Variability of an early developmental cell population underlies stochastic laterality defects

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Summary

Embryonic development seemingly proceeds with almost perfect precision. However, it is largely unknown how much microscopic variability is hidden beneath this macroscopic accuracy. Here, we quantified embryo-to-embryo variability in vertebrate development, by studying cell number variation in the zebrafish endoderm. We noticed that the size of a sub-population of the endoderm, the dorsal forerunner cells (which later form the left-right organizer), is highly variable between individual embryos. We found that the frequency of left-right laterality defects is increased drastically in embryos with a low number of dorsal forerunner cells, and we observed that these fluctuations are largely nonhereditary. Hence, a stochastic variation in early development leads to a remarkably strong macroscopic phenotype. These fluctuations appear to be caused by variable deposition of maternal factors involved in specification of the dorsal forerunner cells. In summary, we here dissect cause and consequence of embryo-to-embryo variability in a vertebrate model.

Introduction

During embryogenesis, cells acquire their identity due to a combination of external signaling cues and internal competence factors. Embryonic development is remarkably robust towards fluctuations of regulatory factors such as morphogen levels (Barkai and Shilo 2009; Briscoe and Small 2015) or genetic variation (El-Brolosy et al. 2019). However, developmental

buffering of fluctuations is not perfect, and phenotypic variation can even be observed in mutants from isogenic *C. elegans* strains raised under identical environmental conditions due to noisy gene expression and stochastic variation in genetic interaction partners (Burga, Casanueva, and Lehner 2011; A. Raj et al. 2010).

Differences in the size of progenitor populations may be another important source of phenotypic variability in higher organisms. However, the degree of variability in cellular ontogenies and their potential phenotypic consequences remain largely unknown. Some notable exceptions are: changes in the subdivision of the primordium that gives rise to the head sensory organs lead to fluctuations in ommatidia number at different levels (inter-individual, inter-strain and inter-specific) in the *Drosophila* genus (Ramaekers et al. 2019; Gaspar et al. 2020); sexual dimorphism and left-right asymmetry of ommatidia number in the ant *Temnothorax albipennis* are related to differences in mating and motion behavior, respectively (Hunt et al. 2018). Here, we set out to systematically quantify the degree of inter-individual cell number variation and its phenotypic consequences, using endoderm specification in the early zebrafish embryo as a model system.

The endoderm, which is induced by high levels of Nodal signaling during early development, contributes to the formation of liver, pancreas, intestine, stomach, pharynx and swim bladder (Warga and Nüsslein-Volhard 1999). The dorsal forerunner cells (DFCs) are a small group of cells, considered a subset of the endoderm (Alexander et al. 1999; Warga and Kane 2018) (Figure 1A). They are the precursors of the Kupffer's vesicle (KV) (Melby, Warga, and Kimmel 1996), the organ that determines left-right asymmetry in the embryo (Essner et al. 2005) (Figure 1B). Since the endoderm is the smallest germ layer during early zebrafish development (Shah et al. 2019) and the DFCs comprise only a few dozen cells (Oteíza et al. 2008; Gokey, Dasgupta, and Amack 2015), we asked if naturally occurring embryo-to-embryo variation in cell numbers in wild-type embryos could have any phenotypic consequences.

53 **Results**

54 To characterize cell number variability during zebrafish early embryogenesis, we counted the
55 number of endodermal cells and DFCs at the 75% epiboly stage using a Tg[*sox17*:GFP] reporter
56 line (Figure S1A-D). We found that the number of endodermal cells exhibits considerable
57 variation, with ~500-800 cells per embryo (Figure 1C). However, the number of DFCs is even
58 more variable, ranging from ~10 to ~50 cells per embryo (Figure 1D). This variation of DFC
59 numbers persists at a later stage when they have formed the KV (Figure 1E, Figure S1E),
60 suggesting that no corrective mechanisms are triggered in the animals with particularly high or
61 low numbers of DFCs. We did not find a positive correlation between total cell number and
62 either the endoderm or the DFCs population (Figure 1C, D), indicating that this variation is not
63 due to staging differences. Furthermore, we found no association between the number of DFCs
64 and the number of endodermal cells (Figure S1F), which suggests that the fluctuations of these
65 two cell populations have independent sources.

66 The surprisingly large fluctuations in DFC numbers prompted us to investigate possible
67 phenotypic consequences of this variation at later developmental stages. Since the DFCs give
68 rise to the KV, the organ establishing left-right asymmetry, we focused on investigating
69 possible laterality defects. Previous experimental studies, as well as mathematical modeling,
70 suggested that the size of the KV needs to exceed a certain threshold in order to enable reliable
71 left-right patterning (Sampaio et al. 2014; Gokey et al. 2015). As the heart is the first organ that
72 is formed during zebrafish development, and since its laterality can be assessed easily in live
73 embryos, we examined the percentage of embryos with defective heart laterality (DHL) in
74 different clutches of embryos (Tüpfel long-fin (TL) wildtype strain) at prim-22 stage (Figure
75 2A; Video 1). We found a DHL average of 3.9% [standard deviation (sd): 3.0%, 53 independent
76 observations (IO), total amount of scored embryos (n): 6081]. Even though this is a remarkably
77 high frequency for a wildtype population, this value is in very good agreement with previous

78 reports based on *in situ* hybridization of lower sample numbers (Wang et al. 2011: 3%, n = 650;
79 Noël et al. 2013: 6%, n = 387).

