Tissue Specific Roles for the Ribosome Biogenesis Factor Wdr43 in Zebrafish Development

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Abstract

During vertebrate craniofacial development, neural crest cells (NCCs) contribute to most of the craniofacial pharyngeal skeleton. Defects in NCC specification, migration and differentiation resulting in malformations in the craniofacial complex are associated with human craniofacial disorders including Treacher-Collins Syndrome, caused by mutations in TCOF1. It has been hypothesized that perturbed ribosome biogenesis and resulting p53 mediated neuroepithelial apoptosis results in NCC hypoplasia in mouse Tcof1 mutants. However, the underlying mechanisms linking ribosome biogenesis and NCC development remain poorly understood. Here we report a new zebrafish mutant, fantome (fan), which harbors a point mutation and predicted premature stop codon in zebrafish wdr43, the ortholog to yeast UTPS3. Although wdr43 mRNA is widely expressed during early zebrafish development, and its deficiency triggers early neural, eye, heart and pharyngeal arch defects, later defects appear fairly restricted to NCC derived craniofacial cartilages. Here we show that the C-terminus of Wdr43, which is absent in fan mutant protein, is both necessary and sufficient to mediate its nucleolar localization and protein interactions in metazoans. We demonstrate that Wdr43 functions in ribosome biogenesis, and that defects observed in fan mutants are mediated by a p53 dependent pathway. Finally, we show that proper localization of a variety of nucleolar proteins, including TCOF1, is dependent on that of WDR43. Together, our findings provide new insight into roles for Wdr43 in development, ribosome biogenesis, and also ribosomopathy-induced craniofacial phenotypes including Treacher-Collins Syndrome.

Introduction

Neural crest cells (NCCs), a transient cell type that is unique to vertebrates, originate from the dorsal aspect of the neural tube during embryogenesis. After undergoing an epithelial-to-mesenchymal transition (EMT), NCCs migrate along well defined pathways, and eventually inhabit peripheral destinations where they differentiate into diverse derivatives, including melanocytes, craniofacial cartilage and bone, smooth muscle, and neuronal lineages. In the head region, cranial neural crest cells (CNCC) give rise to nearly all craniofacial structures, including the facial skeleton and the vast majority of facial connective tissues [1,2]. Defects in CNCC development are associated with craniofacial malformations, one of the most common of human birth defects [3].

Treacher-Collins syndrome (TCS), an autosomal dominant congenital disorder of craniofacial development, characterized by mandibulofacial dysostosis including cleft palate and hypoplasia of the facial bones, is most commonly associated with mutations in the TCOFI gene [4]. Trecale, the protein encoded by the TCOFI gene, is a nucleolar phosphoprotein [5,6] that plays a key role in ribosome biogenesis via involvement in both rDNA methylation and rRNA transcription [7–10]. Extensive research in the mouse model has shown that mutations in Tof1 disrupt ribosome biogenesis, resulting in impaired proliferation and subsequent apoptosis of neuroepithelial and NCC precursors, which in turn results in reduced numbers of NCCs migrating into the developing craniofacial complex [10]. Interestingly, inhibition of p33 function can rescue craniofacial abnormalities in mouse Tof1 mutants, without rescuing ribosome biogenesis defects [11].

The question of how ribosome biogenesis defects can preferentially affect NCC proliferation and differentiation remains to be elucidated. In eukaryotic cells, ribosome biogenesis begins with the transcription of rRNA from rDNA located in the nucleolus, the most prominent visible structure in the nucleus. Ribosome biogenesis is extremely complex, requiring the accurate processing of pre-rRNAs into four different ribosomal RNAs (28S, 18S, 5.8S and 5.8S in vertebrates) and complex formation with about 80 constituent ribosomal proteins. In addition, more than 200 nucleolar ribosome biogenesis factors are required to complete
Here, we describe the identification and characterization of a novel zebrafish craniofacial mutant, fantome (fan), caused by a point mutation in the wdr43 gene. Although previously characterized as UTP5 in yeast, a nucleolar protein functioning in ribosome biogenesis, here we show that Wdr43 also regulates early zebrafish development, including NCC specification and differentiation. Mutations in nucleolar proteins have been found to be causative for a variety of human craniofacial syndromes including Treacher-Collins Syndrome (TCS), often caused by mutations in TCOF1, which also plays important roles in ribosome biogenesis. However, the underlying mechanisms linking ribosomal biogenesis and NCC specification and differentiation into pharyngeal arch cartilages remains poorly understood. Here we describe the fan/wdr43 mutant phenotype, and present functional characterizations of Wdr43 in craniofacial development. We show that WDR43 is required for the proper nucleolar localization of a variety of nucleolar proteins, including TCOF1/Treacel. These studies provide new insight into ribosomal protein function in early zebrafish development, with focus on NCC derived craniofacial development, as a model for human craniofacial neurocristopathies.

Here, we report a novel zebrafish mutant, fantome (fan), characterized by a variety of early developmental defects including eye, hindbrain, forebrain, cardiac, neurocranium, fin, and NCC derived pharyngeal arch cartilages development. NCCs in fan mutants fail to differentiate, and NCC precursors undergo p53 mediated cell apoptosis. fan mutants contain a point mutation in the zebrafish wdr43 gene, which encodes Wdr43, the ortholog of yeast Utp5. We demonstrate that, similar to the yeast ortholog, zebrafish and human WDR43 localize to the nucleolus. We also show that the C-terminal of Wdr43, truncated in fan mutants, mediates its localization to nucleoli, and is both necessary and sufficient to mediate its interaction with other t-UTP subcomplex members. Interestingly, blocking WDR43 expression in HeLa cells results in nucleolar maturation defects, together with abnormal localization of other nucleolar proteins, including TCOF1. Together, our data suggest that loss function of Wdr43 results in ribosome biogenesis defects that induce the p53 signaling pathway, triggering cell death of NCC precursors. We introduce the fan mutant as a valuable model to provide insight into a variety of human craniofacial neurocristopathy diseases.

