CARMIL3 is important for cell migration and morphogenesis during early development in zebrafish

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ABSTRACT

Cell migration is important during early animal embryogenesis. Cell migration and cell shape are controlled by actin assembly and dynamics, which depend on capping proteins, including the barbed-end heterodimeric actin capping protein (CP). CP activity can be regulated by capping-protein-interacting (CPI) motif proteins, including CARMIL (capping protein Arp2/3 myosin-I linker) family proteins. Previous studies of CARMIL3, one of the three highly conserved CARMIL genes in vertebrates, have largely been limited to cells in culture. Towards understanding CARMIL function during embryogenesis in vivo, we analyzed zebrafish lines carrying mutations of carmil3. Maternal-zygotic mutants showed impaired endodermal migration during gastrulation, along with defects in dorsal forerunner cell (DFC) cluster formation, which affected the morphogenesis of Kupffer's vesicle (KV). Mutant KVs were smaller, contained fewer cells and displayed decreased numbers of cilia, leading to defects in left/right (L/R) patterning with variable penetrance and expressivity. The penetrance and expressivity of the KV phenotype in carmil3 mutants correlated well with the L/R heart positioning defect at the end of embryogenesis. This in vivo animal study of CARMIL3 reveals its new role during morphogenesis of the vertebrate embryo. This role involves migration of endodermal cells andDFCs, along with subsequent morphogenesis of the KV and L/R asymmetry.

1. Introduction

1.1. CARMIL regulation of actin assembly via capping protein

CARMILs are one family of capping-protein-interacting (CPI-motif) proteins, reviewed in (Edwards et al., 2014). Vertebrates, including zebrafish, have three conserved CARMIL-encoding genes, called CARMIL1, CARMIL2 and CARMIL3 in humans (Stark et al., 2017; Stark and Cooper, 2015). The zebrafish genes, carmil1, carmil2 and carmil3, have distinct spatial and temporal expression patterns during development (Stark and Cooper, 2015). In human cultured cells, available evidence indicates that the gene products have distinct subcellular locations and functions, even within one cell type (Lanier et al., 2015; Liang et al., 2009; Stark et al., 2017). The functions of CARMIL1 and CARMIL2 include cell migration, macropinocytosis, lamellipodial activity and cell polarity (Edwards et al., 2014; Lanier et al., 2015; Liang et al., 2009; Stark et al., 2017).

In contrast to this information for CARMIL1 and CARMIL2, relatively less is known about the CARMIL3 gene. In mouse, the CARMIL3 protein localizes to developing synapses and spines in neurons, where it recruits actin CP, which is known to cap barbed ends of actin filaments. Depletion of CARMIL3 protein in neurons leads to defects in spine and synapse assembly and function (Spence et al., 2019). In breast and prostate cancer patients, elevated expression of CARMIL3 correlates with poor outcomes, and mouse tumor models reveal a role for CARMIL3 in epithelial-mesenchymal transition, cadherin-based cell adhesions, and cell migration and invasion (Wang et al., 2020).
1.2. Cell migration and morphogenesis in early vertebrate development

Early vertebrate embryogenesis sees massive cell rearrangements that establish and shape the three germ layers—mesoderm, endoderm, and ectoderm—during gastrulation. The most deeply positioned endoderm gives rise to the gut and other alimentary organs. At the onset of zebrafish gastrulation, mesendodermal progenitors are located at the margin of a cup-shaped blastoderm that covers the animal hemisphere of a large yolk cell (Warga and Kimmel, 1990). The mesodermal and endodermal lineages soon separate, and internalized endodermal cells initially disperse on the yolk surface as individuals via a random walk to almost completely cover the yolk cell (Pézéron et al., 2008). Concurrently, mesoderm and ectoderm spread around the yolk cell in the process of epiboly (Warga and Kimmel, 1990). Later during gastrulation, endoderm cells migrate towards the dorsal midline along trajectories that are biased either animal/anteriorly (for cells in animal hemisphere) or vegetally/posteriorly (for cells in vegetal hemisphere), thus simultaneously elongating the nascent endoderm along the AP axis (Schmid et al., 2013). Endodermal cell migration during gastrulation depends on Rac1-regulated actin dynamics (Woo et al., 2012).

On the dorsal gastrula side there is a small cluster of dorsal forerunner cells (DFCs) that travel vegetalward in advance of the spreading germ layers. Later during segmentation DFCs will form an epithelial ciliated vesicle known as Kupffer's vesicle (KV), the left-right (L/R) organizer of...
A Wild type
MZ carmil3sa19830

B Schematic of measurements

C, C' carmil3sa19830+
Endo cells
DFCs
100μm

D Migration path

E, E' MZ carmil3sa19830
Endo cells
DFCs

sox17:EGFP

G

<table>
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<th>Embryos</th>
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<td>MZ carmil3sa19830</td>
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[Graphs showing Speed (μm/m/m) and Migration persistence]

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zebrafish (Amack and Yost, 2004). At the end of epiboly, DFCs form multiple rosette-like epithelial structures whose focal points are enriched for apical proteins. During segmentation, these rosettes arrange into a single rosette lined by a lumen with cilia at the apical membrane of the cells, thereby forming the KV (Oteiza et al., 2008). L/R patterning in zebrafish depends on the motile cilia in the KV that generate an asymmetric fluid flow (Gokey et al., 2015, 2016; Sampaio et al., 2014). Mutations that affect the shape and size of the KV or that affect the number or length of cilia in the KV can impair robust L/R patterning (Amack, 2014).

