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Relationship Between Physiological Umbilical Herniation and Liver
Morphogenesis During the Human Embryonic Period: A Morphological and
Morphometric Study

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Running title: PUH and liver morphogenesis

Grant sponsor: Japan Society for the Promotion of Science;

Grant numbers: 26220004, 16K15535, 17H05294, and 18K07876.

1

2 Abstract

3

4 It is widely hypothesized that physiological umbilical herniation (PUH) in humans
5 occurs because the liver occupies a large space in the abdominal cavity, which
6 pushes the intestine into the extra-embryonic coelom during the embryonic
7 period. We have recently shown the presence of the intestinal loop in the
8 extra-embryonic coelom in embryos with liver malformation. Here, we analyzed
9 the relationship between the liver and PUH at Carnegie stage 21 of four embryos
10 with liver malformation, including two with hypogenesis (HY1, HY2) and two with
11 agenesis (AG1, AG2), using phase-contrast X-ray computed tomography and
12 compared them to two control embryos. The intestinal loop morphology in the
13 malformed embryos differed from that in the control embryos, except in HY1.
14 The length of the digestive tract in the extra-embryonic coelom of the embryos
15 with liver malformation was similar or longer than that of the controls. The rate of
16 intestinal loop lengthening in the extraembryonic coelom compared to that of the
17 total digestive tract in all embryos with liver malformation was similar or higher
18 than that of the controls. The estimated total abdominal cavity volume in the
19 embryos with liver malformation was considerably smaller than that of the
20 controls, while the intestinal volume was similar. The cardia and proximal portion
21 of the pancreas connecting to the duodenum were located at almost identical
22 positions in all the embryos whereas other parts of the upper digestive tract
23 deviated in the embryos with abnormal livers. Thus, our results provided
24 evidence that PUH occurred independently of liver volume.

25

26 Key words: Physiological umbilical herniation, Liver morphogenesis, Liver
27 malformation, Total abdominal cavity volume, Intestinal volume.

1 Introduction

2
3 The human intestine lengthens much more rapidly during the
4 embryonic period (Meckel, 1817) and develops a dorsoventral hairpin fold called
5 the primary intestinal loop at the Carnegie stage (CS) 14 (Kim et al., 2003). The
6 intestine temporarily distributes in both the abdominal coelom and extra
7 embryonic coelom in the umbilicus. Such a condition was classically referred to
8 as physiological umbilical herniation (PUH) (Meckel, 1817; Mall, 1898). A
9 previous study showed that PUH begins from CS 17 (Ueda et al, 2016), and is
10 followed by reduction in which the intestine returns to the abdominal coelom
11 during the early-fetal period (Kim et al., 2003). Finally, the intestine assumes its
12 adult position by fixation of the mesentery. During these processes, the intestine
13 is hypothesized to undergo a counterclockwise rotation of 270°.

14 Mall (1899) advocated the most simple and probable cause of PUH as
15 the liver growing very rapidly and almost entirely filling the abdominal cavity. As a
16 result, it pushes the intestine into the extra-embryonic coelom. Recent textbooks
17 present descriptions consistent with Mall's theorem (Gary et al., 2009; Moore et
18 al., 2008). However, ever since Mall's description was put forth, no data
19 indicating the actual mechanism of PUH have been reported.

20 We have recently shown human embryonic samples at CS 21 with liver
21 malformation in which the intestinal loop is within the extra-embryonic coelom
22 (Kanahashi et al., 2016). We doubted Mall's description because the intestinal
23 loop was even within the extra-embryonic coelom of embryos with hypogenetic
24 and agenetic livers. Detailed analyses of these embryos may be valuable for
25 reconsidering the mechanism of PUH, which could occur due to reasons other
26 than the growth and occupation of the liver in the abdominal cavity. Thus, in the
27 current study we aimed to examine the precise relationship between the liver
28 and PUH using both morphological and morphometric approaches by comparing
29 embryos with liver hypogenesis, liver agenesis, and normal liver. Herein, we
30 discuss the mechanisms underlying the occurrence of PUH.

33 Materials & Methods

Human embryo specimens

Approximately 44,000 human embryos comprising the Kyoto Collection of Human Embryos are currently stored at the Congenital Anomaly Research Center of Kyoto University (Nishimura et al., 1968; Yamada et al., 2006; Shiota et al., 2007). Most of the embryos were obtained after pregnancy termination during the first trimester for socioeconomic reasons under the Maternity Protection Law of Japan. The current study was approved by the ethics committee of the Kyoto University Graduate School and Faculty of Medicine (E986). In the laboratory, the aborted embryos were measured, examined, and staged using the criteria for Carnegie staging described by O’Rahilly and Müller (1987), which is used internationally (Moore, 2008). Staging was based on the external and internal morphological development of the embryo, independent of both age and size.

Four embryos consisting of two embryos with liver hypogenesis (HY1, HY2) and two embryos with liver agenesis (AG1, AG2), as well as two control embryos, all at CS 21, were used in the study. These embryos were selected and described in part for our previous study (Kanahashi et al, 2016). Briefly, 1,156 magnetic resonance imaging (MRI) datasets at CS 14 to CS 23 were selected from the Kyoto collection as externally normal embryos (Matsuda et al., 2003; Shiota et al., 2007; Yamada et al., 2010). The 1,156 embryos were screened using liver volume as the target organ. Embryos with liver volumes 2 standard deviations above or below the mean for the stage of development were selected and examined using MRI. Embryos identified with potentially abnormal livers were further analyzed by using phase-contrast X-ray computed tomography (PXCT). Complications in the organs other than the liver are summarized in Table 1. Four embryos with liver anomaly showed defects partly at the outside of umbilical cord (e.g. Wharton’s Jelly), while the digestive tract, umbilical arteries, vein, and proximal part of the umbilicus and abdominal wall remained intact. The two control embryos were normal, both externally and with respect to liver morphology. The previous study elucidated that the prevalence of

liver malformations in CS18 and CS21 in the intrauterine population of externally normal embryos is approximately 1.7% (Kanahashi et al, 2016). Most of such liver abnormality embryos do not survive because liver function becomes essential (Koga, 1971; Migliaccio et al., 1986; Tavian et al., 1999).