Results

Phenotype of the fantome (fan) mutant

The fan mutant, identified in a large scale ENU chemical mutagenesis screen conducted by the Yelick Lab [29], was notable by its distinctive lack of pharyngeal arch cartilages at 4 days post fertilization (dpf) (Fig. 1D’, arrow). Subsequent developmental analyses showed that fan mutants are first identifiable at 16 hpf by a distinct area of necrotic cells present in the neural ectoderm of the presumptive eye (Fig. 1A’, arrow). At 24 hpf, a larger area of necrosis was detected mainly in neural tissues (Fig. 1B’, bracket), while at 30 hpf, fan mutants displayed distinct hydrocephaly in hind brain ventricles (Fig. 1C’, arrow), incomplete closure of the choroid fissure, and lack of pigmented retinal epithelium in the ventral eye (Fig. 1G’, arrowhead). Preliminary whole mount in situ hybridization (WISH) analyses of NCC marker gene expression revealed that fan mutants exhibited reduced snail2 and dlx2a expression at 12 and 24 hpf, respectively, as compared to age matched wild type sibling (Fig. 1E’, Fig. 1F’, arrow). At 96 hpf, Alcian blue staining revealed that homozygous fan mutants lacked virtually all pharyngeal arch cartilages (Fig. 1D’, arrow) as compared to age matched wild type siblings (Fig. 1D). Early lethal homozygous recessive fan mutants die at approximately 5–6 dpf, while heterozygous fan embryos and adults appear normal and are viable and fertile.

In normal zebrafish development, NCCs originating from the dorsal aspect of the neural tube migrate ventrally to the pharyngeal pouches and give rise to a variety of structures including pharyngeal arch cartilages [30] [31] [32]. To more carefully characterize the pharyngeal arch phenotype observed in fan mutants, we used WISH to examine the developmental expression of additional NCC markers, including: sox9a, essential for proper morphogenesis and differentiation of pharyngeal arch cartilages [33,34]; the pan-NCC marker crestin, expressed in pre-migratory and migratory NCCs [35]; hand2, expressed in branchial arch mesenchyme [36]; and dlx2a, which is expressed in migrating CNC that contribute to the pharyngeal arches [37]. We detected reduced expression of all NCC marker expression in fan mutants as compared to age matched wild type siblings (Fig. 1F’). To better visualize and study NCC defects in fan mutants, we also created a Tg(fish:EGFP)/fan mutant reporter line, which expresses EGFP in the derivatives of the cranial neural crest until at least 7dpf, and in developing vasculature [38]. We found that fan

Author Summary

Here, we describe the identification and characterization of a novel zebrafish craniofacial mutant, fantome (fan), caused by a point mutation in the wdr43 gene. Although previously characterized as UTP5 in yeast, a nucleolar protein functioning in ribosome biogenesis, here we show that Wdr43 also regulates early zebrafish development, including NCC specification and differentiation. Mutations in nucleolar proteins have been found to be causative for a variety of human craniofacial syndromes including Treacher-Collins Syndrome (TCS), often caused by mutations in TCOF1, which also plays important roles in ribosome biogenesis. However, the underlying mechanisms linking ribosomal biogenesis and NCC specification and differentiation into pharyngeal arch cartilages remains poorly understood. Here we describe the fan/wdr43 mutant phenotype, and present functional characterizations of Wdr43 in craniofacial development. We show that WDR43 is required for the proper nucleolar localization of a variety of nucleolar proteins, including TCOF1/Treacel. These studies provide new insight into ribosomal protein function in early zebrafish development, with focus on NCC derived craniofacial development, as a model for human craniofacial neurocristopathies.
mutants exhibited fewer GFP-positive NCCs, and abnormal NCC migration and pharyngeal arch formation, as compared to age matched wild type siblings (Fig. 1L–L’).

fan locus encodes zebrafish Wdr43/Utp5

Using bulk segregant analyses, we determined that the fan locus mapped between SSLPs z8774 and z9831 on zebrafish Linkage Group (LG) 17 (Fig. 2A), to an interval of 3.93 Mb containing 11 genes. Further analyses of cDNA and genomic DNA sequences of these genes identified a cytosine to thymidine mutation at nucleotide 1066 of the wdr43 gene in this interval, which introduced a premature stop codon at amino acid 356 in exon 9 (Arg356Stop) (Fig. 2A). This mutation was confirmed by sequencing full length wdr43 cDNA amplified from fan mutant mRNA, and sequence analysis of exon 9 of the wdr43 gene in PCR amplified genomic DNA isolated from six individual homozygous fan mutants. This mutation was not present in wild type sibling cDNA or genomic DNA, and was detected along with wild type sequence in heterozygous fan family members. Full length zebrafish Wdr43 contains 650 amino acid and is well conserved from yeast to human (Data not shown). Domain analysis of the zebrafish Wdr43 protein reveals that it is composed of three WD40 repeats and one Utp12 domain (http://pfam.sanger.ac.uk) (Fig. 2A). The truncated form of Wdr43 encoded in the fan mutant lack the C-term 294 aa, including the Utp12 domain (Fig. 2A).

The identified gene mutation in fan/wdr43 mutants was confirmed using two approaches. We used single cell injections of titrated amounts of wdr43 antisense morpholino oligomers (MOs) to test whether targeted depletion of Wdr43 in wild type embryos resulted in embryos that phenocopied the fan mutant. We first confirmed the functional targeting of anti-sense wdr43 MOs by demonstrated quenching of the wdr43-GFP mRNA construct fluorescence in vivo (Fig. S1). We next injected wdr43 MOs into clutches of fan mutants and wild type single cell stage embryos, which were then raised to 3–5 dpf and stained with Alcian blue to examine pharyngeal arch cartilage formation. Our results showed that wild type embryos injected wdr43 MOs exhibited early neural tissue necrosis similar to that observed in fan mutants (Fig. 2C, E, arrows). When wdr43 MOs were injected into single cell stage fan mutant embryos, we observed no apparent exacerbation of the fan mutant phenotype, suggesting that the fan mutation is a functional null (data not shown). WISH analyses of wdr43 MO injected embryos revealed similar reduction in NCC marker gene expression, as observed in fan mutants (Figure S2).

Secondly, we performed rescues by injecting full length wild type wdr43 mRNAs into single cell stage fan mutant and wild type sibling embryos. Analyses of injected embryos at 5 dpf via Alcian blue staining revealed rescue of the pharyngeal arch cartilage formation, although full rescue was not observed (Fig. 2, I versus G, F). Together, these data provide strong evidence that the identified wdr43 gene mutation results in the fan mutant phenotype.

Figure 1. Phenotype of fan mutants. (A–C’) Live images of developmentally staged wild type (A–C) and fan mutant (A’–C’) zebrafish. Arrow in A’ indicates necrotic cells in the presumptive eye region. Bracket in B’ shows necrosis in neural and pharyngeal arch tissues. Arrow in C’ points to the distinct hydrocephaly in fan mutant hind brain ventricles, arrowhead indicates incomplete chorid fissure closure and craniofacial defects. (D–D’) Alcian blue stained pharyngeal arch cartilages in 4dpf wild type (D) and fan mutant (D’). (E–K’) Whole mount ISH images of wild type (E–K) and fan mutants (E’–K’) for neural crest markers at indicated developmental stage. Arrows in (E’–G’) and asterisks in (I’–K’) indicate reduced gene expression in fan mutant embryos. Interestingly, fan mutant embryos exhibit similar crestin expression in the trunk NCCs (asterisks in H, H’) but reduced expression in cranial NCCs. (L–L’) Tg(fli1a:EGFP)/fan mutants (L’) exhibit reduced GFP expression in the pharyngeal arch region as compared to wild type embryos (L). 