To investigate the function of CARMIL3 in vertebrate development, we examined phenotypes resulting from the disruption of the gene encoding CARMIL3 in zebrafish. We found defects of variable penetrance and expressivity, in endodermal cell migration, DFC migration and clustering, KV morphogenesis, the number of cells and cilia in the KV, and L/R asymmetry.

2. Materials and methods

2.1. Zebrafish lines and husbandry

Animal protocols were approved by the Institutional Animal Care and Use Committees at University of Iowa and Washington University in St. Louis. At University of Iowa, zebrafish were maintained as described previously (Xu et al., 2011) and embryos were obtained by natural spawning and staged according to morphological criteria or hours post fertilization (hpf) at 28°C, 32°C unless otherwise specified, as described previously (Kimmel et al., 1995). At Washington University, zebrafish and embryos were maintained at 28.5°C using the standard operating procedures and guidelines established by the Washington University Zebrafish Facility, described in detail at http://zebrafishfacility.wustl.edu/documents.html. The following zebrafish lines were used in this study: AB*/Tuebingen, Tg(sox17:EGFP) (Mizoguchi et al., 2008), carmil3sa19830 (irc166sa19830), and carmil3t413.

Zebrafish line carmil3sa19830 (irc166sa19830) was obtained from the Zebrafish International Resource Center (Eugene, OR) (described at https://zfin.org/ZDB-ALT-131217-14950) (Kettleborough et al., 2013). The mutation was created by N-ethyl-N-nitrosourea (ENU) treatment of adult males, and the mutated gene has a G to T conversion at an essential splice site of intron 27–28, which introduces multiple stop codons beginning at amino acid residue 832 (Kettleborough et al., 2013). Failure to splice at this site is predicted to change amino acid residue 831 from E to D, with the next codon being ochre TAA, and thus truncating the protein to 832 residues from its normal length of 1384 residues (Fig. 1A). This residue, 832, is predicted to lie within the third alpha-helical segment of the homodimerization domain (https://www.uniprot.org/uniprot/A0A0PCP424protvista).

Zebrafish line carmil3t413 was created by TALEN-mediated mutagenesis (Boch et al., 2009; Moscou and Bogdanove, 2009). TALEN target sequences used were 5'-TGACAAGCATCACTCAGT and 5'-TTTGGCACCCTGTCTTCG (corresponding to bases 72063-72082 and 72101-72120, respectively, for the sequence of the genomic locus FQ377760). TALEN cleavage led to several different independent mutations in founder fish. The largest deletion was an 11-bp segment of exon 2, 5'-AGCTATCAAGA. This deletion eliminates the Alu1 restriction site found in exon 2, and the deletion leads to a premature stop codon at amino-acid residue 101. This residue, 101, is predicted to lie within the non-canonical PH domain (https://www.uniprot.org/uniprot/A0A0PCP424protvista) Fig. 1A). Fish carrying this mutation were selected for outcrossing and further study.

carmil3sa19830 and carmil3t413 were genotyped by restriction enzyme digestion of PCR amplicons containing the mutations. For carmil3sa19830, the forward primer was 5'-ACGAGGTGTCCTCTTCTCCAC, and the reverse primer was 5'-GATGAGGTTGATGAGTAC. Msel digested distinguish WT (191 bp band) from sa19830 mutant (bands at 123 bp and 68 bp). For carmil3t413, the forward primer was 5'-AGATAGTTGAAATCTACATTTTTAACC, and the reverse primer was 5'-AGGAGAGGTGTAATACCTTTAAGTCTTCA (corresponding to bases 71935-71965 and 72249-72278, respectively, for the sequence of the genomic locus FQ377760). Alu digestion produced two bands at 157 bp and 187 bp from WT genomic DNA, and a single band at 333 bp from the mutant. Mutant founders were outcrossed into WT AB fish to produce heterozygous fish, which were fully viable. Mutant founders were outcrossed into WT AB fish and the resulting heterozygotes were crossed to produce in the next generation zygotic and maternal-zygotic (MZ) homozygous mutants, which were fully viable. Alternatively, genotyping of carmil3sa19830 and carmil3t413 mutants was performed by Transnetyx (Cordova, TN) using real-time PCR with allele-specific probes.

2.2. Protein expression and purification

Capping protein (CP, mouse alpha1beta2) was expressed and purified as described (Johnson et al., 2018). Glutathione-S-transferase (GST)-tagged CP binding region (CBR) fragments of human CARMIL1a (E964-S1078, plasmid pBJ 2411), human CARMIL3 (S955-S1063, plasmid pBJ 2449), zebrafish Carmil3 (S943-N1040, plasmid pBJ 2451) (Fig. 1A) and zebrafish Carmil3 CPT-mutant (S943-N1040, with point mutations H964A and R966A, plasmid pBJ 2452) were expressed from pGEX-KG vectors in E. coli BL21 Star (DE3). The fusion proteins were affinity-purified on Glutathione Sepharose® 4 Fast Flow (GE Healthcare), and then bound to POROS GoPure XS (Applied Biosystems) in 20 mM NaH2PO4, 100 μM EDTA, 1 mM NaN3, 5 mM DTT, 4M urea (pH 7.5). After elution with a gradient to 1 M NaCl in the same buffer, the purified GST-CBR fragments were concentrated and stored at –70°C.