Acquisition of 3D images with PXCT

The 3-D PXCT images of the human embryos were obtained using a radiographic imaging system (BL14-C, 17.8 keV) from Photon Factory, Institute of Materials Structure Science, High Energy Accelerator Research Organization (KEK, Tsukuba, Japan). The data provided a resolution of 18 $\mu\text{m}/\text{pixel}$. The mechanism and conditions used to acquire the PXCT images of embryos have been described elsewhere (Yoneyama et al., 2011).

Image analysis and morphometry

The structures of the digestive tract, including the esophagus, stomach, duodenum, intestinal loop, and colorectum, were reconstructed for each image set using the Amira software suite (version 5.5.0; Visage Imaging, Berlin, Germany). The length of the digestive tract was measured as described previously (Ueda et al., 2016). Briefly, the AmiraSkel software module was used to determine the centerline length of the digestive tract. The lengths of the total digestive tract (Total length) were set from the pyloric sphincter to the end of the intestinal loop. Since the end of the intestinal loop may correspond to the point at which the length of the mesentery drastically changes, as described by Soffers et al. (2015), it was defined in the current study as the most obvious dorsal inflection point on the 3D image. The length of intestinal loop in the extraembryonic coelom (extra length) was measured in the 3-D image from the point where the intestinal loop first entered the extraembryonic coelom to the last point of return to the abdominal coelom. The estimated total abdominal cavity volume and liver volume were measured by manual segmentation of the transverse section on sequential 2-D images using the Amira software. For the

total abdominal cavity volume, the slices including the cranial end of the diaphragm through the lower pole of the metanephros were used. The estimated intestinal volume was calculated by subtracting the liver volume from the estimated total abdominal cavity volume.

Results

Morphological features of the intestinal loop

The digestive tracts, livers, and other intra-abdominal organs were clearly detectable in the PXCT images of all embryos (Fig. 1). No internal damage was detected in any of the embryos. The elongated intestines entered the extra-embryonic coelom in all the embryos examined, both in the controls and in the embryos with liver hypogenesis or agenesis (Figs. 1 and 2).

The cecum was located to the left of the superior mesenteric artery (SMA) in the controls and in all malformed embryos except AG1 in which the cecum was located close to tip of the intestinal loop (Fig. 1). The intestinal loop had a secondary loop in the extra-embryonic coelom (Figs. 1 and 2). The number of secondary loops was the same in all embryos except HY2, which showed a greater number of secondary loops compared to that in the controls (Table 1, Fig. 2).

The proximal part of the intestinal loop ran straight across at the border between the abdominal coelom and extra-embryonic coelom in the two control embryos and AG1 (see red circle in Fig. 1 and red dashed line in Fig. 2). The secondary loop was located on the border between the abdominal coelom and extra-embryonic coelom in HY1, HY2, and AG2. The distal part from the cecum of the intestinal loop, which is equivalent to the large intestine, ran straight across the border between the abdominal coelom and the extra-embryonic coelom in all embryos except AG2 in which the intestinal tract from the cecum to the colorectum was not recognized (see * in Fig. 2).

Length measurements of the digestive tract

The Total length (a) and extra length (b) were measured (Fig. 3A) and the b/a ratio was calculated (Fig. 3B, Table 1). The Total length in control 1 and control 2 was 29 mm and 26 mm (mean 28 mm), respectively. In HY1 and HY2, the Total length was similar to that of the controls or longer (HY1, 27 mm; HY2, 33 mm). In AG1 and AG2, the Total length was shorter than that of the controls (AG1, 23 mm; AG2, 21 mm).

The extra length in control 1 and control 2 was 20 mm and 17 mm, respectively, with a mean of 19 mm. In HY1 and HY2, the extra length was longer than that of the controls (HY1, 23 mm; HY2, 22 mm). In AG1 and AG2, the extra length was similar to that of the controls (AG1, 19 mm; AG2, 18 mm).

The ratios of extra length to Total length in the controls were 69% and 65% (mean 67%). In HY1 and HY2, the ratios of extra length to Total length were similar to or higher than those in the controls (HY1, 84 %; HY2, 67 %). In AG1 and AG2, the ratios of extra length to Total length were higher than those in the controls (AG1, 82%; AG2, 87%).

Estimated abdominal volume

Images of the mid-sagittal sections of the abdominal cavity and its contents of control 1, HY1, and AG1 are shown in Fig. 4A. The liver and digestive tract occupied almost the entire abdominal space. The liver occupied the dominant amount of space in the controls. In contrast, the digestive tract occupied most of the abdominal space in the embryos with liver malformation.

Total estimated volumes of the abdominal cavities and intestines were calculated (Fig. 4B, Table 1). The estimated total abdominal cavity volumes in the controls were 63 mm³ and 50 mm³ (mean 57 mm³). Compared to those in the controls, the estimated total abdominal cavity volumes were markedly lower in the embryos with liver malformation (HY1, 30 mm³; HY2, 36 mm³; AG1, 21 mm³; and AG2, 18 mm³). Decreased liver volumes directly contributed to the decreases observed in the estimated total abdominal cavity volumes in the embryos with malformations.

1 In the controls, the intestinal volumes were 24 mm³ and 16 mm³ with a
2 mean of 20 mm³. In HY1 and HY2, the intestinal volumes were consistent with
3 those in the controls or higher (HY1, 23 mm³; HY2, 28 mm³). In AG1 and AG2,
4 the intestinal volumes were consistent with those in the controls or lower (AG1,
5 21 mm³; AG2, 18 mm³).