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Developmental and tissue specific expression of \textit{wdr43}

We next examined the developmental expression pattern of \textit{wdr43} mRNA via whole mount and sectioned in situ hybridization (ISH). \textit{wdr43} is maternally expressed, and maintains a fairly ubiquitous expression pattern during the first 24 hours of development (Fig. 3A–J). \textit{wdr43} expression becomes restricted to neural and pharyngeal arch tissues between 24 and 48 hpf (Fig. 3K), consistent with the pharyngeal arch defects observed in \textit{fan} mutants. Sectioned ISH demonstrated discrete \textit{wdr43} mRNA expression in neurepithelium and pharyngeal arches at 48 and 72 hpf (Fig. 3, N, N\textsuperscript{9}, P, P\textsuperscript{9}, arrows), consistent with the observed hindbrain and pharyngeal arch defects observed in \textit{fan} mutants. Strong expression was also observed in the gut epithelium (Fig. 3, N, P, arrows). We examined the expression of \textit{wdr43} mutant \textit{wdr43} mRNA using RT-PCR analysis of developmentally staged \textit{fan} mutant and wild type sibling embryos (Fig. 3, M). These analyses showed that wild type \textit{wdr43} was detectable at all stages examined, and also that mutant \textit{wdr43} was detectable in \textit{fan} mutants at 48 and 72 hpf, and thus was apparently not targeted for nonsense mediated decay (Fig. 3, M).

\textit{fantome} mutants exhibit increased apoptosis and reduced cell proliferation

To better characterize tissue necrosis and cell proliferation in \textit{fan} mutants, TUNEL assay and phosphohistone H3 (pH3) IF analyses were performed, respectively, on developmentally staged \textit{fan} mutant and wild type sibling embryos (Fig. 4). TUNEL revealed significantly upregulated apoptosis in \textit{fan} mutants at all developmentally staged examined (Fig. 4A, arrows). In contrast, cell proliferation, indicated via pH3 antibody staining, was decreased in \textit{fan} mutants as compared to age matched wild type siblings (Fig. 4B). Quantification of TUNEL and pH3 immunofluorescence showed significantly increased apoptosis in \textit{fan} mutants as compared to age matched wild type siblings at all stages examined, and significantly decreased cell proliferation at 48 hpf (Fig. 4C). Together, these results are consistent with the observed lack of NCC derived pharyngeal arch tissue formation in \textit{fan} mutants.
Nucleolar localization of wild type and fan mutant Wdr43

To better understand the molecular nature of mutant Wdr43 protein, we next examined the subcellular localization of wild type and fan mutant zebrafish Wdr43 in human cells. First, we performed immunofluorescence analyses of cultured HeLa cells using the anti-human WDR43 antibody and demonstrated that endogenous human WDR43 localized to nucleoli, as shown by co-localization with the nucleolar marker protein, B23 (Fig. 5, A1–A4). Next, we generated N-terminal EGFP-tagged wild type and truncated fan mutant zebrafish Wdr43 constructs driven by the CMV promoter, which we transfected into cultured HeLa cells, and then visualized chimeric fusion protein using anti-GFP antibody. The EGFP-tagged wild type Wdr43 protein showed perfect overlapping expression pattern with B23 (Fig. 5, B1–B4),
were performed using Student’s t-test. (p = for each genotype at each developmental stage. Statistical analyses

consistent with the previously characterized nucleolar localization of the yeast ortholog for Wdr43, Utp5 [39]. In contrast to the full length EGFP-Wdr43 protein, EGFP-tagged fan mutant Wdr43 (amino acids 1–364) lost its exclusive nucleolar localization, and rather exhibited expression throughout the nucleus (Fig. 5, C1–C4). To correlate these in vitro results to in vivo expression studies in zebrafish, the same EGFP-tagged zebrafish wild type and fan mutant wdr43 constructs were injected into single cell stage zebrafish, which were then analyzed via confocal microscopy (Fig. S3). These analyses also showed that while full length Wdr43 co-localized with mCherry-B23 to nucleoli (Fig. S3, A–C), truncated fan mutant Wdr43 remained dispersed throughout the nucleus (Fig. S3, D–F).

Protein-protein interactions between wild type, fan mutant Wdr43 and t-Utp subcomplex proteins

Utp5, the yeast ortholog of Wdr43, has been shown to function in the yeast t-Utp subcomplex, which mediates both pre-ribosomal RNA (rRNA) transcription and processing [16] [17]. Previously, it has been shown that yeast Utp5 interacts with Utp4 and Utp15 (Freed & Baserga 2010) [40]. We used yeast two-hybrid analyses (Y2H) to examine the protein-protein interactions between yeast and zebrafish wild type and fan mutant Wdr43 with other t-Utp complex proteins. We found that both yeast and zebrafish full length Wdr43 interacted with Utp4 and Utp15 (Fig. 5D and Fig. S4), consistent with previously published yeast Utp5/Wdr43 binding studies [17]. We also determined that the C-terminal portion of Wdr43 protein is both necessary and sufficient to mediate this interaction. Zebrafish and yeast truncated fan mutant Wdr43 did not bind to either Utp4 or Utp15, while the C-terminal fragment of Wdr43 alone was able to bind to both Utp4 and Utp15 (Fig. 3D zebrafish and Fig. S4 yeast). Together, these data revealed the conserved interaction of yeast and zebrafish full length Wdr43 proteins with Utp4 and Utp15, and also suggest that the C-terminal portion of Wdr43, which contains the Utp12 domain, is required for protein interaction of Wdr43 with other t-UTP subcomplex member proteins.