2.3. Actin polymerization assays

Pyrene-actin polymerization assays were performed as described (Carlsson et al., 2004). Pyrene-labeled and unlabeled gel-filtered rabbit muscle actin stocks were mixed to produce a total actin monomer concentration of 1.5 μM in the cuvette. Pyrene-actin filament seeds were prepared as described (Ramabhadran et al., 2012). CP at 5 nM and GST-CBR at varied concentrations were added at the start of the experiment (0 s). Pyrene-actin fluorescence was measured using time-based scans on a steady-state fluorometer (QuantaMaster, PTI, Edison, NJ) with excitation at 368 nm and emission at 386 nm.

2.4. Whole-mount RNA in situ hybridization (WISH)

Digoxigenin-labeled antisense RNA probes for myl7 (cardiac myosin regulatory light chain) (Ye and Lin, 2013; Yelon et al., 1999), sox17 (sex determining region Y-box 17) (Alexander et al., 1999; Hu et al., 2018), and carmil3sa19830 were used in our Tg(sox17:EGFP) fish to label cardiac myosin filaments in the KV. (A) Representative images illustrating the patterns of cardiomyocyte labeled in Tg(sox17:EGFP) fish at 28 hpf. Red dotted line indicates the leading edge of the endoderm, used to measure migration distance. (B) Schematic diagram of zebrafish gastrula illustrating the distance of the leading edge of the migrating endoderm cells to DFCs. In the plot, each data point corresponds to one embryo. Values for the distributions were as follows (mean ± s.d.): WT 74 ± 25 (number of embryos, N = 50), MZcarmil3sa19830 mutant 144 ± 28 (N = 44). Asterisks (*) indicate p value of <0.0001 in Student's t-test. (C-G) In vivo migration analysis of EGFP-labeled endodermal cells performed by epifluorescence time-lapse experiments on carmil3sa19830/° or MZcarmil3sa19830 mutant embryos in the Tg(sox17:EGFP) background. (C, E) Snapshots at 80% epiboly stage from the time-lapse movie, with the tracked cells labeled. Scale bar: 100 μm. (C’, E’) Snapshots at 95% epiboly stage. (D, F) Migration tracks from the 80%-95% epiboly stage delineate routes of endodermal cells. Solid magenta squares denote the endpoint of migration. (G) Total speeds, net speeds, and migration persistence. The number of embryos and cells analyzed is indicated above the graphs. Asterisks (**) indicate p value of <0.0001 in Student's t-test.
Fig. 3. Patterns of dorsal forerunner cell (DFC) distribution in MZcarmil3sa19830 mutant embryos compared to control heterozygous MZcarmil3sa19830/– embryos. Panels A to E illustrate the patterns observed, with results quantified in panel F. In most control embryos, DFCs form a single tight cluster of normal size (light blue arrow in panel A and light blue bars in panel F). In some embryos, DFCs display three patterns of defects: B) normal size with mild splitting (i.e., two clusters); C) normal size with severe splitting (more than two clusters); D) small size with no splitting; E) small size with splitting. (F) Percentages of embryos displaying the patterns of DFC distribution in control gastrulae and both MZcarmil3sa19830 and MZcarmil3sa19830 mutants. The percentage of embryos in each group is indicated above each bar, and the total number of embryos in each group (N) is listed underneath the labels on the abscissa.

2.7. Rescue experiments

We attempted to rescue the reduced KV size phenotype by injection of a subphenotypic dose of synthetic WT carmil3 RNA into MZcarmil3sa19830/Tg(sox17:EGFP) 1-cell stage embryos. To define a subphenotypic dose, quantities from 2.5 to 100 pg were injected into WT Tg(sox17:EGFP), embryos and morphology assessed during gastrulation, 6-somite stage and at 24hpf. Doses equal to or less than 5 pg produced morphological normal embryos at these stages. All embryos were injected with a total of 12.5 pg of RNA. The difference between this and the dose of carmil3 RNA was made up with Histone2B-RFP RNA. Injected embryos were fixed at the 6-somite stage and imaged on a confocal microscope at Washington University as described below. Z-stacks were collected and maximum intensity Z-projected to measure area of the KV. The number of cells in the KV were counted in Z-stacks.

2.8. Microscopy and image analysis

For still epifluorescence images, live or fixed embryos were mounted in 2% methylocellulose and photographed using a Leica DMI 6000 microscope with a 5x NA 0.15 objective or a 10x NA 0.3 objective. For WISH images in Fig. 3, embryos were injected in 80% glycerol/PBS and photographed using a Leica DMI 6000 microscope with a 5x NA 0.15 objective or a 10x NA 0.3 objective. For Figs. 4 D,E and 5A-E, fluorescence images were collected on a spinning disk confocal microscope (Quorum, Canada) using an inverted Olympus IX-81 microscope, a Hamamatsu EMCCD camera (C9100-13) and Metamorph acquisition software. For Figs. 2A, 6 and 7, WISH images were collected and maximum intensity Z-projected to measure area of the KV. The number of cells in the KV were counted in Z-stacks.
were collected on a Nikon Macroscope with a Nikon AZ100 objective, a 1x lens N.A. 0.1, and a 4x lens N.A. 0.4.

For time-lapse imaging of endodermal cells in Fig. 2 C-G and Fig. 4 A-C, Tg(sox17:EGFP) embryos were embedded in 0.8% low-melting agarose in a dorsal-mount imaging mold as previously described (Ye et al., 2015). Time-lapse imaging was taken in the dorsal region of endoderm at 25 °C, at 5-min interval with a 5 × /NA 0.15 objective on an inverted Leica DMI 6000 microscope. Images were processed and cell tracking analyzed using ImageJ (Schneider et al., 2012). Data were exported to Excel where cell migration speed, paths, and direction were determined as previously reported (Lin et al., 2005).