6 The volumes of the bilateral adrenal glands and metanephros were
7 also calculated. In the controls, the volumes of bilateral adrenal glands were 1.7
8 mm³ and 2.0 mm³ (mean 1.9 mm³) and the volumes of the bilateral metanephros
9 were 1.0 mm³ and 0.6 mm³ (mean 0.8 mm³). In the embryos with liver
10 malformation, the volumes of the bilateral adrenal glands were similar to those in
11 the controls or at most 1.0 mm³ higher (HY1, 2.0 mm³; HY2, 3.0 mm³; AG1, 2.6
12 mm³; AG2, 2.1 mm³). The volumes of bilateral metanephros in the embryos with
13 liver malformation were almost the same as those of the controls (HY1, 1.0 mm³;
14 HY2, 1.0 mm³; AG1, 0.8 mm³; AG2, 0.8 mm³). Morphological abnormality of the
15 bilateral adrenal glands and metanephros in embryos with liver malformation
16 were not detected (data not shown).

17 18 Deviation of the upper digestive tract (stomach, duodenum) and pancreas 19

20 To compare the deviation of the upper digestive tract and pancreas in
21 embryos with liver malformation as that of controls, we evaluated reconstructed
22 3-D images of the digestive tract in all the embryos (Fig. 5). Two anatomical
23 references, the cardia and proximal part of the pancreas connecting to the
24 duodenum, were located at almost identical positions in all the embryos with liver
25 abnormality and in the controls, indicating that no deviation developed around
26 these two landmarks. The SMA was located just dorsal to the proximal part of
27 the pancreas. The duodenum was connected to the posterior abdominal wall via
28 short dorsal mesentery except at the end of the duodenum. The duodenum was
29 directly connected to the posterior abdominal wall at the end of the duodenum
30 (Fig. 5).

31 However, other regions of the upper digestive tract, such as the body of
32 the stomach, the pylorus, the descending and horizontal parts of the duodenum,
33 and the body and tail of the pancreas, were deviated in the embryos with liver

malformation (Table 1). In HY2, the body of the stomach deviated ventrally and cranially, the descending part of the duodenum deviated caudally, and the horizontal portion deviated ventrally (data not shown). In AG1 and AG2, the body of the stomach deviated ventrally and cranially and was thus inverted. In addition to the inversion, deformation was also observed in body of the stomach of AG1 and AG2. In the embryos with agenesis (AG1, AG2), the digestive tracts from the pyloric antrum to the upper region of the duodenum were straight and ran vertically in the right ventral quadrant. As for the pancreas, the body and tail deviated ventrally and cranially in the embryos with liver malformation. The degree of deviation was greater in the embryos with agenesis (AG1, AG2) compared to that in the embryos with hypogenesis (HY1, HY2).

Discussion

Mall (1899) proposed rapid liver growth in the abdominal space as the simplest and most probable cause of PUH. Many studies have examined intestinal loop morphology and morphometry in the digestive tract during embryonic and fetal periods (Frazer and Robbins, 1915; Dott, 1923; Snyder and Chaffin, 1952, 1954; Kim et al., 2003; Metzger et al., 2011; Soffers et al., 2015). However, other than Mall's theory, there has been no cause of PUH described. The current study demonstrated an elongated intestinal loop distributed in both the abdominal coelom and the extra-embryonic coelom of CS 21 embryos with liver malformation, as well as in control embryos with normal liver morphology. To reconsider the cause of PUH, we examined the morphology and morphometry of the digestive tracts in embryos with liver malformations and in controls and estimated the abdominal volumes.

Ueda et al (2016) first measured the rapid elongation of the intestinal loop in the human embryonic period. Primarily owing to the small intestinal part of the intestinal loop, the intestinal tract elongated rapidly over this period. Most of the intestinal loop was morphologically observed in extra embryonic coelom, though the actual length for the extra embryonic part was not measured in that study. The present study revealed the intestinal length and morphological

1 features of the intestinal loop in embryos with liver malformations. Total length
2 and extra embryonic length were almost the same (21-33 mm and 17-23 mm) in
3 all the embryos, including those with liver malformation and the controls.
4 Furthermore, the ratio of extra length/total length was also almost same in the
5 embryos with liver malformations. These findings indicated that the elongation of
6 the intestinal tract as well as distribution of intestinal tract between extra coelom
7 and abdominal coelom was independent of liver development.

8 Mekonen et al (2015) described the development of the ventral body
9 wall using historical collections of embryos and fetuses. At CS 20, the dense
10 mesenchyme below the umbilicus is still identifiable but has split ventromedially
11 to surround the abdominal rectus muscles. They observed that the abdominal
12 muscles, including the rectus muscles, reach the hip bone at CS 23. Thus, the
13 structures of the abdominal wall remain immature at CS 21 and therefore the
14 developing abdominal muscles and mesenchymal tissue may still be soft and
15 prone to change in their form. On the other hand, the abdominal organs such as
16 the liver grow rapidly during the embryonic development (Hirose et al., 2012;
17 Kanahashi et al., 2016). Among the abdominal organs, the liver was the most
18 likely to expand the abdominal wall and to increase the abdominal volume.
19 There was less of a possibility that the abdominal organs other than the liver
20 were responsible for the increased abdominal volume. For instance, the volumes
21 of adrenal glands in the embryos with liver malformation were at most 1.0 mm³
22 larger than that of the controls. Such an increase seemed to be small and was
23 likely negligible. We speculate that it was primarily the liver that pushed to
24 change the form of abdominal wall. Correspondingly, a reduction in liver volume
25 may result in a decreased abdominal volume.