80 We then evaluated whether a change in the environmental conditions could unmask an
81 underlying sensitivity in individuals that would otherwise present normal organ laterality.
82 Previous reports have observed an influence of temperature on the penetrance of zebrafish
83 mutant phenotypes (Imai et al. 2001). The incubation temperature for zebrafish embryos ranges
84 from 25°C to 33°C, with 28.5ºC being the (Kimmel et al. 1995). Increasing the temperature to
85 33ºC led to a remarkable increase of the DHL average to 26.4% (sd = 9%, IO = 89, n = 9557)
86 (Figure 2A). We observed no difference in the mortality rate, and the fraction of abnormal
87 embryos (which were excluded from analysis in any condition) remained moderate (4.6% and
88 12.1% of the embryos showed a swollen heart cavity or tail defects at 28.5°C and 33°C,
89 respectively). Taken together, these observations suggest that establishment of left-right
90 laterality in zebrafish is relatively unstable, which we hypothesized might be due to the variable
91 number of DFCs.

92 To test this idea, we set out to determine whether the measured fluctuations in DFC
93 numbers are directly linked to the observed laterality variation. To do so, we counted the
94 number of DFCs by live microscopy at 60% epiboly stage and assessed the laterality of the
95 heart and the liver at later stages, after incubation at 33ºC. We found that the number of DFCs
96 at 60% epiboly is strongly predictive of laterality defects by long pec stage (Figure 2B), which
97 establishes a direct association between DFC numbers and laterality defects. Of note, we
98 observed that the fraction of embryos with laterality defects is ~50% for those individuals with
99 less than ~20 DFCs, suggesting that laterality is established randomly (and hence half of the
100 time correctly) in embryos with low DFC number. In summary, we found that an early
101 fluctuation of the size of a small cell population is correlated with a macroscopic defect at later
102 stages, rather than be being corrected over the course of development.

103 We then aimed to exploit the influence of temperature as a tool to better understand the
104 mechanism of how variability in DFC numbers influences the frequency of heart laterality
105 defects. To this end, we tested several temperature shift treatments spanning early development
106 (1-cell to bud stage), early somitogenesis (bud to 14-somites stage) and organogenesis
107 (following 14-somite stage) (Figure 2C). Interestingly, we found that treating the embryos
108 during early somitogenesis alone, the time at which the KV is formed, was sufficient to generate
109 a similar DHL frequency as with continuous incubation at 33°C (Figure 2C, III). This result
110 suggests that the temperature treatment mostly influences KV function (and hence global left-
111 right patterning). However, temperature does not have a major influence on specification of
112 DFCs, which happens before somitogenesis, or heart looping, which happens after
113 somitogenesis. To corroborate this hypothesis, we decided to investigate potential defects of
114 global left-right patterning mediated by the KV. Rotational movements by cilia create a
115 directional fluid flow in the extracellular space that triggers the activation of Nodal signaling
116 on the left lateral plate mesoderm by inducing degradation of the Nodal antagonist *dand5* on
117 the left side. This leads to expression of the Nodal ligand *southpaw* (*spaw*) only on the left side
118 of the lateral plate mesoderm (Hashimoto et al. 2004), which is the molecular signal used to
119 establish organ laterality. Indeed, we found that asymmetric expression of *dand5* was reduced
120 at 33ºC compared to 28.5ºC (Figure S2A,B), suggesting that the elevated temperature interferes
121 with the patterning activity of the KV.

122 To gain further insight into how the number of cells is related to the function of the KV,
123 we measured the size of the KV as well as the number of cells (Figure S2C-E). We found that,
124 as expected, the size of the KV and the number of cells are correlated (R^2 = 0.78 for 28.5ºC,
125 0.58 for 33ºC). However, while the number of cells remained similar, the size of the KV lumen
126 was significantly decreased at 33ºC compared to 28.5ºC. Taken together, these results suggest
127 that an elevated temperature leads to a lowering of the cell threshold required for reliable

128 functioning of the KV, which is mediated at least partially via a reduction of the size of the
129 lumen.

130 In line with these observations, we found a gradient of expression patterns for the early
131 left-right marker gene *spaw*, ranging from the expected pattern (i.e. *spaw* only on the left) to
132 bilateral expression (*spaw* visible on both sides, although not necessarily at the same level) and
133 complete reversal (i.e. *spaw* only on the right) (Figure 2D). We hypothesized that these defects
134 on the level of *spaw* expression should produce concordant laterality defects in different organs
135 that exhibit left-right asymmetry, such as the heart and liver. Indeed, we found that in most, but
136 not all, cases the embryos with reversed heart laterality also exhibited reversed liver laterality
137 (Figure 2E). We speculate that discordant organ laterality might occur in case of almost
138 perfectly balanced bilateral *spaw* expression (i.e., the same expression levels on both sides),
139 leading to a scenario where in some cases the two organs would develop their laterality largely
140 randomly and independently from each other.