2.9. Statistical analysis

Data were compiled from two or more independent experiments and were presented as the mean ± SEM or ± SD as indicated in figure legends. Statistical analyses were performed in GraphPad Prism (GraphPad Software) using unpaired two-tailed Student’s t-tests with unequal variance. The numbers of cells and embryos analyzed in each experiment and significance levels are indicated in graphs and/or the figure legends.

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**Fig. 4.** Morphology of Kupffer’s vesicle (KV) and cilia in carmil3 mutant embryos. (A) Line drawing of a 6-somite stage embryo in a lateral view with KV outlined in green. On the right side is an image of a Tg(sox17:GFP) labeled KV. (A’). Line drawing of a posterior view of the 6-somite stage embryo. This view was used to collect confocal images. On the right side is a maximum intensity z projected image showing KV. Panels B through C are from one analysis, based on wide-field fluorescence images showing the morphology of sox17-EGFP labelling of the KV in Plus - Slash - Minus. carmil3<sup>stl413/+</sup> control (B) and MZcarmil3<sup>stl413</sup> mutant (B’ and B’’) embryos at the 6-somite stage. The KV is outlined by the dashed yellow line. (C) Area of the KV in Plus - Slash - Minus. carmil3<sup>stl413/+</sup> control and MZcarmil3<sup>stl413</sup> mutant embryos. All the data points are shown. The red lines indicate the mean and one standard error of the mean. The results are statistically significant with a p value of <0.0001 based on Student’s t-test. The number of embryos analyzed was 15 for control and 30 for mutant. In this set of experiments, the control was a heterozygous strain. Panels D and E show results of the experiments testing if injected WT carmil3 RNA can rescue KV area in MZcarmil3 mutants. KV area was scored similarly to Fig. 4 panel B. (D) Data from individual clutches are presented in the same color, e.g., orange colors are injections into a WT clutch of a control RNA, 2.5, or 5 pg of WT carmil3 RNA. Variability in ranges correlates with different clutches. Clutches used for only one injection are in grays. (E) Unpaired Student’s T-test with unequal standard deviation for rescue experiments. We plotted average area for each injection condition, with Standard Error of the Mean. Comparisons with significant p values are indicated by bars.
3. Results

As part of our interest in how actin assembly and actin-based motility contributes to morphogenesis and cell movement during development, we examined the role of CARMIL3. The CARMIL family of proteins regulate the heterodimeric actin capping protein (CP) that controls polymerization of actin filaments at barbed ends, in vitro and in cells (Stark et al., 2017). To advance our understanding of the functions of the three CARMIL isoforms conserved across vertebrates, we first assayed biochemical activities of zebrafish Carmil3 in comparison to human CARMIL3 and Carmil2. Second, we altered the activity of the zebrafish carmil3 gene, previously known as lrrc16b and sick211-204d18.1, by creating and/or interrogating lines carrying homozygous loss-of-function mutations.

3.1. CARMIL3 inhibits actin capping activity of CP

The biochemical activities of the vertebrate CARMIL1 and CARMIL2 isoforms with respect to CP and actin have been studied in mouse and human systems (Stark et al., 2017). The ~100-aa capping protein binding region (CBR) of CARMIL1 and CARMIL2 are known to bind directly to CP and partially inhibit its actin capping activity, via an allosteric mechanism. Comparative studies of CARMIL3 had not been previously done and so we performed them here.

We prepared recombinant proteins corresponding to the CBR region of CARMIL3 from human and zebrafish (Fig. 1A), and we tested their ability to inhibit the capping activity of mouse non-sarcomeric CP (alpha1beta2) (Edwards et al., 2013). In actin polymerization assays seeded with barbed ends of actin filaments, CP inhibited actin polymerization (Fig. 1B). A human CARMIL3 (C3) CBR fragment partially reversed the inhibitory effect of CP, with a potency slightly greater than that of human CARMIL1 (C1) (Fig. 1B). Zebrafish Carmil3 (C3) CBR also inhibited CP (Fig. 1C), with a potency approximately half that of human CARMIL3. Specific point mutations known to impair the ability of human CARMIL3 to interact with CP (Edwards et al., 2013; Lamier et al., 2015) were also created in the CBR fragment of zebrafish Carmil3 (C3-HVR>-AVA) (Fig. 1A). The mutant CBR failed to inhibit CP activity (Fig. 1C). Thus, the CBR region of CARMIL3 from both human and zebrafish, is able to inhibit the capping activity of CP, in a manner consistent with the direct-binding mechanism described for other CARMILs (Stark et al., 2017). This result supports the hypothesis that functional properties of CARMIL3 in vertebrates may involve direct interaction with CP.