26 Two anatomical references, the cardia and the proximal part of the
27 pancreas connecting to the duodenum, were located at almost identical
28 positions in the controls and all the embryos with liver malformation, whereas the
29 body of the stomach and the body and tail of the pancreas were deviated in the
30 embryos with liver malformation. Deviation may be anatomically restricted in
31 these two references. The cardia was located close to the diaphragm in all
32 embryos. The septum transversum differentiates into the diaphragm after CS 17
33 (Hirose et al., 2012). Therefore, at CS 21, the movement of the cardia should be

1 restricted by the point at which the esophagus crosses the diaphragm. In
2 contrast, the position of the proximal part of pancreas may be restricted by
3 duodenum and SMA, whereas the passage of duodenum and orientation of
4 pancreas can still be deviated to some extent. Such deviations might occur by a
5 decrease (loss) in volume of the liver.

6 The present study demonstrated that the total and extra intestinal length
7 and its ratio were almost same in all embryos irrespective of the liver volume. In
8 contrast, a decrease in liver volume leads to a decrease in abdominal cavity
9 volume. These findings strongly indicate that PUH occurs due to near-normal
10 growth of the midgut and a near-normal increase in abdominal space for the
11 midgut. Therefore, Mall's theory should be reconsidered.

12 13 14 15 16 17 18 19 20 Acknowledgements

21 We are deeply indebted to Associate Professor Kazuyuki Hyodo of Photon
22 Factory for the use of the synchrotron radiographic equipment at BL-14C. We
23 thank Ms. Chigako Uwabe at the Congenital Anomaly Research Center for
24 technical assistance in handling the human embryos. We thank Dr. Tohoru
25 Takeda at Kitasato University for acquisition of the PCXT images. This work was
26 performed with the approval of the Photon Factory Program Advisory Committee
27 (Proposal Nos. 2013G514, 2012G138, 2014G018, 2015G574).

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Figure legends

Fig. 1 Three-dimensional (3-D) reconstructed PXCT images and representative 2-D images of control 1, HY1, and AG1 embryos demonstrating physiological umbilical herniation. The red circle indicates the border between the abdominal coelom and extra-embryonic coelom. The scale bar represents 1 mm. The arrowhead indicates the cecum. The cecum was behind the intestinal loop in HY1. Ad, adrenal gland; CR, colorectum; Du, duodenum; Es, esophagus; In, intestine; Li, liver; Ms, mesonephros; Mt, metanephros; Pa, pancreas; St, stomach; UC, umbilical cord.

Fig. 2 Three-dimensional reconstructed images of the intestinal loop in control 2, HY2, and AG2 embryos. The secondary intestinal loop was confirmed in the extra-embryonic coelom. The red dashed line indicates the border between the abdominal coelom and extra-embryonic coelom. The arrowhead indicates the cecum. The asterisk (*) indicates the end of the intestinal loop in AG2. The scale bar represents 1 mm. CR, colorectum; Du, duodenum; St, stomach.

Fig. 3 Length of the intestinal loop. (A) The blue bar indicates the total digestive tract length (Total length). The red bar indicates the intestinal length within the extra-embryonic coelom (extra length). (B) Ratio of physiological umbilical herniation. The ratio of the intestinal loop length in the extraembryonic coelom (extra length) to the total digestive tract length (Total length).

Fig. 4 Estimated total abdominal cavity volume and intestinal volume. (A) Representative mid-sagittal sectional PXCT images representative of the abdominal cavity in control 1, HY1, and AG1 embryos. The blue line indicates the diaphragm. The red dashed line indicates the border between the abdominal

1 coelom and extra-embryonic coelom. (B) The estimated total abdominal cavity
2 volume was the sum of the liver volume (Blue bar) and intestinal volume (Red
3 bar). The intestinal volume was estimated as described in Materials and
4 Methods. He, heart; In, intestine; Li, liver; Lu, lung; St, stomach; UC, umbilical
5 cord.

6
7
8 Fig. 5 Representative three-dimensional reconstructed images of the stomach,
9 duodenum, and pancreas in control 1, HY1, and AG1 embryos. The circle
10 indicates the cardia. The square indicates the proximal region of the pancreas
11 connected to the duodenum. The arrow indicates the end of the duodenum,
12 where the duodenum is directly attached to the posterior abdominal wall. The
13 scale bar represents 1 mm. Du; duodenum; Pa, pancreas; SMA, superior
14 mesenteric artery; St, stomach.

20 Abbreviations

21 AG = liver agenesis; CS = Carnegie stage; extra length = length of intestinal loop
22 in extraembryonic coelom; HY = liver hypogenesis; PUH = physiological
23 umbilical herniation; PXCT = phase-contrast X-ray computed tomography; Total
24 length = length of total digestive tract

Table 1. Description of liver malformation cases and controls

Case (ID)	CRL (mm)	Intestinal length (mm)			Estimated abdominal vol. (mm ³)			Deviation			Other complications	Related figures
		TDTL	exL	b/a (%)	Total	Liver	Intestine	Stomach	Duodenum	Pancreas		
control 1 (25858)	19	29	20	69	63	40	24	(-)	(-)	(-)	(-)	F1, F4, F5
control 2 (28066)	18	26	17	65	50	34	16	(-)	(-)	(-)	(-)	F2
HY1 (31528)	21	27	23	84	30	7	23	(-)	(-)	(+)	Gastroschisis (suspicious)	F1, F4, F5 *(SG3)
HY2 (22909)	22	33	22	67	36	8	28	(+)	(+)	(+)	More frequent secondary loops Gastroschisis (suspicious)	F2 *(SG5)
AG1 (31874)	19	23	19	82	21	0	21	(+)	(+) ^a	(+)	(-)	F1,F4, F5 *(SG7)
AG2 (28153)	19	21	18	87	18	0	18	(+)	(+) ^a	(+)	Intestinal atresia between the cecum and colorectum. No right mesonephros and genital ridge.	F2 *(SG6)

CRL, crown rump length; TDTL, total digestive tract length; exL, length of extra-embryonic coelom; b/a, Ratio of exL/TDTL; *, indicate the sample ID described in Kanahashi et al., 2016.

