in ft clones. ftΔ249 mutant clones, marked by the absence of GFP expression (green channel), were generated in third-instar eye imaginal discs in which Ft mutant transgenes were expressed by TubP-Gal4. β-gal staining was shown in red. (A) ft mutant clones show autonomous increases in β-gal. Ubiquitous expression of Ft-FL (B), and Ft-intra (C) rescues the elevated β-gal levels in ft clones, whereas MH2 (D) and MAH2 (E) do not rescue (see result section). (PDF)

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Author Contributions
Conceived and designed the experiments: XZ CHY MAS. Performed the experiments: XZ CHY MAS. Analyzed the data: XZ CHY MAS. Wrote the paper: XZ MAS.
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