3.2. Genomic mutations of zebrafish carmil3: creation and characterisation

A carmil3 mutant line from the Zebrafish Mutation Project (Kettleborough et al., 2013), carmil3sa19830, carries a single-nucleotide change at an essential splice site, predicted to truncate the protein to 832 residues from its WT length of 1384 residues (Fig. 1A). We created a second carmil3 mutant line by TALEN-mediated mutagenesis, carmil3stl413; this mutation produces an 11-bp deletion within exon 2 that causes a frameshift followed by a nonsense mutation, predicted to produce a truncated protein of 101 residues (Fig. 1A). Both mutant alleles are predicted to lack the Capping protein Binding Region (CBR), essential for Carmil3′s biochemical interaction with CP. Heterozygous and homozygous zygotic mutants of both alleles were viable as embryos and adults and did not present any overt phenotypes. When fish homozygous for either of the two alleles were crossed, the resulting embryos, deficient in both maternal and zygotic carmil3 function, MZcarmil3sa19830/a19830 (thereafter MZcarmil3sa19830) or MZcarmil3stl413/a19830 (thereafter MZcarmil3stl413), also completed epiboly and gastrulation and developed into morphologically normal embryos. Using qRT-PCR we determined that during gastrulation the level of carmil3 transcripts in MZcarmil3stl413 mutants was not significantly different from that of wild-type (WT) embryos (data not shown).

3.3. Migration of endodermal cells

To assess potential subtle or transient developmental defects, we analyzed the expression of germ layer markers in MZcarmil3 mutants at mid-gastrulation by whole-mount in situ hybridization (WISH). When examining the positions of endodermal cells and DFCs, marked by expression of sox17, at mid-gastrulation (70–80% epiboly), we noted that the leading edge of the vegetally migrating endodermal cells in MZcarmil3sa19830 mutant embryos lagged behind that of WT embryos (Fig. 2A). To quantify the effect, we measured the distance from the front edge of the endodermal cells to the edge of the aggregation of DFCs, as diagrammed on the right side of Fig. 2B. The distance was larger for the MZcarmil3stl413 mutant, compared to WT embryos (Fig. 2B), indicating that carmil3 is important for endodermal migration. The mean distance values were 67 μm (95% confidence interval of 60–74, number of embryos, N = 61) for WT embryos and 142 μm (95% confidence interval of 135–150, N = 55) for mutant embryos. The difference between the two sets of data had a two-tailed p value of <0.0001 in an unpaired t-test. These data were combined from clutches obtained on separate days.

To test this observation directly in living embryos, we then examined the migration of endodermal cells by conducting epifluorescence microscopy time-lapse experiments using a transgenic line expressing eGFP fluorescent protein from the sox17 gene promoter Tg(sox17:Egfp) (Fig. 2) (Mizoguchi et al., 2008). Cell tracking revealed that in control embryos (heterozygous carmil3p[a19830]/a19830), endodermal cells migrate toward the vegetal pole and converge dorsally along fairly straight paths (Schmid et al., 2013) (Fig. 2C and D). In contrast, in carmil3-deficient mutants (MZcarmil3stl413), the endodermal cells took less direct paths (Fig. 2E and F). Further cell tracking analyses showed that both cell movement, i.e. (total speed: movements in all directions) and migration efficiency (net speed: along straight line between the start and endpoint) were impaired in mutants compared to WT controls, as was the persistence of migration (ratio of net: total speed) (Fig. 2G). Thus, Carmil3 is required for efficient migration of endodermal cells.

3.4. Aggregation of dorsal forerunner cells (DFC)

Static images from these time-lapse experiments with a Tg(sox17:Egfp) background revealed the migration and cluster formation by dorsal forerunner cells (DFCs), which strongly expressed eGFP. DFCs in WT embryos usually formed a single large cluster (Fig. 2C). In mutant embryos, the DFC cluster was fragmented; we observed this in MZcarmil3stl413 (Fig. 2E) and MZcarmil3stl413 mutant embryos (data not shown). In addition, we noted bright eGFP-labeled cells that resided in an apparent gap between the DFC cluster and the leading edge of the migrating endodermal cells (Fig. 2E) in the mutant, but not WT embryos, which we suggest correspond to DFCs exhibiting abnormal migration.

We further examined DFC cluster morphology and DFC distribution during gastrulation by performing WISH for sox17, which also labels DFCs in embryos at 70–80% epiboly (midgastrulation; Fig. 3). In WT embryos, DFCs usually formed a single coherent cluster (Fig. 2C). In mutant embryos, the DFC cluster was fragmented; we observed this in MZcarmil3stl413 (Fig. 2E and F). Further cell tracking analyses showed that both cell movement, i.e. (total speed: movements in all directions) and migration efficiency (net speed: along straight line between the start and endpoint) were impaired in mutants compared to WT controls, as was the persistence of migration (ratio of net: total speed) (Fig. 2G). Thus, Carmil3 is required for efficient migration of endodermal cells.
3.5. Morphology of the Kupffer's vesicle and cilia

The cluster of DFCs that forms during gastrulation, undergoes a morphogenetic transition into an epithelial sac known as Kupffer's vesicle (KV), the ciliated organ that determines L/R asymmetry during zebrafish development (Eissner et al., 2005; Gokey et al., 2016). Because we observed a deficit in DFC aggregate formation, we examined the morphology of the KV and its cilia during segmentation (Fig. 4).

To measure the size of the KV, we labeled the KV and endoderm with Tg(sox17:EGFP) and collected images using widefield epifluorescence or confocal fluorescence microscopy at the 6-somite stage. As the Tg(sox17:-EGFP) transgenic line expresses eGFP in the cytoplasm of the KV epithelial cells, it allowed us to measure a total KV area including its epithelium (Fig. 4A, A’, B, B’). A posterior view (A’) provided the largest area, which we measured with FIJI/ImageJ. At the 6-somite stage, the KVs in MZcarmil3stl413; Tg(sox17:EGFP) mutant embryos were significantly smaller than those in Plus - Slash - Minus. carmil3sa19830 control embryos (Fig. 4C), revealing a requirement for Carmil3 in the KV development.