^aThe pyloric antrum to the upper part of the duodenum was straight and ran vertically in the right ventral abdomen.

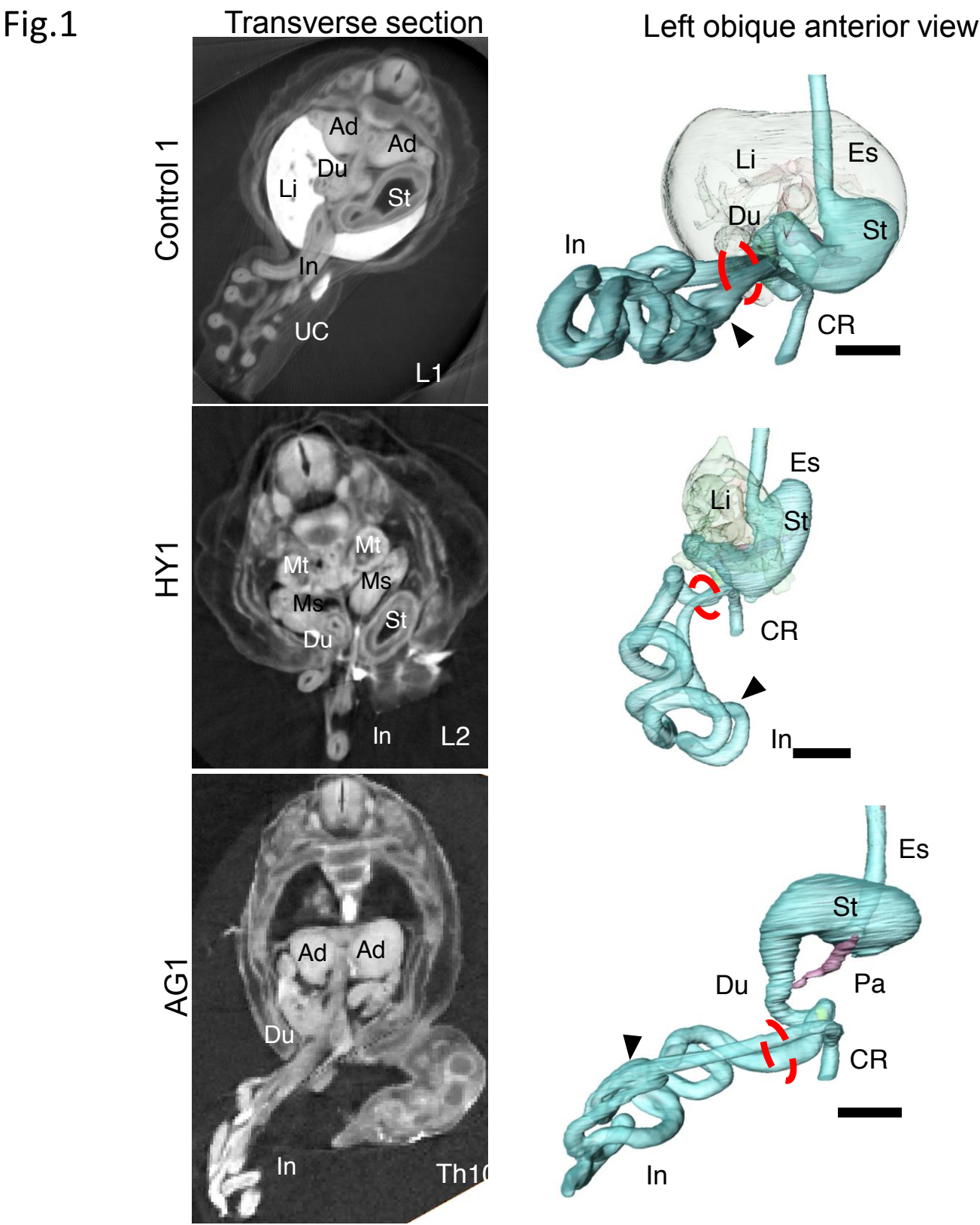


Fig. 1 Three-dimensional (3-D) reconstructed PXCT images and representative 2-D images of control 1, HY1, and AG1 embryos demonstrating physiological umbilical herniation. The red circle indicates the border between the abdominal coelom and extra-embryonic coelom. The scale bar represents 1 mm. The arrowhead indicates the cecum. The cecum was behind the intestinal loop in HY1. Ad, adrenal gland; CR, colorectum; Du, duodenum; Es, esophagus; In, intestine; Li, liver; Ms, mesonephros; Mt, metanephros; Pa, pancreas; St, stomach; UC, umbilical cord.