141 After observing this strong and macroscopically visible phenotype caused by cell
142 number fluctuations at very early developmental stages, we wanted to understand the origin of
143 the DFC number fluctuations. First, we set out to distinguish stochastic fluctuations from
144 genetically determined variation of cell numbers. To identify possible genetic factors, we raised
145 embryos with a reversed heart loop to adulthood and evaluated the DHL frequency on their
146 progeny. As a control, embryos with normal heart laterality were raised as well. To our surprise,
147 we detected no significant differences in the percentage of reversed heart laterality in the
148 offspring compared to the control group, at either 28.5°C or 33°C (Figure 3A), suggesting that
149 this phenotype is largely non-genetic.

150 The stochastic origin of the observed laterality defects is a striking finding, with
151 potentially important conceptual consequences for our interpretation of variable disease
152 phenotypes (see Discussion). However, the stochastic nature of the phenotype numbers makes
153 it more difficult to identify the molecular origin of DFC number fluctuations. We therefore tried

154 to identify general principles that underlie DFC number variability. As both the DFCs and the
155 surrounding dorsal domain are induced by high levels of Nodal in a non-cell-autonomous
156 manner (Oteíza et al. 2008; Hagos and Dougan 2007), we reasoned that there could be a direct
157 correlation between the number of cells forming the dorsal organizer and the DFCs number;
158 however, we didn't find such an association (Figure S3A,B). Furthermore, we only found a
159 weak association between maternal age and DHL frequency at 28.5ºC (Figure S3C).

160 Next, since DHL has been considered a spontaneous strain-dependent defect (Malicki
161 et al. 2011), we compared the frequencies of reversed heart laterality between embryos with TL
162 genetic background (the one used so far) versus AB. Interestingly, we found significantly lower
163 DHL frequencies for AB embryos at 28.5ºC (1.3%, sd = 1.3%, IO = 22, n = 2714) and 33°C
164 (7.7%, sd = 4.1%, IO = 34, n = 3353) compared to TL (Figure 3B). In line with this observation,
165 we found that the number of DFCs and KV cells was higher in AB versus TL (Figure 3C), and
166 that the antero-posterior cilia distribution is more asymmetric in AB than in TL (Figure S3D,E).

167 The strain-specific differences in DFC numbers gave us a handle to explore the
168 molecular mechanism in more detail. Specifically, we hypothesized that the fluctuations might
169 be maternally controlled, since DFCs begin to be emerge 1 hour after maternal-to-zygotic
170 transition. To test this hypothesis, we crossed AB females with TL males and vice versa. Indeed,
171 we found that the number of DFCs depends mostly on the mother's genetic background, as the
172 cell numbers of the two hybrid crosses were different from each other (Figure 3D , with the TL
173 (female) / AB (male) cross resembling the TL/TL cross, and the AB (female) / TL (male) cross
174 resembling the AB/AB cross (Figure 3C,D). Consequently, a major source of the observed
175 fluctuations in the embryo must lie in the development of the oocytes in the mother. When
176 comparing TL and AB embryos by RNA-seq, we found 94 genes that were consistently
177 differentially expressed in the period before and after the zygotic genome activation (2.25 hpf
178 and 3.25 hpf, respectively) as well as the time window during which the DFCs are specified
179 (4.25 - 5.25 hpf) (Figure 3E, Supplemental Figure 4, Table 1). 12 of the differentially expressed

180 genes are reported to have an early expression in specific cell types or structures (Figure 3F;
181 ZFIN database of gene expression patterns). It is very likely that most of these differentially
182 expressed genes are related to processes other than DFC specification that differ between the
183 two strains. However, we were intrigued to find that 7 of these 12 genes with a reported early
184 spatial pattern are expressed in cell types related to DFC specification – the enveloping layer
185 (EVL), from which the DFCs are derived (Cooper and D'Amico 1996; Oteíza et al. 2008), and
186 the yolk syncytial layer (YSL), which remains connected to the DFCs until 4hpf (Cooper and
187 D'Amico 1996).

188 We found a high correlation in the expression levels of several of the differentially
189 expressed YSL/EVL genes across the TL embryos (Figure 3G, Supplemental Figure 4). This
190 raises the possibility that variation of some upstream factor (or a combination of factors) acting
191 during oocyte development may be responsible for the observed variable but correlated
192 expression. Interestingly, the mutant for one of these genes, *ctsba,* has been reported to have a
193 reduced number of EVL cells (Langdon et al. 2016). However, it is unlikely that the genes
194 identified here are the only ones responsible for variable DFC numbers, since additional factors
195 (such as fluctuations in protein levels) may contribute equally. Furthermore, additional genes
196 without a reported YSL/EVL expression might also be involved in this phenomenon. In
197 summary, these experiments suggest that variable deposition of maternal factors, possibly
198 involved in YSL/EVL specification and function, may be responsible for the observed
199 fluctuations in DFC numbers.