To test if Carmil3 could restore KV area, we injected different doses of carmil3 synthetic RNA into one-celled WT and MZcarmil3ed413 mutant embryos and analyzed their overall morphology during gastrulation and measured the KV size at 6-somite stage (Fig. 4D). As a control, embryos were injected with an equal mass of Histone2B-RFP RNA. These experiments revealed that zebrafish embryos are highly sensitive to carmil3 RNA dose. We observed that injections of 2.5 pg/embryo of carmil3 RNA significantly increased the KV size in WT embryos. However, higher doses (5 and 10 pg) did not further increase the KV size (Fig. 4D and E) and injections of 10 pg carmil3 RNA caused defects during epiboly and abnormal embryo morphology at 1dpf (not shown). Injections of carmil3 RNA also increased the KV size in MZcarmil3ed413 mutant embryos, but only at 10 pg/embryo (Fig. 4D and E). These results indicate that Carmil3 promotes KV size in both WT and MZcarmil3ed413 mutant embryos, but at a higher dose in the latter. Moreover, in contrast to WT embryos, injections of 10 pg of carmil3 RNA into MZcarmil3ed413 mutant embryos did not cause any morphological defects. The difference in sensitivity of MZcarmil3 mutants and WT embryos to carmil3 RNA is consistent with the levels of functional carmil3 transcripts being lower in the mutants.

To visualize the epithelial KV morphology and the KV cilia during later development, we stained embryos fixed at the stage of 10–14 somites with antibodies against the epithelial marker atypical PKC (aPKC) (Amack et al., 2007) and acetylated tubulin (Ac-tubulin) (Piperno and Fuller, 1985), resolving the cilia as individual structures, allowing us to count their number and measure their length. In these confocal microscopy images, the hollow empty center of the KV corresponding to its lumen appears as a relatively dark unstained region in the aPKC channel and as staining positive for cilia in the Ac-tubulin channel. In instances where the KV did not “inflate” (MZcarmil3ed413 in Fig. 5A, bottom panel), an unstained region was not observed. As observed for the DFC cluster in earlier stages (Fig. 3), and the total KV area at the 6-somite stage (Fig. 4C), the KV lumen was often smaller or fragmented, resulting in a decreased KV area for embryos homozygous for either of the two mutant alleles (Fig. 5B). The decrease in KV lumen area was statistically significant (Fig. 5B), and the magnitude of the decrease was sufficiently large that the values were at or below the value for the area of the KV at this stage found to be necessary for robust L/R patterning by Gokey and colleagues (Gokey et al., 2016). Values for the two mutants differ from the value for wild type based on Student's t-test (p < 0.005 (actual = 0.0006 and 0.0005)). (B) Area of cilia per KV. Horizontal dotted line corresponds to the value for area of functional cilia per KV (30) found to be necessary for robust L/R patterning by Gokey and colleagues (Gokey et al., 2016). Values for the two mutants differ from the value for wild type based on Student's t-test (p < 0.005 (Actual = 0.0001 and 0.001)). (D) Length of cilia. Values for the two mutant lines do not differ from the value for WT embryos based on Student's t-test (p = 0.3277 and > 0.9999). Values of N (embryos counted) as follows: WT, 9; MZcarmil3ed413, 8; MZcarmil3ed413, 4 for panels B to D. (E) Panel E shows a second experimental series with a different set of animals than those in panels A–D. Left side: The number of cells in the KV at the 6-somite stage is reduced in MZcarmil3ed413 (29.4) relative to WT (32.5), without statistical significance. Each was injected with a control RNA (~12.5 pg of Histone2B-RFP). Right side: The total area of KV is proportional to the number of cells. The same KVs examined for the cell number on the left were measured for area on the right.
The decreased number of cilia in the KV of MZcarmil3 mutants could be due to a reduced number of the KV constituent cells. To test this hypothesis, we imaged, at the 6-somite stage, WT and MZcarmil3 mutants that harbored the Tg(sox17:EGFP) transgene and were injected with synthetic H2B-RFP RNA, so that they were injected with the same mass of RNA as embryos in the following rescue experiments (Fig. 4A).

While the difference does not reach statistical significance, there is a trend toward fewer cells in the assembled KV in the mutant (Fig. 5E, left), and there are more mutant KVs of small cell number and small area (Fig. 5E, right). Finally, our measurements of the KV size and count of cell number revealed that the KV area was proportional to the number of KV cells in both WT and mutant conditions (Fig. 5E, right), suggesting that KV with fewer cilia (Fig. 5B and C) are also smaller in area.

### 3.6. Left-right patterning

Kupffer's Vesicle is the embryonic organ of laterality in vertebrates (Essner et al., 2005). Because we observed defects in formation of the KV, including its morphology, size and the number of its constituent cells and cilia; we asked whether further development revealed defects in L/R asymmetry pathways and outcomes. Since the MZcarmil3 mutants displayed KV area and number of cilia below the thresholds shown to be required for robust L/R patterning by previous studies (Gokey et al., 2016; Sampaio et al., 2014), we hypothesized that there would be defects in L/R patterning in these mutants.