Fig.2

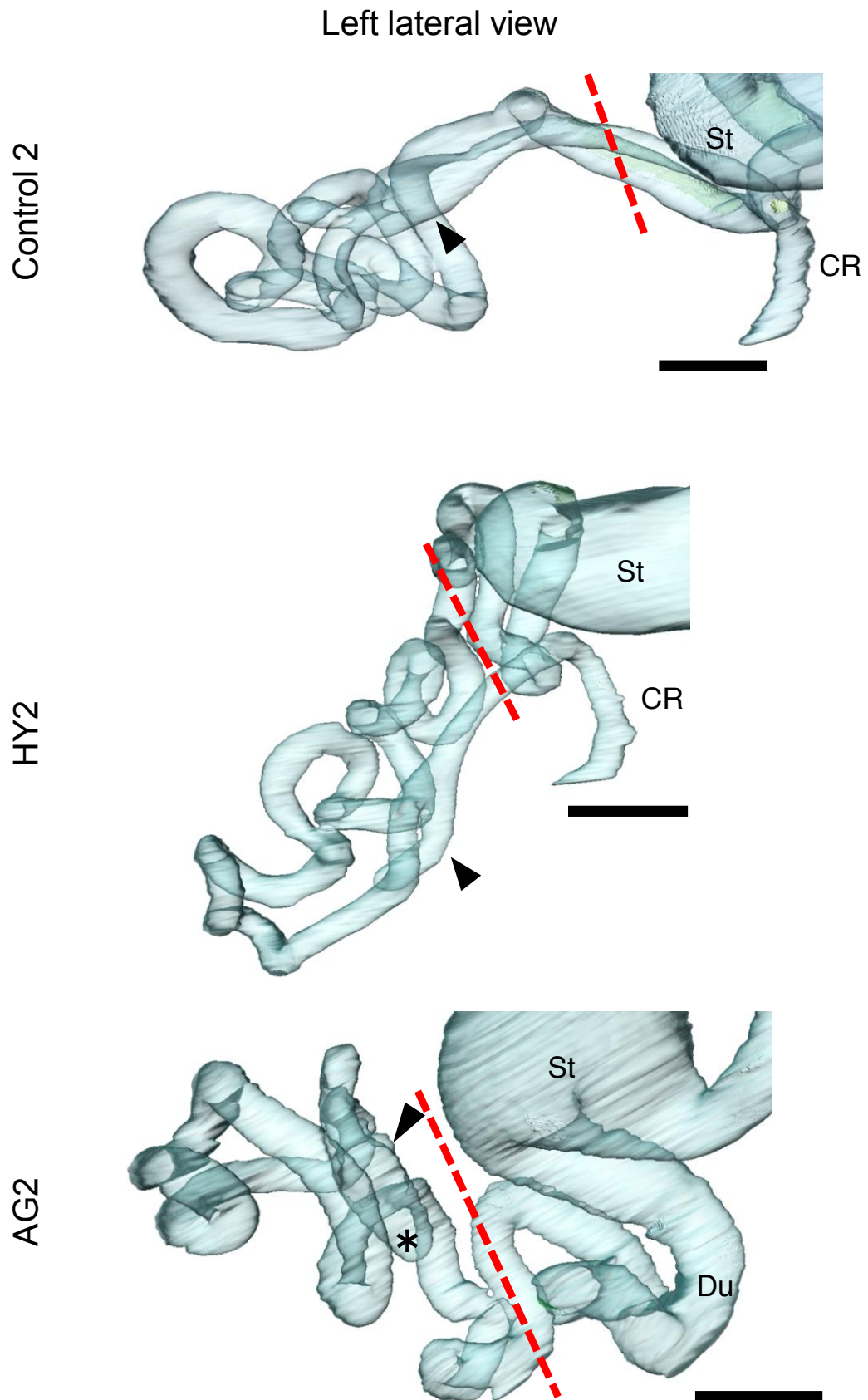


Fig. 2 Three-dimensional reconstructed images of the intestinal loop in control 2, HY2, and AG2 embryos. The secondary intestinal loop was confirmed in the extra-embryonic coelom. The red dashed line indicates the border between the abdominal coelom and extra-embryonic coelom. The arrowhead indicates the cecum. The asterisk (*) indicates the end of the intestinal loop in AG2. The scale bar represents 1 mm. CR, colorectum; Du, duodenum; St, stomach.