Wdr43 is required for optimal pre-rRNA transcription and proper ribosomal protein sub-nucleolar localization

Based on our results, and those of previously published reports, we anticipated that Wdr43 would play an important role in ribosome biogenesis. We therefore investigated pre-rRNA synthesis and processing in 30 hpf and 50 hpf fan mutant and wild type sibling embryos via Northern blot analysis, using a probe specific for the 5′ external transcribed spacer (5′ETS) region of the pre-rRNA at the start site of transcription (Fig. 6A). These analyses showed reduced levels of the primary transcript (labeled a) in fan mutants (M) at both 30 hpf and 50 hpf as compared to that of wild type (W) siblings, consistent with a defect in pre-rRNA transcription in fan mutants. Quantification of the ratio of full length primary transcript (a) to the processed pre-18S rRNA (b) showed reduced pre-rRNA in fan mutants relative to age matched wild type siblings. These results are consistent with previously published results showing that siRNA knockdown of human UTP5 resulted in defects in pre-rRNA transcription and processing [19], and suggest conserved functions for vertebrate Wdr43 and yeast Utp5/ Wdr43 in pre-rRNA transcription [17].

Having defined an important role for zebrafish Wdr43 in ribosome biogenesis, we further examined its function in cultured human HeLa cells, which we transfected with human WDR43 small interfering RNA (siRNA) (Sigma MISSION esiRNA) to silence WDR43 expression. GFP esiRNA was used as a negative control for these studies. Analysis of WDR43 protein and mRNA expression in siRNA treated cells using both Western blot (Fig. 6B) and qRT-PCR (Fig. 6C) analyses, respectively, confirmed that endogenous WDR43 expression was significantly reduced with
WDR43 siRNA treatment. We next examined how WDR43 depletion affected the localization of another t-UTP complex Wdr43 interacting protein, UTP15. Due to the lack of available antibody for UTP15, we transfected N-terminal mCherry tagged zebrafish UTP15 into HeLa cells, and monitored its localization via fluorescent confocal microscopy. Consistent with a role for UTP15 in pre-rRNA processing, we found that mCherry tagged UTP15 localized to nucleoli in control GFP siRNA treated cells (Fig. 6D). In contrast, in HeLa cells depleted of WDR43 using WDR43 siRNA, mCherry-tagged UTP15 did not localize to nucleoli, but rather appeared to exhibit a perinuclear expression pattern (Fig. 6E). These results indicate that WDR43 is required for entry into the nucleus, as well as for proper nucleolar localization of UTP15.

It was intriguing to us that many of the observed phenotypes observed in fan mutant zebrafish have also been reported in humans (and mice) with mutations in TCOF1/Treacle, the gene commonly mutated in Treacher-Collins Syndrome (TCS). TCS results in aberrant NCC specification and differentiation, craniofacial dysmorphologies, increased cell apoptosis, and upregulated p53 signaling [10,11]. Based on these common characteristics, we investigated whether the localization of TCOF1 and other nucleolar proteins was affected in human HeLa cells depleted of WDR43 protein. For reference, we examined the expression of the nucleolar proteins Mpp10, Nucleolin and Fibrillarin, which have been associated with distinct nucleolar functions. Mpp10 is normally found in the dense fibrillar component (DFC) and in the boundary between the DFC and the fibrillar center (FC), sites of rDNA transcription and pre-rRNA splicing and modification by snRNPs, while Fibrillarin is normally found in association with snoRNAs throughout the DFC [41]. Nucleolin/C23 is normally localized to the outer layer of nucleoli with fainter expression at the center [42]. Our investigation of the expression of these nucleolar proteins, and TCOF1, in control and WDR43 siRNA treated HeLa cells (Figure 7) showed that TCOF1 exhibited a reduced and perinucleolar expression pattern in WDR43 depleted cells (Fig. 7 B vs. A, arrows). Mpp10 expression appeared reduced, but was expressed throughout the smaller nucleoli (Fig. 7 B vs. D, arrows). Nucleolin, although barely detectable in WDR43 siRNA expressing cell lines as compared to control GFP siRNA treated cells, was also expressed in a perinucleolar fashion, similar to that of TCOF1 (Fig. 7 F vs. E, arrows). In contrast, Fibrillarin expression appeared relatively less affected in WDR43 depleted cells, and was detected throughout the smaller nucleoli (Fig. 7H vs. G, arrows). These results are indicative of disrupted nucleolar organization and rRNA transcription, consistent with the observed
defects in pre-rRNA transcription observed in fan mutants. To confirm and more reliably study these observations, we next examined TCOF1 localization in stable HeLa cell lines expressing stable short hairpin RNAs (shRNAs) for GFP control and WDR43 (Fig. 8). We first tested five shRNAs against WDR43 gene and found that all of them showed somewhat reduced WDR43 expression using qRT-PCR analysis (Fig. S5). Western blot analyses showed that two of the shRNA WDR43 cell lines, sh9 and 2a1, exhibited the most efficient inhibition of protein levels. As observed in WDR43 siRNA treated cell lines, stable WDR43 shRNA expressing cell lines sh9 and 2a1 exhibited mislocalized TCOF1 expression at the periphery of nucleoli, as compared to control shRNA stable cell lines (Fig. 8, E, I, N, O vs. A, M, arrows). Together, these results suggest that WDR43 expression is required for proper nucleolar organization and the subnucleolar localization of a variety of nucleolar proteins including TCOF1, and for optimal pre-rRNA transcription.

WDR43 is required for proper nucleolar morphology, maturation and assembly

We also found that WDR43 depletion had an effect on nucleolar size and shape. WDR43 depleted cells had larger numbers of mini nucleoli as compared to control cells, which exhibited fewer numbers of larger sized, mature nucleoli. To quantitate this observation, we monitored nucleolar number and size in control and WDR43 depleted cultured HeLa cells immunostained for TCOF1 (Fig. 8). We found that in normal and control shRNA cultured HeLa cells, nucleoli reassembled after mitosis, with several small nucleoli fusing into 1–4 larger, mature nucleoli per HeLa cell. In contrast, TCOF1 expressing nucleoli failed to fuse together in WDR43 shRNA expressing HeLa cells, but rather remained as small unfused mini nucleoli, or “nucleolar caps”, which also appeared spherical shape as compared to the irregular shaped nucleoli present in control HeLa cell cultures (Fig. 8 F, J vs. B). We also found that the
average number of nucleoli was increased, and the number of fused nucleoli was reduced, in WDR43 shRNA expressing HeLa cells as compared to control shRNA treated HeLa cells, as shown using the nucleolar marker B23 (Fig. 8, F, J vs. B). Quantification of these results revealed that WDR43 depleted cells exhibited increased numbers of smaller nuclei as compared to control cells (Fig. 8, Q, R). Together, these results demonstrate that depletion of WDR43, an essential ribosome biogenesis factor, affects nucleolar maturation and assembly.