First, we examined the pattern of expression of the lateral plate mesoderm marker southpaw (spaw) in embryos at the 18–20 somite stage with spaw staining distribution at the 18–20 somite stage in carmil3 mutant embryos compared with WT embryos. (A) Representative images illustrate observed patterns, which is purple. (B) Percentage of spaw staining patterns, comparing WT embryos with embryos of two different carmil3 alleles, MZcarmil3stl413 and MZcarmil3sa19830. The color scheme is indicated in panel A, below the images. In cases scored as “absent,” no staining was observed. Values are listed in Table 1.
Table 1
Pattern of expression of spaw at 18–20 somites, comparing carmil3 maternal zygotic (MZ) mutant embryos with WT embryos. The percentage of the total number of embryos with different patterns are listed, along with 95% confidence intervals (CI) and the number of embryos scored (N). Statistics, including confidence intervals, were calculated with GraphPad Prism.

<table>
<thead>
<tr>
<th>Genotype</th>
<th>Phenotypic Pattern, Percent of Total (95% CI)</th>
<th>N</th>
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<tbody>
<tr>
<td>Wild type</td>
<td>98 (94–99) 1 (0–3) 2 (0–5) 0 172</td>
<td></td>
</tr>
<tr>
<td>MZcarmil3stl413</td>
<td>54 (46–62) 9 (5–15) 26 (20–34) 11 (7–17) 137</td>
<td></td>
</tr>
<tr>
<td>MZcarmil3stl10630</td>
<td>53 (43–64) 7 (3–14) 31 (22–41) 9 (5–17) 88</td>
<td></td>
</tr>
</tbody>
</table>

(60) (Long et al., 2003). Whereas WT embryos uniformly displayed spaw staining on the left, the two carmil3 mutant lines showed substantial percentages of embryos with bilateral or right-sided spaw staining. Representative examples are shown in panel A of Fig. 6, with quantitation in panel B and Table 1. Results for the two mutant lines were similar to each other and differed significantly from results for WT embryos. These embryos were generated from the same set of mutant and WT animals as those used in the experiments illustrated in panels A–C of Fig. 5, where the KV area and number of cilia per KV were below the critical threshold in the mutant. When a different generation of mutant animals was used in a separate set of experiments to generate MZcarmil3embryos, in which the KV area was not below the critical value illustrated in Fig. 5B, little or no defects in spaw staining distribution were observed (data not shown), consistent with the findings of Gokey and colleagues (Gokey et al., 2016), and Sampaio and colleagues (Sampaio et al., 2014) noted above.

spaw regulates FGF signaling (Neugebauer and Yost, 2014), and spaw mutants are defective in L/R asymmetry, as revealed by the position of the heart (Ahmad et al., 2004; Long et al., 2003). Therefore, we asked whether mutant embryos had a phenotype related to heart position. Indeed, mutant embryos at ~30 hpf displayed a defect in the L/R positioning of the heart (Fig. 7). In a noticeable and significant number of embryos, the heart was in the middle or on the right side of the embryo, based on staining for cardiac myosin light chain (myl7), for which representative examples are shown in panel A of Fig. 7. A combined count of embryos with altered heart position showed defects of 6–9% for each of the two mutant alleles and a value less than 1% observed in WT embryos (Fig. 7, Panel B). The defects in the two mutant lines did not differ from each other by a statistically significant degree, and the differences between the two mutant lines and the WT line was highly significant in both cases.

As above for spaw staining, these data for heart position are from a set of animals and experiments where the KV area and number of cilia per KV in the mutant were below the critical threshold (Fig. 5A–D). In a separate set of experiments, using MZcarmil3stl413 mutants, in which the KV area was not below the critical value (Fig. 4B and C), little or no defects in heart position were observed (data not shown). Therefore, the penetrance and expressivity of the KV phenotype in MZcarmil3stl413 mutants correlated well with those of the spaw asymmetric staining defect during mid-segmentation stages and the L/R heart positioning defect at the end of embryogenesis, consistent with the findings of Gokey and colleagues (Gokey et al., 2016) and Sampaio and colleagues (Sampaio et al., 2014).

4. Discussion
We investigated the role of the actin assembly regulator, CARML3, in cell migration and morphogenesis during early zebrafish development. First, we confirmed that in actin polymerization assays, human CARML3 and zebrafish Carmil3 interact with and regulate the activity of capping protein with respect to actin polymerization. More important, we discovered that Carmil3 is required for normal migration of endodermal cells and the migration and/or aggregation of dorsal forerunner cells (DFCs) during zebrafish gastrulation. Impaired aggregation of DFCs led to defects in the formation of the KV in terms of its shape and size. These KV defects were associated with decreased numbers of cells and cilia present in mutant KVs; as a consequence, L/R asymmetry was impaired in some mutants, manifested by malposition of the marker spaw during segmentation and later in the position of the zebrafish heart tube. These KV and L/R phenotypes displayed variable but correlated penetrance and expressivity.