Fig.3

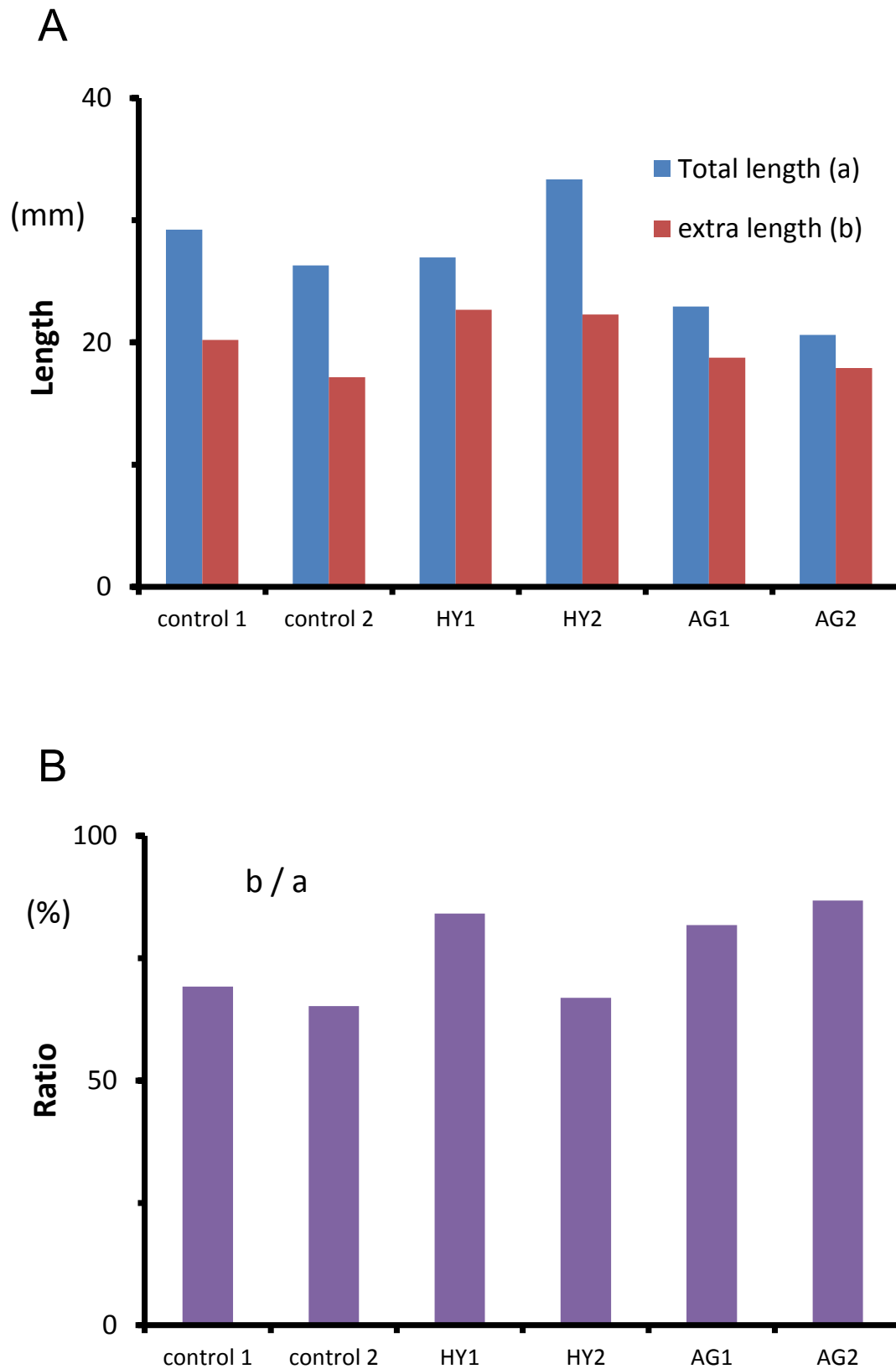
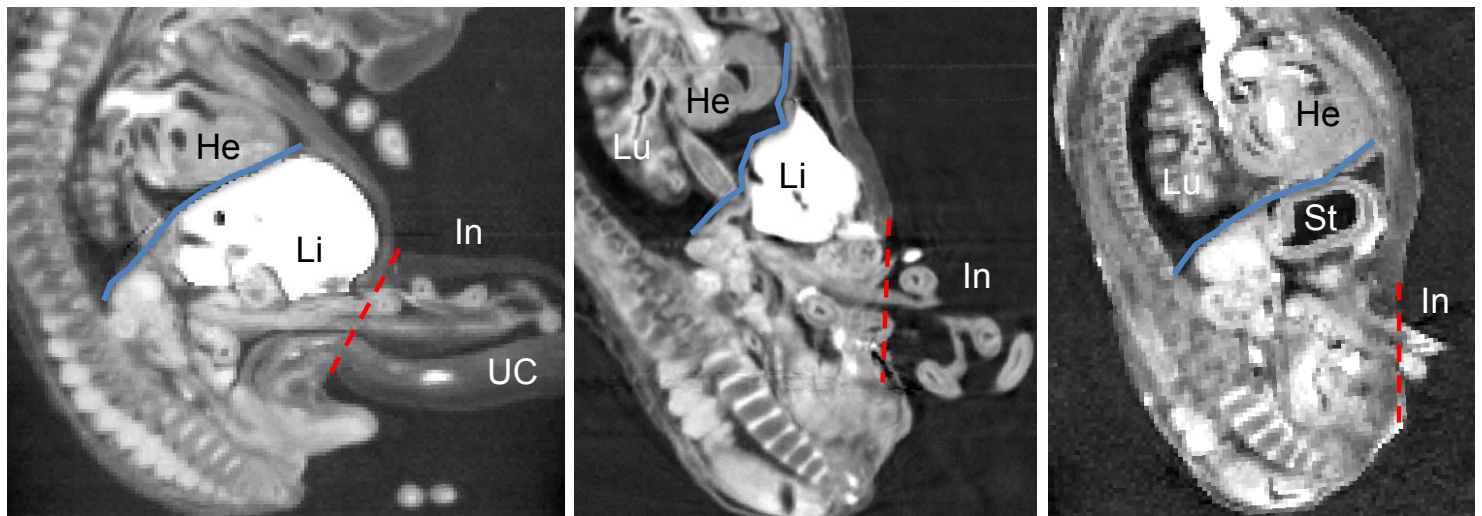


Fig. 3 Length of the intestinal loop. (A) The blue bar indicates the total digestive tract length (Total length). The red bar indicates the intestinal length within the extra-embryonic coelom (extra length). (B) Ratio of physiological umbilical herniation. The ratio of the intestinal loop length in the extraembryonic coelom (extra length) to the total digestive tract length (Total length).