200

201 **Discussion**

202 We found that stochastic fluctuations of a small cell population generated during the earliest
203 stages of embryogenesis (the DFCs) are not corrected over the course of development, but
204 instead persist and give rise to a macroscopic phenotype (defects of organ laterality). While it
205 was reported before that left-right patterning in wildtype zebrafish is surprisingly variable (Noël

206 et al. 2013; Wang et al. 2011) and that there is a size threshold for reliable functioning of the
207 KV (Sampaio et al. 2014; Gokey et al. 2015), we now show that these phenomena are linked
208 and originate in fluctuations of DFC numbers that are at least partially caused by variable
209 deposition of maternal factors.

210 Here we show that the number of cells in the KV is important for proper function.
211 However, other studies have shown that the internal architecture is equally critical, since a
212 higher density of cilia in the anterio-dorsal region is needed to create a proper leftward flow
213 (Kreiling et al. 2007; Okabe, Xu, and Burdine 2008). We observed differences in cilia density
214 ratio between AB and TL strains (Figure S3E), but no correlation to the cilia number (data not
215 shown). If and how variation in DFC number is linked to morphological asymmetries in the
216 KV is an interesting question, which would require additional live experiments.

217 The sizes of progenitor populations in vertebrate embryos are inherently variable due to
218 environmental, genetic and stochastic fluctuations. Without feedback control, the size of the
219 final cell population would dependent linearly on the size of the progenitor pool and hence be
220 highly volatile (Lander et al. 2009). It is therefore remarkable that in this case there seem to be
221 no corrective mechanisms that reduce cell number variability and thereby ensure robust left-
222 right patterning. Furthermore, while mouse embryos pass a size checkpoint at around the time
223 of gastrulation (Snow and Tam 1979; Lewis and Rossant 1982), the externally developing
224 zebrafish embryos don't seem to pass such a checkpoint and are apparently evolved to
225 maximize the speed of development, even at the cost of occasional laterality defects.

226 Here we investigated the consequences of developmental variability in a specific cell
227 population, the DFCs, and it remains to be answered how general such phenomena are in other
228 cell populations. How variable vertebrate development is, and which degree of fluctuations is
229 still compatible with normal development, are important open questions, which will require
230 novel experimental approaches such as high-throughput lineage tracing (Spanjaard et al. 2018;
231 B. Raj et al. 2018; Alemany et al. 2018).

While stochastic developmental defects of this frequency and severity are probably not very common in wildtype mammals, it is likely that similar mechanisms as the one described here may underlie incomplete penetrance of mutations in model organisms (A. Raj et al. 2010; Burga, Casanueva, and Lehner 2011) as well as in humans. Furthermore, there is increasing evidence that disease phenotypes that manifest late in life, like Alzheimer's disease, may already originate in development (Arendt, Stieler, and Ueberham 2017). A similar phenomenon may occur in type I diabetes – an emerging view is that the initial pool of beta cells at the end of development is variable between individuals, leading to increased disease susceptibility in a subset of the population (Atkinson et al. 2015).We therefore speculate that early stochastic fluctuations of progenitor cell pools might at least partially contribute to a variety of human disease phenotypes.

Author contributions

J.P.J. and R.M.A. conceived and designed the project. R.M.A performed experiments and analyzed the data, with support by P.O.C. and R.S.. J.P.J. guided experiments and data analysis. All authors discussed and interpreted the results. R.M.A. and J.P.J. wrote the manuscript, with input from P.O.C..

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Declaration of interests

The authors declare no competing interests.

284 **Figures**

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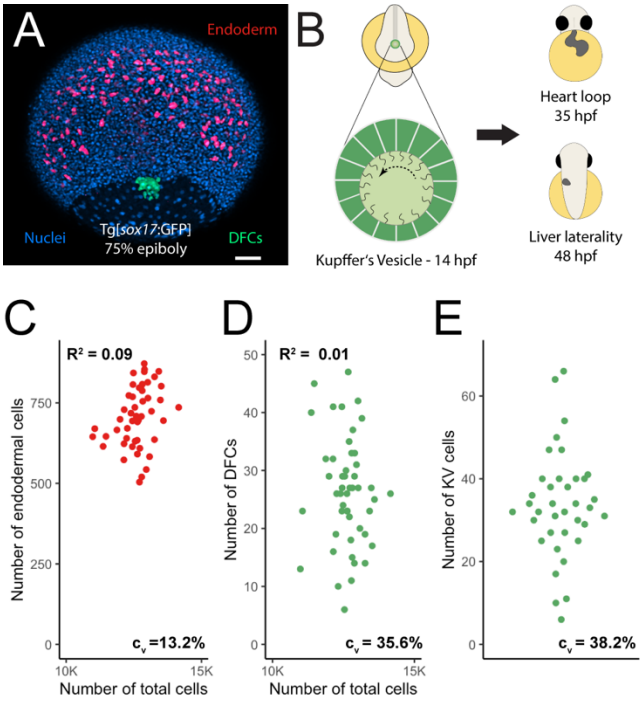

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287 **Figure 1. Cell number variability during early embryogenesis.** (A) Maximum projection of
288 confocal images of a Tg[*sox17*:GFP] embryo at 75% epiboly stage showing the endoderm (red),
289 DFCs (green) and nuclei (blue). Scale bar = 80µm. (B) Graphical representation of and 8-somite
290 stage embryo and the KV (based on figures from Dasgupta et al. 2018), together with its role
291 in determining organ laterality. (C-E) Cell number in endoderm (n = 49), DFCs (n = 49) and
292 KV (n = 37) for individual embryos. The coefficient of variation (C_V) is indicated at the bottom
293 right, (C-D) also show the total cell number on the x-axis (C_V = 4.8%) and the coefficient of
294 determination (R^2).