fan mutants exhibit upregulated p53 signaling

Ribosome biogenesis defects, such as those observed in fan mutants, have been reported to be associated with upregulation of the p53 signaling pathway, and cellular apoptosis [11,25] [43] [44] [45] [46]. Based on the increased apoptosis observed in fan mutants, we next examined p53 signaling pathway gene expression in developing fan mutant and wild type sibling embryos. Immunohistochemical analysis using the zebrafish p53 antibody (kind gift of David Lane) [47] revealed high levels of expression of p53 in fan mutants, while p53 was virtually undetectable in wild type sibling embryos at 5 dpf (Fig. 9, B vs. A). Next, we used qRT-PCR analyses to show that the expression of p53 downstream target genes, including the N-terminal truncated p53 isoform delta113p53, mdm2, and cyclin G1 were all upregulated in 24, 48 and 72 hpf fan mutants as compared to wild type sibling controls (Fig. 9, C, D and data not shown). We next tested whether knockdown of p53 signaling via injection of p53 anti-sense MOs could rescue the fan mutant phenotype. Similar to previous reports in the Treacher-Collins mouse model [11], we found that the fan mutant pharyngeal, neural and eye defects were largely rescued in p53 MO injected fan mutants (Fig. 9, G vs. F, and H), and NCC marker gene expression was also rescued in fan mutants (Fig. S2). TUNEL analyses revealed rescue of apoptosis in p53MO injected fan mutants (Fig. 9, K vs. J, I), indicating that increased apoptosis observed in fan mutants was mediated via upregulated p53 signaling pathways.
proliferation, migration and differentiation. In Treacher-Collins NCCs may be more sensitive to ribosome biogenesis defects as and Wdr43 in polymerases I and III in Treacher-Collins Syndrome [54] [55], proteins – TCOF1/Treacle and a common subunit of RNA and differentiation are similarly affected by mutations in nucleolar with Treacher-Collins Syndrome. The fact that NCC specification resembles the craniofacial malformations observed in individuals diverse readouts in different cell types. mRNA competition for timely translation, which could result in spectrum of translation products in different cell types due to increased translation in NCC [21]. Finally, decreased efficiency of most significantly affect those tissues exhibiting a high demand for subsequent anticipated reduced protein translation efficiency will patterns. Another theory is that ribosome biogenesis defects and apoptosis may similarly exhibit tissue specific expression possible that additional ribosomal proteins that regulate cell cycle or apoptosis may similarly exhibit tissue specific expression patterns. One possibility is that there are tissue specific, developmental requirements for ribosome biogenesis proteins. For example, certain ribosome biogenesis factors may have tissue specific, developmental expression patterns. In fact, we show here that the expression of zebrafish vdr43 mRNA becomes localized to neural and pharyngeal arch tissues starting at ~24 hpf, which is consistent with the observed fan/vdr43 mutant phenotype. It is possible that additional ribosomal proteins that regulate cell cycle or apoptosis may similarly exhibit tissue specific expression patterns. Another theory is that ribosome biogenesis defects and subsequent anticipated reduced protein translation efficiency will most significantly affect those tissues exhibiting a high demand for protein synthesis. This may include NCC and erythropoiesis progenitor cells, although recent data does not find evidence for increased translation in NCC [21]. Finally, decreased efficiency of cellular translation machinery may affect a wide and varied spectrum of translation products in different cell types due to mRNA competition for timely translation, which could result in diverse readouts in different cell types.

The craniofacial phenotype exhibited by zebrafish fan mutants resembles the craniofacial malformations observed in individuals with Treacher-Collins Syndrome. The fact that NCC specification and differentiation are similarly affected by mutations in nucleolar proteins – TCOF1/Treacle and a common subunit of rRNA polymerases I and III in Treacher-Collins Syndrome [54] [55], and Wdr43 in fan mutants - leads to the intriguing question of why NCCs may be more sensitive to ribosome biogenesis defects as compared with other tissues. Based on our data presented here and on the published reports of others, we hypothesize that high protein translation levels must be maintained by progenitor and differentiating NCCs in order to support their extensive cell proliferation, migration and differentiation. In Treacher-Collins Syndrome, TCOF1 mutant induced defects in ribosome biogenesis are characterized by stimulation of the nucleolar stress response, which in turn activates the p53 apoptosis pathway, resulting in the depletion of the neural crest precursor pool [10]. We observe a similar upregulation of p53 signaling and depletion of NCCs in fan mutants. Although beyond the scope of the present study, it will be interesting in future studies to compare pre-rRNA transcription, ribosome biogenesis and protein translation efficiency in developmentally staged NCC versus non-NCC populations harvested from fan mutant and wild type siblings.

Our Northern blot results indicated that pre-rRNA levels are significantly decreased in developmentally staged zebrafish fan/wdr43/utp5 mutants, consistent with the previously characterized role for yeast Up5 in pre-rRNA transcription [16,19]. We suggest that a variety of nucleosomal proteins are required for optimal pre-rRNA transcription. Novel findings from this report include the fact that blocking WDR43 function in HeLa cells resulted in the distinct mislocalization of nucleolar proteins including UTP15, Mpp10, nucleolin and to a lesser extent fibrillarin, suggesting that Wrd43/UTP5 is required for proper subnucleolar organization and function. Interestingly, TCOF1 also mislocalized to the outer periphery of nucleoli, rather than exhibiting its normal distribution throughout the nucleolus, suggesting that WDR43 may also be required for proper TCOF1 subnucleolar localization and function. Although we have not detected direct binding between TCOF1 and WDR43/UTP5 using Y2H, we have detected interactions between WDR43/UTP5 and other rRNA transcription component proteins (data not shown). Together, these results suggest roles for Wdr43/UTP5 in ribosomal protein sub-nucleolar localization and function of other ribosome biogenesis factors, and raise the intriguing possibility that manipulation of WDR43 expression could be used to correct the localization and improve the function of TCOF1 in Treacher-Collins Syndrome patients.

Nucleolar mis-localization phenotypes have also been observed in HeLa cells treated with actinomycin D, an inhibitor of RNA Pol I [56], which is a TCOF1/Treacle interacting protein [57]. It is possible that WDR43 may also function together with TCOF1 and Nopp140 to recruit proteins to the nucleolar organizer regions (NORs) and the upstream binding factor (UBF), an RNA Pol I transactivator [19]. Wdr43 could also mediate rRNA transcription by binding to rDNA and UBF directly, as shown by other Utps [58]. These functions for Wdr43 remain to be elucidated.