Fig.4

A

Mid-sagittal section



B

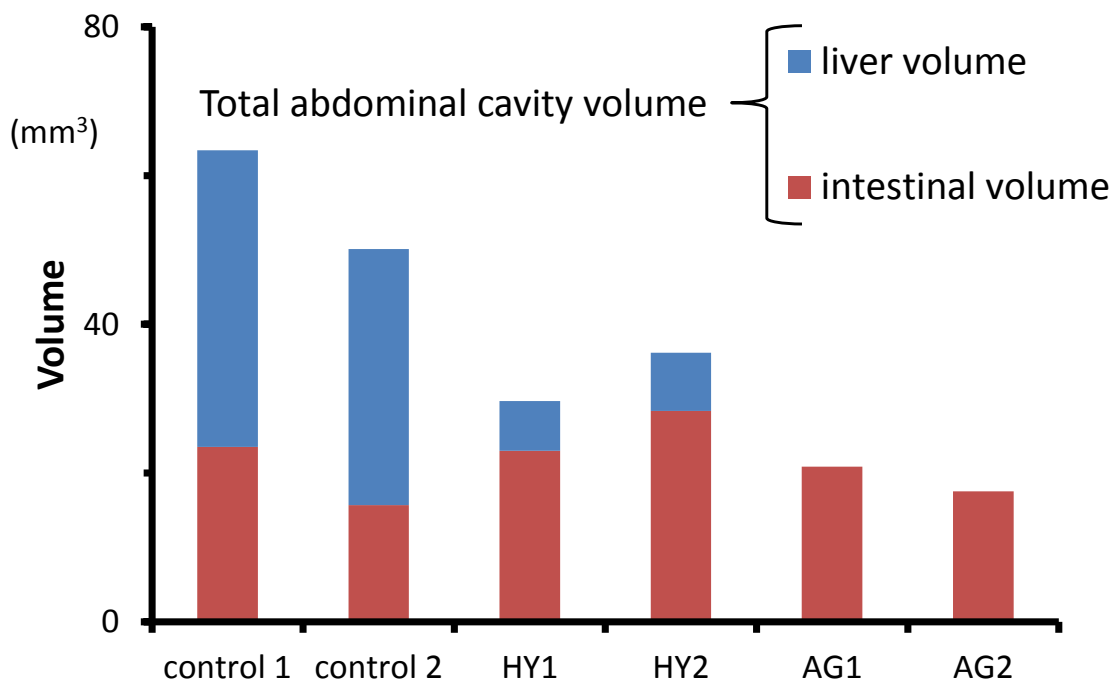


Fig. 4 Estimated total abdominal cavity volume and intestinal volume. (A) Representative mid-sagittal sectional PXCT images representative of the abdominal cavity in control 1, HY1, and AG1 embryos. The blue line indicates the diaphragm. The red dashed line indicates the border between the abdominal coelom and extra-embryonic coelom. (B) The estimated total abdominal cavity volume was the sum of the liver volume (Blue bar) and intestinal volume (Red bar). The intestinal volume was estimated as described in Materials and Methods. He, heart; In, intestine; Li, liver; Lu, lung; St, stomach; UC, umbilical cord.

Fig.5

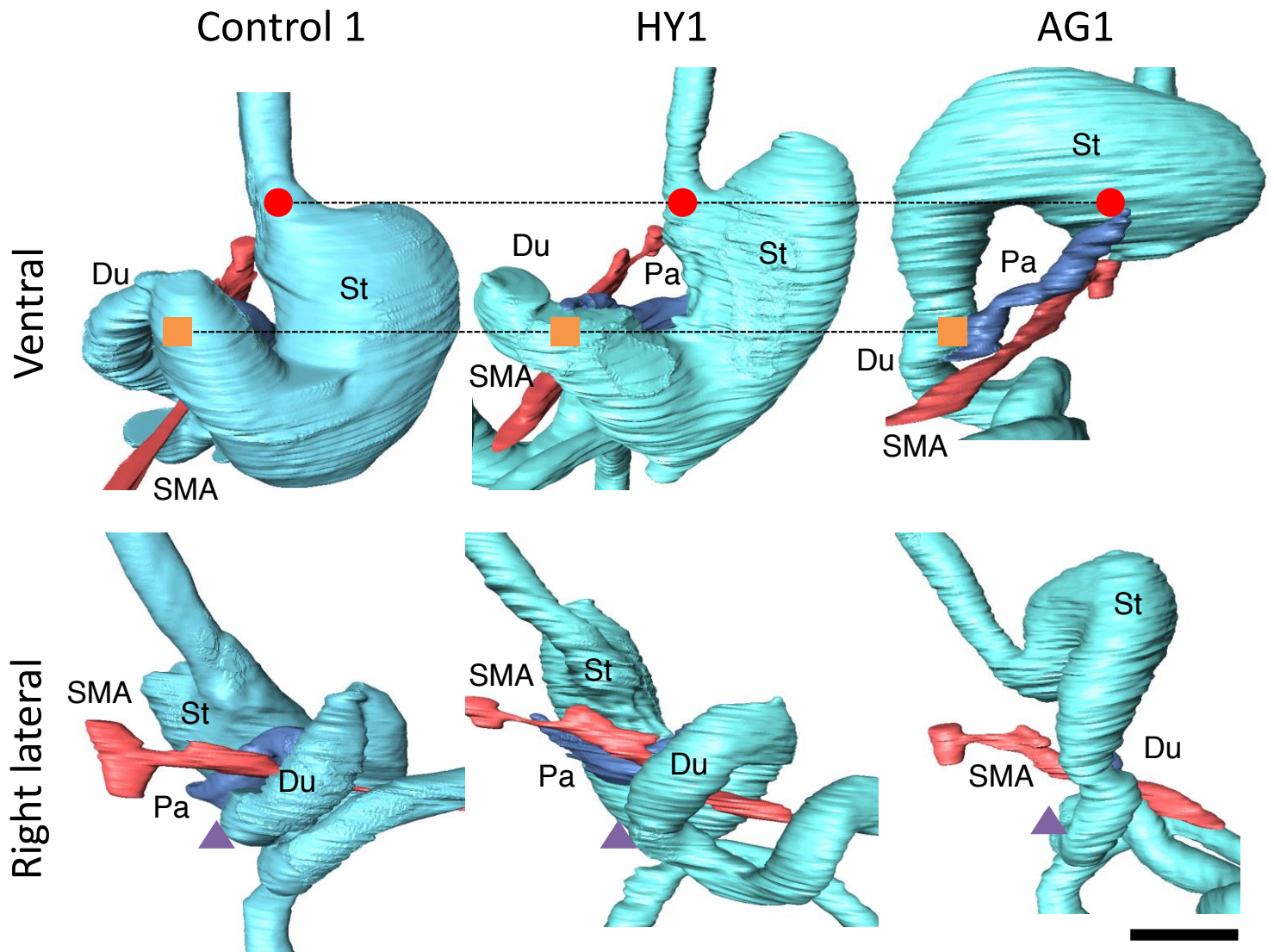


Fig. 5 Representative three-dimensional reconstructed images of the stomach, duodenum, and pancreas in control 1, HY1, and AG1 embryos. The circle indicates the cardia. The square indicates the proximal region of the pancreas connected to the duodenum. The arrow indicates the end of the duodenum, where the duodenum is directly attached to the posterior abdominal wall. The scale bar represents 1 mm. Du; duodenum; Pa, pancreas; SMA, superior mesenteric artery; St, stomach.