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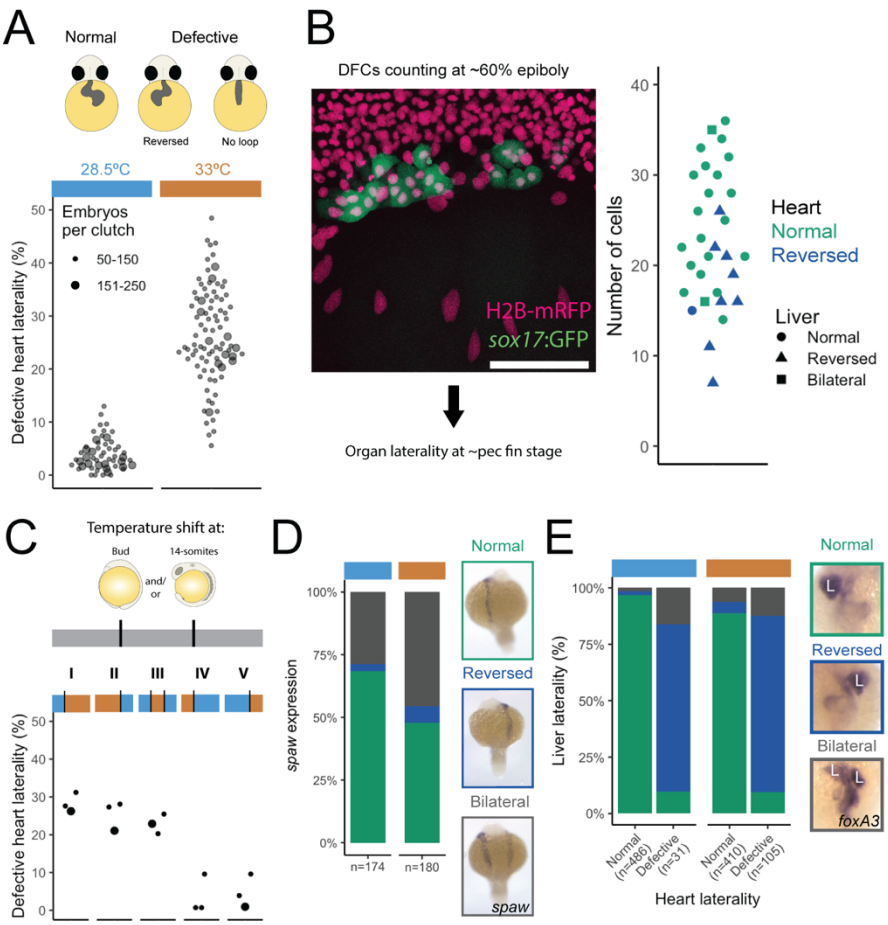

Figure 2. Fluctuations of DFC numbers lead to defects of organ laterality. (A) Heart laterality phenotypes scheme observed at prim-22 stage: normal (d-loop) and defective, which includes reversed (s-loop) and no heart loop, observed in embryos incubated at 28.5ºC (blue) and 33ºC (orange). The column scatter plots show the percentage of embryos with defective heart laterality per individual clutch analyzed at 28.5º (marked by the horizontal light blue bar) and 33ºC (orange bar). The small circles indicate a clutch size of 50-150 and the bigger ones a range of 151-250 embryos. (B) Confocal z-projection of the dorsal side of a Tg[*sox17*:GFP] (green) live embryo injected with H2B-mRFP (pink). The plot shows the DFC number distribution at 60% epiboly and the resulting heart and liver laterality in the individual embryos assessed by *in situ* hybridization (n = 31). (C) Graphical representation of the different temperature shift treatments: the first third of the gray bar represents the period between 1-cell and bud stage, the second until the 14-somite stage, and the third until the period of collection (long pec stage). Circle size as in (A). Bud and 14-somite schemes are based on Kimmel et al.