We also used both siRNA and shRNA WDR43 silencing methods to confirm the function of WDR43 in nucleolar fusion in cultured HeLa cells. At the present time, mechanisms regulating nucleolar fusion remain poorly understood. It has been shown that after mitosis, multiple small nucleoli form around transcriptionally active NORs, and as cells progress through the cycle, these small nucleoli fuse to form larger nucleoli [59] [60]. Although the mechanism of WDR43 function in nucleolar fusion is not clear, preventing nucleolar fusion may not be common to all ribosome biogenesis protein mutations based on the fact that inhibition of NOL11 resulted in the formation of one large (not small) nucleolus [20]. One possible explanation is that WDR43 depletion may result in structural changes to rDNA, which in turn could interfere with nucleolar fusion [61]. Such a proposed function for WDR43

Discussion

Proper ribosome biogenesis is required for the production of functional ribosomes, the primary site of protein synthesis. Most if not all ribosomal proteins (RPs) are thought to be essential for ribosome biogenesis and cell survival. It is therefore surprising that ribosome biogenesis defects caused by mutations in certain RPs can lead to variable and seemingly tissue-specific defects in vertebrate development. For example, mutations in several RPs are associated with human congenital hypoplastic Diamond-Blackfan anemia (DBA), and similar DBA phenotypes were observed when DBA associated RP mutations were expressed in zebrafish [48–51]. In addition, several reports on zebrafish ribosome biogenesis protein mutants describe a variety of diverse phenotypes, ranging from tumors, to central nervous system degeneration, to organogenesis defects [23,25,52,53].

Here we present data showing that defects in the zebrafish ribosome biogenesis protein Wdr43 result in early developmental defects in a variety of tissues, including neural, heart and pharyngeal arches, while later developmental defects appear fairly localized to NCC derived pharyngeal arch cartilages. These observations raise the question of how a defect in what is thought to be a universally required ribosomal biogenesis protein, Wdr43, can result in a rather specific craniofacial tissue-specific phenotype? One possibility is that there are tissue specific, developmental requirements for ribosome biogenesis proteins. For example, certain ribosome biogenesis factors may have tissue specific, developmental expression patterns. In fact, we show here that the expression of zebrafish vdr43 mRNA becomes localized to neural and pharyngeal arch tissues starting at ~24 hpf, which is consistent with the observed fan/vdr43 mutant phenotype. It is possible that additional ribosomal proteins that regulate cell cycle or apoptosis may similarly exhibit tissue specific expression patterns. Another theory is that ribosome biogenesis defects and subsequent anticipated reduced protein translation efficiency will most significantly affect those tissues exhibiting a high demand for protein synthesis. This may include NCC and erythropoiesis progenitor cells, although recent data does not find evidence for increased translation in NCC [21]. Finally, decreased efficiency of cellular translation machinery may affect a wide and varied spectrum of translation products in different cell types due to mRNA competition for timely translation, which could result in diverse readouts in different cell types.

The craniofacial phenotype exhibited by zebrafish fan mutants resembles the craniofacial malformations observed in individuals with Treacher-Collins Syndrome. The fact that NCC specification and differentiation are similarly affected by mutations in nucleolar proteins – TCOF1/Treacle and a common subunit of rRNA polymerases I and III in Treacher-Collins Syndrome [54] [55], and Wdr43 in fan mutants - leads to the intriguing question of why NCCs may be more sensitive to ribosome biogenesis defects as compared with other tissues. Based on our data presented here and on the published reports of others, we hypothesize that high protein translation levels must be maintained by progenitor and differentiating NCCs in order to support their extensive cell proliferation, migration and differentiation. In Treacher-Collins Syndrome, TCOF1 mutant induced defects in ribosome biogenesis are characterized by stimulation of the nucleolar stress response, which in turn activates the p53 apoptosis pathway, resulting in the depletion of the neural crest precursor pool [10]. We observe a similar upregulation of p53 signaling and depletion of NCCs in fan mutants. Although beyond the scope of the present study, it will be interesting in future studies to compare pre-rRNA transcription, ribosome biogenesis and protein translation efficiency in developmentally staged NCC versus non-NCC populations harvested from fan mutant and wild type siblings.

Our Northern blot results indicated that pre-rRNA levels are significantly decreased in developmentally staged zebrafish fan/wdr43/utp5 mutants, consistent with the previously characterized role for yeast Up5 in pre-rRNA transcription [16,19]. We suggest that a variety of nucleosomal proteins are required for optimal pre-rRNA transcription. Novel findings from this report include the fact that blocking WDR43 function in HeLa cells resulted in the distinct mislocalization of nucleolar proteins including UTP15, Mpp10, nucleolin and to a lesser extent fibrillarin, suggesting that Wrd43/UTP5 is required for proper subnucleolar localization and function. Although we have not detected direct binding between TCOF1 and WDR43/UTP5 using Y2H, we have detected interactions between WDR43/UTP5 and other rRNA transcription component proteins (data not shown). Together, these results suggest roles for Wdr43/UTP5 in ribosomal protein sub-nucleolar localization and function of other ribosome biogenesis factors, and raise the intriguing possibility that manipulation of WDR43 expression could be used to correct the localization and improve the function of TCOF1 in Treacher-Collins Syndrome patients.

Nucleolar mis-localization phenotypes have also been observed in HeLa cells treated with actinomycin D, an inhibitor of RNA Pol I [56], which is a TCOF1/Treacle interacting protein [57]. It is possible that WDR43 may also function together with TCOF1 and Nopp140 to recruit proteins to the nucleolar organizer regions (NORs) and the upstream binding factor (UBF), an RNA Pol I transactivator [19]. Wdr43 could also mediate rRNA transcription by binding to rDNA and UBF directly, as shown by other Utps [58]. These functions for Wdr43 remain to be elucidated.

We also used both siRNA and shRNA WDR43 silencing methods to confirm the function of WDR43 in nucleolar fusion in cultured HeLa cells. At the present time, mechanisms regulating nucleolar fusion remain poorly understood. It has been shown that after mitosis, multiple small nucleoli form around transcriptionally active NORs, and as cells progress through the cycle, these small nucleoli fuse to form larger nucleoli [59] [60]. Although the mechanism of WDR43 function in nucleolar fusion is not clear, preventing nucleolar fusion may not be common to all ribosome biogenesis protein mutations based on the fact that inhibition of NOL11 resulted in the formation of one large (not small) nucleolus [20]. One possible explanation is that WDR43 depletion may result in structural changes to rDNA, which in turn could interfere with nucleolar fusion [61]. Such a proposed function for WDR43
may be dependent or independent of its function in the t-Utp complex. A recent study using Xenopus oocytes showed that nucleoli exhibit fluid dynamics similar to that of liquid droplets, and that nucleolar fusion requires dynamic exchange between nucleoli and the nucleoplasm [62]. In future studies, it will be interesting to determine whether WDR43 is also involved in this process.