The two carmil3 mutant alleles we report here were generated by reverse genetic approaches: carmil3stl413 by TILLING for ENU-induced nonsense mutations (Kettleborough et al., 2013) and carmil3d413 by deploying TAL effectors (Boch et al., 2009; Moscou and Bogdanove, 2009). Both are premature stop codons predicted to produce truncated proteins lacking the Capping protein Binding Region (CBR), essential for Carmil3’s interaction with CP (Fig. 1), and thus should represent strong or null alleles. Consistent with that view, the phenotypes are recessive. However, our study does not include an analysis of the protein products that would test this view definitively. The observation that only maternal zygotic but not zygotic carmil3 mutants presented gastrulation phenotypes, is consistent with known significant maternal carmil3 expression, which is likely to compensate for the lack of zygotic expression during gastrulation (Solnica-Krezel, 2020; Stark and Cooper, 2015). The lack of full penetrance for the later phenotypes indicates a substantial level of robustness in the processes of cell migration and morphogenesis, consistent with observations for many other genes regulating early embryogenesis (Chen et al., 2018; Kelly et al., 2000; Li-Villarreal et al., 2015; Solnica-Krezel and Driever, 2001). CARML3s are encoded by three conserved genes in vertebrates, including zebrafish, so redundant and overlapping functions contributed by carmil1 or carmil2 may also account for the variable level of penetrance seen for downstream phenotypes. In addition, as the two carmil3 mutant alleles are nonsense mutations, the observed phenotypes could represent only partial loss-of-function and variability due to genetic compensation triggered by RNA degradation (El-Brolosy et al., 2019; Ma et al., 2019). However, against this notion, our qRT-PCR studies indicated that carmil3d413 mutant transcripts were expressed at relatively normal levels in MZ mutants during gastrulation.

Our results provide new information about the function of CARML3, especially in the in vivo context of a whole vertebrate organism during the process of embryogenesis. Previous studies on CARML3 have used cell culture and mouse tumor models to uncover roles in neuronal synapse formation and cancer cell migration, based on actin assembly (Hsu et al., 2011; Lanier et al., 2016; Spence et al., 2019; Wang et al., 2020).

Previous studies, with human cultured cells that express both CARML1 and CARML2, found overlapping but distinct functions for the two proteins, based on subcellular localization and knockdown phenotypes (Liang et al., 2009; Stark et al., 2017). While the early phenotypes observed here display strong penetrance, the later phenotypes are far less penetrant, raising the question of compensatory and overlapping function among the CARML1-encoding genes. In support of the view, preliminary observations in our laboratories have revealed stronger L/R asymmetry phenotypes in double mutant zebrafish embryos that carry mutations in genes for both CARML2 and CARML3 (Stark, Solnica-Krezel and Cooper, unpublished); these observations will merit further study in the future.

We discovered that Carmil3 is important for the migration of endodermal cells during zebrafish gastrulation. Mutant endodermal cells had both reduced overall motility as well as reduced persistence. Recent work demonstrated cultured cells lacking CARML3 had reduced migration in the classical scratch assay and in trans-well migration. KO cells were less polarized, had less polymerized actin and fewer focal adhesions (Wang et al., 2020). Interestingly, the Rac-specific guanine nucleotide exchange factor, Prex1, was implicated in regulation of Nodal-dependent actin dynamics and random endoderm cell motility during gastrulation (Woo et al., 2012). Future experiments will determine if endodermal migration defects in MZcarmil3 mutants are associated with abnormal actin organization and/or focal adhesion formation.

We found that Carmil3 is important for the aggregation of DFCs during gastrulation. DFCs typically migrate vegetalward ahead of the

germ layers in one cluster of cells. Occasionally a few cells may separate from the cluster. DFC coalescence and migration as a cohesive cluster is strongly dependent on cell-cell adhesion. cdh1/E-cadherin (halfbaked/volcano) mutants in which cell adhesion is reduced, exhibit delayed epiboly of all germ layers and frequently fragmented DFC clusters (Shimizu et al., 2005; Solnica-Krezel et al., 1996). A recent study showed loss of CARML3 in cultured cells inhibits cadherin-based adhesion through transcriptional downregulation of epithelial type gene expression (Wang et al., 2020). Such a mechanism might account for the reduced adhesion of DFC in vivo. We observed smaller and malformed KV, both its total area and lumen, likely the direct result of the smaller DFC clusters. However, we cannot exclude a more direct role of CARML3 in KV morphogenesis.

CARMLs appear to regulate actin via direct biochemical interactions with and effects on the actin-capping properties of CP (Edwards et al., 2013; Lanier et al., 2015; Stark et al., 2017). Indeed, CP itself is known to have an important role in morphogenesis. Mutations in humans and in zebrafish of the gene encoding the beta subunit of CP, known as capzb in zebrafish, were found to cause craniofacial and muscle developmental defects, with effects on cell morphology, cell differentiation and neural crest migration (Mukherjee et al., 2016). In addition, the same study found that capzb overexpression produced embryonic lethality.

Among regulators of CP, CARMLs are only one family of proteins with CPI motifs. Among other CPI-motif protein families, which are unrelated to each other outside of their CPI motifs, several have been shown to have roles in actin-based processes of development. The CPI-motif protein CKIP-1 is important for myoblast fusion in mammalian and zebrafish systems (Baas et al., 2012). CapZIP, known as duborysia/dub in zebrafish, is important in zebrafish development for actin organization in the kidney glomerulus, in mammals and zebrafish (Hentschel et al., 2007; Tossidou et al., 2019). Zebrafish CIN85, a homologue of CD2AP encoded by the gene sh3kb1, is also important for glomerular podocyte function (Teng et al., 2016), and has a role in the formation and maintenance of the vascular lumen (Zhao and Lin, 2013). Finally, the twinfilin family of CPI-motif proteins, encode by four genes in zebrafish - twf1a, twf1b, twf2a and twf2b- has not been as well-studied, but may also affect actin assembly, as may the WASHCAP/Fam21 family of CPI-motif proteins, encoded by wasch2c in zebrafish.

The carml3 mutant lines generated here provide a valuable tool for studying the regulation of actin dynamics during endoderm and DFC migration and KV morphogenesis in the context of a developing vertebrate embryo, and for testing potential functional interactions with CARML3 interacting proteins.

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References