315 1995. (D) Relative frequency of different *spaw* expression patterns in embryos incubated at
316 28.5º and 33ºC. *in situ* hybridization photographs of normal (green), reversed (dark blue) and
317 bilateral (gray) *spaw* expression in 18-somite stage embryos are shown to the right. (E) Relative
318 frequency of embryos with normal, reversed or bilateral liver, separated into embryos with
319 normal or defective heart laterality (incubated at 28.5º or 33ºC). To the right: Liver laterality as
320 measured by *foxA3 in situ* hybridization.
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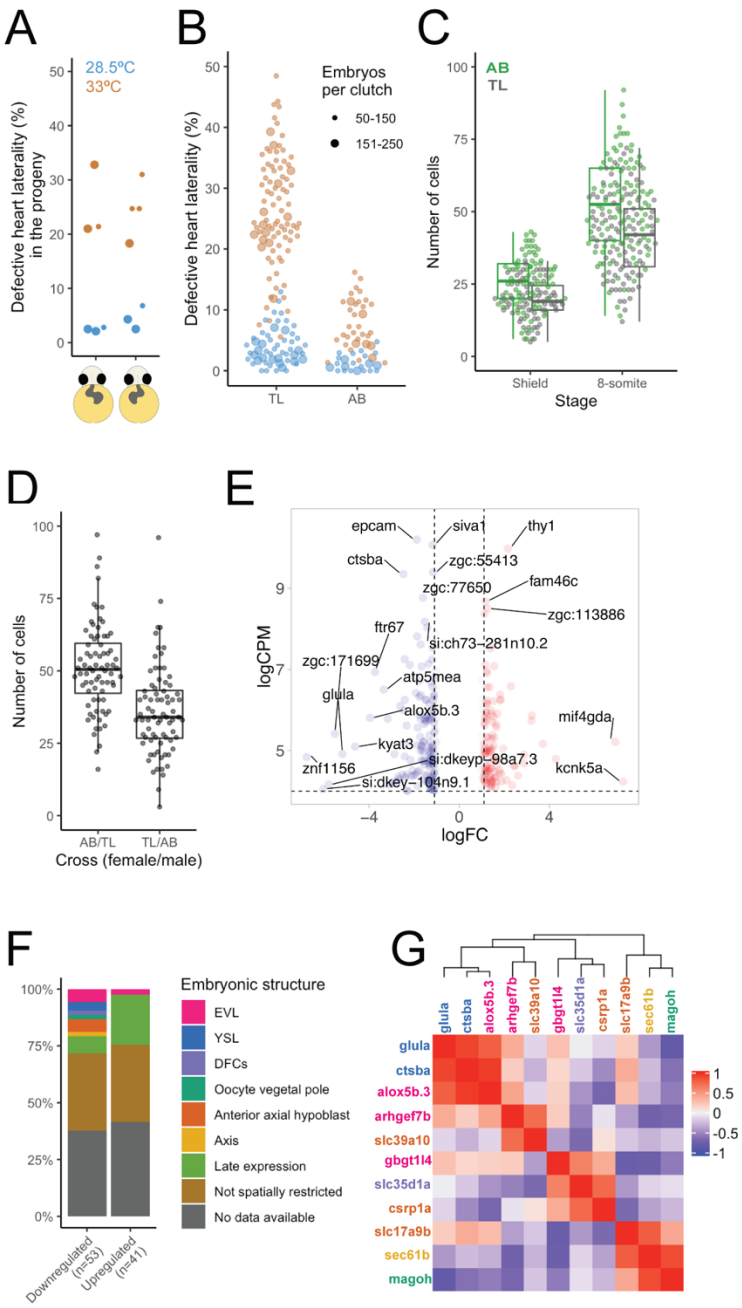

Figure 3. Heart laterality defects are a stochastic phenomenon that is linked to variable deposition of maternal factors. (A) Percentage of embryos with defective heart laterality in the progeny of individuals that showed either normal or reversed heart laterality at prim-22 stage. Incubation temperature 28.5ºC (light blue) or 33ºC (orange). The small points indicate a clutch size ranging of 50-150 and the bigger points for a range of 151-250. (B) Percentage of embryos with defective heart laterality per individual clutch analyzed at 28.5º and 33ºC in TL (same data as Figure 2A) and AB embryos. (C) Number of DFCs / KV cells at shield and 8-

349 somite stage; for shield stage: n = 78 and 75 for AB and TL, respectively, $p < 0.001$; for 8-
350 somite stage: n = 94 and 97 AB and TL, respectively, $p < 0.001$). (D) KV cell numbers at 8-
351 somite stage in crosses between individual AB and TL males and females, AB/TL vs TL/AB p
352 <0.001 (n=76 for both); AB vs AB/TL p = 0.461; TL vs TL/AB p = 0.01; AB vs TL/AB p
353 <0.001; TL vs AB/TL $p < 0.001$. (E) Volcano plot ($\log_2$ of the counts per million (CPM) and
354 fold change (FC)) showing differentially expressed genes in TL compared to AB embryos at
355 2.25 hpf determined by RNA-seq. The top 20 DE genes are shown. (F) Summary of reported
356 embryonic expression patterns for the down- and upregulated genes. (G) Pairwise Pearson
357 correlation between the downregulated genes at 2.25hpf for TL embryos. The gene names are
358 color-coded by embryonic structure as in (F). Color scale to the right. The boxplots in (C) and
359 (D) display the median; the hinges represent the first and third quartiles and the whiskers
360 represent 1.5 of the inter-quartile range from the hinge.
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375 **Methods**

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377 **Zebrafish Husbandry and Staging**

378 All animal procedures were conducted according to local authorities guidance (LAGeSo,
379 Berlin, Germany). Adult zebrafish were maintained and bred under standard conditions.
380 Embryos were left to develop in egg water (0.6 g/L dissolved in dH2O; Red Sea Salt, Red Sea,
381 containing methylene blue, 0.002 g/L) to the desired stage at 28.5C unless otherwise stated.
382 Staging was done based on Kimmel et al. 1995. It was not possible to record the sex of the
383 embryos examined due to their developmental stage.