Recent reports emphasize the apparent tissue specific functions for ribosomal proteins previously thought to exhibit functions in all cells and tissues. The results presented here suggest previously unrecognized roles for Wdr43/UTP5 in craniofacial development. The fact that Wdr43/UTP5 is needed for proper formation of nucleoli and for sub-nucleolar organization and function indicates important roles for Wdr43 as a key participant in ribosome biogenesis. As such, the zebrafish mutant fan may provide a valuable vertebrate developmental model and tool to continue in depth functional studies of RPs and ribosome biogenesis factor proteins in NCC differentiation, including the identification of effective tools for reducing the incidence of craniofacial birth defects.

Materials and Methods

Zebrafish husbandry

AB and WIK fan/mutant and wild type zebrafish were raised in the Tufts Zebrafish Facility at 28.5°C and developmentally staged as previously described (Westerfield, M., 1995). For whole mount in situ hybridization analyses, pigmentation was inhibited by treating embryos with 1-phenyl-2-thiourea (PTU) at a final concentration of 0.2 mM as previously described (The Zebrafish Book, U. Oregon Press).

Ethics statement

All experimental procedures on zebrafish embryos and larvae were approved by the Tufts University Institutional Animal Care and Use Committee (IACUC) and Ethics Committee.

Positional cloning of the fantome mutant locus

The fan mutant was identified in a large-scale ENU-mutagenesis screen conducted by the Yelick Laboratory [29]. Genetic mapping strains were created by crossing identified homozygous fan mutants to polymorphic WIK wild type zebrafish. Embryos were collected from pairwise matings of mapping strain fan/WIK heterozygotes, and scored at 48 hpf for fan specific phenotypes. Genomic DNA was extracted from individual fan mutant and wild type embryos, and bulk segregant analyses were performed using primers designed to amplify SSLP markers from the Massachusetts General Hospital Zebrafish Server website (http://zebrafish.mgh.harvard.edu). The fan mutation mapped to zebrafish linkage group 17 (LG17), to a region spanning 11 genes. Nucleotide sequence analyses of all 11 genes identified a premature stop codon in the region spanning 11 genes. Nucleotide sequence analyses of all 11 genes identified a premature stop codon in the region spanning 11 genes. Nucleotide sequence analyses of all 11 genes identified a premature stop codon in the region spanning 11 genes.

In situ hybridization

Whole-mount and sectioned in situ hybridizations were performed as previously described [Thissie et al., 1993], using a probe generated via PCR using the following primers \(\text{wdr43-forward: 5'}-\text{CAGTTGACAAAAGTTGGTGA-3'}\); \(\text{wdr43-reverse: 5'}-\text{AAAGTTCTGGTGGCCTGCA-3'}\). All other probes were obtained from zfin.org. Embryos were analysed using Zeiss Axioshot and M2Bio microscopes, and imaged using Zeiss Axioshot Imager digital camera (Munich, Germany). Digital images were processed using Adobe Photoshop software.

Targeted protein depletion of Wdr43/UTP5 using antisense \text{wdr43} morpholino oligomer (MO) injections

Antisense morpholino oligonucleotides (MO)s targeted to the initiation of translation codon of \text{wdr43} mRNA (\(5\text{’TCCGTCGCC-CCGCATCTTACCCGTC-3’}\)) were injected into the yolk of 1 cell stage wild type or fan mutant embryos. 2 nL of MO at a concentration of 10 ng/µL was used to knockdown \text{wdr43} translation.

Quantitative reverse transcriptase polymerase chain reaction (qRT-PCR)

Total RNA was extracted from 20 wild type and 20 fan mutant embryos at 24, 48 and 72 hpf, or from HeLa cells 48 hours after transfection using RNEasy Plus Kit (Qiagen, Valencia, CA). DNA was removed using the DNA-free DNase Treatment & Removal Kit (Ambion,Life Technologies, Grand Island, NY) to remove genomic DNA contamination. cDNA was synthesized using a SuperScript III First-Strand Synthesis System (Invitrogen, Life Technologies, Grand Island, NY) with random primers. Gene expression was quantified by qRT-PCR using Quantitect SYBR Green PCR Master Mix (Qiagen, Valencia, CA) and real-time cycler Mx3000P (Stratagene, Agilent Technologies, Santa Clara, CA). Primers for zebrafish \(p33\) isofoms, \(mdm2\), and \(cyclinG1\) were used as described [63]. The following primers were used to amplify the human WDR43 gene: Forward: CCGTCCGCCGAACCT-CAGTGGTGAC; Reverse: AACTGGGGTGCTACGTCGGTGA. Primers for \(\beta\)-actin, used to normalize the expression levels, were as described [64].

Yeast two-hybrid analysis

Yeast two-hybrid assays for interaction between yeast and zebrafish Utp proteins were performed as previously described (Freed and Baserga, 2010). Briefly, yeast \(utp5\) cDNA encoding full length, N-terminal (1–343aa) and C-terminal (344–643aa) proteins were cloned into the pGADT7 prey vector. Additional yeast UTP genes of the t-Utp subcomplex (Utp8, Utp9, Utp10, Utp15 and Utp17) cloned into the bait vector were as previously described (Freed and Baserga 2010). Both bait and prey vectors were transformed into AH109 yeast strain and interactions were identified based on the ability of transfectants to grow on AHTL dropout medium after 3–5 days of incubation at 30°C. To test the interaction between zebrafish Wdr43/Utp3, Utp4 and Utp15, full length zebrafish \(utp4\) and \(utp15\) cDNAs were purchased (Openbiosystems, Lafayette, CO) and cloned into pGADT7 prey vector. Zebrafish \text{wdr43} cDNAs encoding full length, truncated fan mutant Wdr43 (1–356aa) and C-terminal portion of Wdr43 (357–600aa) proteins were cloned into the pGABT7 bait vector. Interactions were tested by growth in triple dropout medium after 3–5 days of incubation at 30°C. The Y2H studies of zebrafish Wdr43 protein interactions were tested in both bait and prey vectors.

Wdr43 subcellular localization

To determine the subcellular localization of wild type and mutant Wdr43 proteins, \(GFP\) cDNA was cloned onto the N-terminal end of full length or \(fan\) mutant zebrafish \text{wdr43} cDNA under the direction of the CMV promoter, using multi-site Gateway reactions [65]. These constructs were then transfected into HeLa cells using Lipofectamine 2000 reagent (Invitrogen, Life Technologies, Grand Island, NY). After 36 hours of growth at 37°C, transfected cells were fixed with 4% PFA, and subjected to standard immunofluorescence (IF) analyses using the anti-B23 antibody (1:200, Santa Cruz Biotechnology, Inc., Santa Cruz, CA). The rabbit polyclonal anti-WDR43 antibody (1:100, Abcam, Cambridge, MA) was used to detect the endogenous WDR43. To
check the expression in zebrafish embryos, the same constructs were injected into single cell zebrafish embryos, which were harvested at 24 hpf, and analyzed for GFP expression using a Leica TCS SP2 confocal microscope.

RNA interference
For siRNA experiments, HeLa cells were transfected with WDR43 MISSION esiRNA (Sigma, EHUC004691) using Lipofectamine 2000 reagent (Invitrogen, Life Technologies, Grand Island, NY). GFP esiRNA (Sigma, EHUEGFP) was used as a negative control. Treated cells were harvested 48 hours after transfection for Western blotting and immunofluorescence experiments. shRNA constructs were purchased from Openbiosystems and used to establish stable shRNA expressing HeLa cells following the manufacturer’s protocol. The following shRNA constructs were used: pGIPZ-WPI1-2, RHS4430-98535022; pGIPZ-non-targeting control RMS4348.

Northern blot analysis
Total RNA was extracted from 20 wild type and 20 fan mutant embryos at 30 and 50 hpf using standard Trizol protocol for RNA isolation. Northern blot analysis was carried out as described in Freed et al, 2012 [20]. For each sample, 2 μg of RNA was separated by electrophoresis on a 1% agarose/1.25% formaldehyde gel. Pre-rRNA species were detected by methylene blue staining and hybridization with a 32P-radio labelled oligonucleotide probe to the 5′ETS: CGGCCAGAGTGGAGGAGAAGCTTTCCTCCTGCTG-TCA. Quantification of pre-rRNA processing was performed using Image J (National Institutes of Health).

Apoptosis and cell proliferation analyses
Developmentally staged wild type and fan mutant embryos, fixed and processed for cryosectioning, were sectioned at 10 μm. Apoptosis TUNEL assay was performed using the In situ cell death detection kit. Fluorescein (Roche Applied Science, Indianapolis, IN, USA). Cell proliferation was assayed with phospho-Histone H3 immunofluorescence analysis, using anti-phospho Histone H3 (Ser10) antibody (Cell Signaling, Danvers, MA) and anti-rabbit goat antibody conjugated with Alexafluor 594 (Life Technologies, Grand Island, NY).

Supporting Information
Figure S1 Knockdown efficiency of wdr43 Morpholino. (A) Diagram of the wdr43 reporter construct used to test wdr43 MO knockdown efficiency. GFP (green box) was fused in frame to the 3′ end of a portion of the 5′ end of the wdr43 cDNA including the 5′ UTR (purple line) and first two exons of wdr43 gene (purple box). Red line indicates the MO target region. The chimeric gene was driven by the CMV promoter and followed by the 3′ SV40 polyA signal (PA). (B–B′) Fluorescence of shield stage zebrafish embryos injected with the wdr43-GFP reporter construct and control MO (CMO) (B, fluorescent microscopy; B′ bright field, B″ merged fluorescent and bright field). (C–C′) Lack of fluorescence in shield stage zebrafish embryos injected with wdr43-GFP reporter constructs and wdr43 MO (C, fluorescent microscopy; C′ bright field; C″ merged fluorescent and bright field). (TIF)

Figure S2 WISH analyses of wdr43 MO injected embryos, and p53 MO injected fan mutants. WISH was performed to examine NCC markers crestin and dlx2a expression in 24 hpf wild type embryos (WT), wild type embryos injected with wdr43 MO, fan mutants, and fan mutants injected with −53 MO, as indicated. wdr43 MO injected embryos exhibited down regulated expression of all NCC markers, similar to that observed in fan mutants (arrows). Injection of p53 MO into single cell stage fan mutants rescued NCC marker gene expression (arrows). (TIF)

Figure S3 Subcellular localization of EGFP-tagged wild type or fan mutant Wdr43 in zebrafish embryos. Confocal images taken from 24 hpf old zebrafish embryos injected at single cell stage with EGFP tagged wild type (A1–4, B1–4, C1–4) or fan mutant (D1–3, E1–3, F1–3) wdr43 mRNA. Co-injection of mCherry tagged zebrafish B23 mRNA was used to label nuclei (A2, B2, C2, D2, E2, F2). Each row of panels indicates different types of cells imaged in the whole embryo. GFP expression was manually saturated in panels (A4, B4, C4) to reveal the entire nucleus. (TIF)

Figure S4 Yeast two hybrid analysis of yeast Wdr43/UTP5 and yeast t-UTP subcomplex proteins. Primers for full length, and fan mutant truncated yeast UTP5 were used to amplify and subclone these yeast cDNA constructs into pGADT7 vector. The remaining yeast expression constructs were obtained from Dr. S. Baerga. P: permissive medium (-Leu and -Trp). S: selective medium (-Ade, -His, -Leu and -Trp). (TIF)

Figure S5 qRT-PCR analysis of WDR43 mRNA expression in WDR43 shRNA treated HeLa cells. The relative expression levels of WDR43 were normalized to human β-actin gene. Bar graph shows data from three independent experiments. (TIF)

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Author Contributions
Conceived and designed the experiments: PCY CZ VA YG SB YZ. Performed the experiments: CZ VA YG ML VL SP YZ KLM. Analyzed the data: PCY VA YG SB YZ KLM SL. Contributed reagents/materials/analysis tools: PCY SB LZ. Wrote the paper: CZ VA SB PCY. Performed the majority of the functional characterization experiments described in this manuscript, including Y2H, subcellular localizations, shRNA, shRNA, Northern blot and Western blot analyses: CZ. Supervised Melissa LaBonty in her experiments: CZ. Significantly contributed to writing the manuscript and generating figures: CZ. Conducted the qRT-PCR analyses, WISH and sectioned ISH, TUNEL, IHC/IF, and participated in writing and editing the manuscript: VA. Performed the positional cloning to identify the wdr43 mutation in fan mutant zebrafish: YG. Performed Northern and Western blot analyses: CZ ML KLM. Performed RT-PCR of developmentally staged fan mutant and wild type siblings, MO phenocopy of the fan mutants, and wild type mRNA rescue experiments: VA VL SP. Helped analyze the Northern blot studies, and significantly contributed to editing and improving the final manuscript: SB. Supported the positional cloning laboratory, and provided comments on the manuscript: LZ. Conducted the screen that resulted in the identification of the fan mutant, oversaw all of the experiments performed in this study, participated in writing and editing the manuscript and figures, and supported the laboratory and zebrafish facility: PCY.
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