384

385 **Zebrafish strains**

386 AB and Tüpfel Long Fin (TL) strains were obtained from the European Zebrafish
387 Research Center (EZRC). The Tg[*sox17*:GFP] strain was reported in (Sakaguchi et al. 2006).
388 To change its genetic background to TL, the fish were crossed with TL fish. By crossing the
389 resulting males with TL females, we obtained similar DHL percentages as in the TL wildtype
390 strain.

391

392 **Immunolocalization**

393 Embryos at the desired stage were fixed overnight at 4ºC in PFA 4% in PBS. The
394 following day, they were washed 3x with PBSTx (1x PBS with TritonX 1%) for 5 minutes and
395 dechorionated. They were incubated in blocking buffer (2% BSA, 5% Goat Serum in PBSTx)
396 for 2 hours at room temperature (RT), followed by an overnight incubation at 4ºC with one or
397 a combination of the following primary antibodies diluted in blocking buffer: mouse anti-
398 acetylated Tubulin (1:200, Sigma Aldrich T6793), chicken or rabbit anti-GFP (1:1000, Abcam
399 ab13970 and ab290, respectively), rabbit anti-Flh (1:200, Sigma Aldrich SAB2702443) and
400 rabbit anti-Laminin-α1β1γ1 (1:100, Sigma Aldrich L9393). On the next day, they were washed

401 3x with PBSTx and 1x with blocking buffer, 30 minutes each at RT, followed by an overnight
402 incubation at 4ºC with Hoechst 1:1000 and secondary antibody diluted in blocking buffer
403 (1:200): goat anti-chicken Alexa-488 (Thermo Fisher A11039), goat anti-mouse Alexa-647
404 (Thermo Fisher A32728) or goat anti-rabbit Alexa-568 (Thermo Fisher A11011). Finally, 3x
405 PBSTx washes, 30 minutes each at RT.
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407 **Imaging**
408 Endodermal, DFCs and total cell number in 75% epiboly embryos: after anti-GFP
409 immunolocalization, the embryos were washed with 50% glycerol/50% PBSTx for ten minutes
410 and then with 100% glycerol overnight at 4ºC. For flat mount, a layer of laboratory labeling
411 tape (13mm wide) was attached to each side of a coverslip to make a bridge and leave enough
412 room for the flat embryo (~120µm). A single embryo was put on a coverslip with a drop of
413 glycerol. For flattening, a closed forceps was introduced in the vegetal pole and then opened to
414 break and split the yolk cell in half. As many yolk granules as possible were removed to avoid
415 damaging the blastoderm, then another coverslip was put on top. The tissue was oriented facing
416 the bottom to be imaged with an SP8 inverted confocal microscope. After obtaining an image
417 stack for the endodermal and total cell number, another stack was obtained for the DFCs region
418 with lower laser power and higher zoom factor, since the GFP signal intensity in these cells is
419 considerably higher than in the endodermal cells. SP8 microscope acquisition parameters: 20x
420 multi-immersion objective, format = 1024 x 1024, speed = 600, zoom factor = 0.75 (except for
421 DFCs, 2 or 4 when all cells fit), line average = 2, Z-step size = 3, gating = 0.3 - 6 for both
422 channels, laser intensity Z-compensation for each image, tiling = 2x2 in most images, with 15%
423 overlapping. The images were automatically stitched with the Leica software.
424 For imaging DFCs at shield, cilia at 8-somite stage and total amount of cells at different
425 stages (3.25-5.25hpf), the embryos were mounted in 1% low melting point agarose, and imaged
426 on an upright Zeiss 880 confocal microscope, with the following settings: 20x water-dipping

427 objective, format = 512 x 512, speed = 8, zoom factor = 1, line average = 2, Z-step size = 3. For
428 the total amount of cell counting a 2-photon Chameleon laser was used, tiling = 2x2, with 15%
429 overlapping. The images were automatically stitched with the Zeiss software.
430
431 **Live imaging**
432 For KV measurements at 8-somite stage, embryos incubated at 28.5º or 33ºC were
433 mounted on a 1.5% agarose injection dish with little liquid. The embryos were oriented inside
434 the chorion to image the KV. Afterwards, 3% methylcellulose in E3 embryo medium (5mM
435 NaCl, 0.17mM KCl, 0.33mM $CaCl_2$, 0.33mM $MgSO_4$) was added to cover the embryos and,
436 thus, retain the order of the individual embryos. The embryos were incubated at the initial
437 temperature until prim-22 for heart laterality analysis. Per session, around 30 embryos were
438 imaged for each condition.
439 For DFC counting, individual Tg[*sox17*:GFP] (injected with ~60 pg of H2B-mRFP
440 mRFP1 at one-cell stage) shield stage embryos were transferred to glass-bottom Petri dishes,
441 most of the liquid was removed and ~0.5 mL of 3% methylcellulose in E3 embryo medium was
442 added. Then, the embryos were manually dechorionated with forceps and oriented with the
443 shield facing the bottom of the dish for imaging on the Leica SP8 confocal microscope. After
444 imaging, the dish was filled with E3 embryo medium and incubated at 33ºC until long pec stage
445 and fixed for *in situ* hybridization.
446
447 ***in situ* hybridization**
448 Whole-mount in situ hybridizations were performed essentially as described previously
449 (Thisse & Thisse, 2008). The following probes were used: *sox32* (Bjornson et al. 2005); *foxA3*
450 and *myl7* (Noël et al. 2013); and *spaw* and *dand5* (Sampaio et al. 2014).The *amhc* probe was
451 kindly provided by Daniela Panáková.
452

453 **mRNA synthesis**

454 The H2B-mRFP1 mRNA was synthesized using the mMESSAGE mMACHINE kit
455 (Thermo Fisher AM1345) according to the manufacturer's recommendations.

456

457 **Temperature shift treatment**

458 After collection, the embryos were put immediately in warmed water, either to 28.5ºC
459 or 33ºC. For the temperature shift, the embryos were transferred to a Petri dish filled with water
460 at the desired temperature.

461

462 **Single embryo RNA-seq**

463 A total of 6 AB and 6 TL embryos were individually collected per stage (2.25, 3.25,
464 4.25 and 5.25 hpf) in LoBind tubes (Eppendorf) in two independent experiments (each of which
465 contained half of the samples for each condition). 0.5mL Trizol and 0.5µl GlycoBlue were
466 added to each samples, and RNA extraction was carried according to the manufacturer's
467 recommendations. Each sample was barcoded, pooled and the libraries were prepared according
468 to the CEL-seq2 protocol (Hashimshony et al. 2016) with different RPI indices for each
469 timepoint and paired-end sequenced on an Illumina NextSeq 500 (read length 12 nt for barcode
470 read and 63 nt for transcript read).

471

472 **Sequencing data analysis**

473 Basecalling was done with bcl2fastq v2.19.0.316. The resulting reads were
474 demultiplexed with *scruff* (Wang et al. 2019). Mapping was done with STAR 2.7.1a (Dobin et
475 al. 2013) using quantMode GeneCounts and the DanRer11v96 transcriptome as reference.
476 Differentially expressed (DE) genes were obtained with edgeR (Robinson, McCarthy, and
477 Smyth 2009). Genes with a False Discovery Rate < 0.01, log Counts Per Million > 4 and log
478 Fold Change < -1 for downregulated and > 1 for upregulated genes were considered DE. Spatial

479 and temporal expression was obtained on The Zebrafish Information Network (Bradford et al.
480 2011).
481
482 **Plots**
483 All plots were generated in R using the following additional libraries: ggplot2 graphing
484 package, ggbeswarm for column scatter plots, ggExtra for marginal histograms and
485 ComplexHeatMap for heatmaps; for which, a logarithmic transformation of the counts per
486 million (CPM) values obtained with edgeR was applied first. Volcano plots were generated
487 with VolcaNoseR (Goedhart and Luijsterburg 2020).
488
489 **Cell counting**
490 The endodermal cells were counted with the Imaris (Bitplane) Surface module:
491 background subtraction: on, diameter of largest sphere: 15 µm, splicing touching: on, seed
492 points diameter: 8 µm. For each sample, the number was visually corroborated. The total cell
493 number was estimated with the Fiji (Schindelin et al. 2012) ITCN plugin. A Z-projection was
494 obtained for each stack, the ITCN parameters were: width 12, minimum distance 6, threshold:
495 0.5.
496 For quantification of dorsal domain cells and DFCs, cell number was estimated
497 manually for cilia with the Cell Counter Fiji plugin. The cilia number was used as a proxy for
498 the KV cell number, since the cells are monociliated. DFCs numbers were first estimated with
499 ITCN (width 18, minimum distance 9, threshold: 0.5) and visually corrected afterwards. For
500 the total amount of cell estimation, the cells were counted with Imaris, same parameters
501 mentioned above.

502 The Flh positive cells were counted with ITCN plugin (same parameters used for DFCs mention
503 above). In all cases the embryos were imaged immediately after immunolocalization. The

504 number of dorsal domain cells was obtained by subtracting the DFCs number from the number
505 of Flh positive cells, since Flh is also expressed in the DFCs.

506 **Statistics**

507 The coefficients of determination were obtained with R. The p-values are obtained with a
508 randomization test done with PlotsOfDifferences (Goedhart 2019). All statistical parameters,
509 including samples numbers and median are shown in the figures, described in the figure legends
510 or in the main text.

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529 **Supplemental files**

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531 **Video 1. Heart loop phenotypes observed at prim-22 stage.** Heart laterality phenotypes on
532 live embryos; normal/dextral loop (d-loop); defective includes: reversed/sinistral loop (s-loop)
533 and no loop, shown side by side on a ventral view.

534

535 **Table 1. Shared DE genes.** Function, earliest expression, spatial expression, DE expression,
536 and embryonic structure information from ZFIN for the differentially expressed genes found
537 across the timepoints selected for AB vs TL comparison.

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