Nonrotation of the intestine: embryological and clinical correlation
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INTRODUCTION
The Embryology of the gastrointestinal tract is complex and involves a particularly well-orchestrated rotation of midgut. The complex steps result in assembly of the parts of the gut in the normal anatomical location. Error at the stage of midgut rotation may result in an array of variations in the abdominal organs. The extreme form of malrotation of the midgut is non-rotation and is a very rarely reported variation with the incidence of less than 1:500 (1). The completely altered architecture of the organs in the peritoneal cavity poses a surgical risk and deserves embryological reasoning. The aim of this study is to report an extremely rare anatomical variant. The altered arrangement of viscera is combined peritoneal relations also and hence may lead to complications such as intestinal obstruction, volvulus, and misdiagnosis. For the benefit associated with its clinical implications. This report might aid in the understanding of this anatomical variant and optimize patient management.

CASE REPORT
During routine dissection of the abdomen in a 93-Year old, Caucasian female cadaver, abnormal location of the small intestine was noted. The cadaver was donated and embalmed with formalin based on appropriate ethical principles. A detailed analysis of the intestines was done by tracing from pyloric end of the stomach to the rectum. The vasculature of the abdominal organs was dissected. The liver, pancreas, spleen and the genitourinary tract were dissected to note any variation. The Stomach was normal in a location with the pyloric end just to the right of the midline. The duodenum was oriented vertically, coiled, lacked its usual ‘C’ loop and to the right of the vertebral column. The entire small intestine was found to lie on the right side of the abdominal cavity. The duodenum was located to the right of superior mesenteric artery. The ileocecal junction was traced to the hypogastrium. The colon formed a loop on the left half, and sigmoid colon was typically located. A fibrous Ladd’s band extended from posterior abdominal wall to the colon, crossing anterior to the duodenum. The clinical presentation of nonrotation ranges from vague intermittent pain to acute bowel obstruction. Adequate knowledge of Embryology and resultant variations aid in understanding the abnormal findings during diagnostic interventions and prevents surgical complications.

Key Words: Malrotation; Nonrotation; Embryology; Midgut; Ladd’s band

DISCUSSION
Early in embryonic life, before approximately 4 weeks of gestation, the to the right of superior mesenteric artery. The ileocecal junction was traced to the hypogastrium. The colon formed a loop on the left half, and sigmoid colon was typically located. A fibrous Ladd’s band extended from posterior abdominal wall to the colon, crossing anterior to the duodenum. The clinical presentation of nonrotation ranges from vague intermittent pain to acute bowel obstruction. Adequate knowledge of Embryology and resultant variations aid in understanding the abnormal findings during diagnostic interventions and prevents surgical complications.

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Figure 2) Reversed relation of superior mesenteric vessels. 1. Pancreas; 2. Abdominal aorta; 3. Superior mesenteric vein; 4. Superior mesenteric artery

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Snyder WH, Chaffin L. Embryology and surgical aspects of the umbilical cord to continue its growth. During this time, the duodenum rotates counterclockwise another 90°, while the colon undergoes no rotation. At ten weeks, the embryonic bowel returns to the abdominal cavity, and both the duodenum and the colon rotate again. At this time, the duodenum completes its final 90° counterclockwise rotation, until the duodeno-jejunal junction is located to the left of the spine, while the colon rotates counterclockwise 180° until the cecum is located in the right lower quadrant of the abdomen. The completed normal rotation of the bowel produces a long mesenteric attachment for the small bowel (4).

A defect in stage II of midgut rotation had resulted in the presentation described in this case report. Stage I defect results in omphalocele. Stage II defect leads to malrotation, nonrotation or reversed rotation. Stage III defect results in the unattached duodenum, mobile cecum, unattached mesentery (5). Malrotation tends to occur when the umbilical ring is wider, and the normal sequence of the return of the prearterial and postarterial segment is lost (6). The following anatomical variations may be associated with malrotation namely, Absence of kidney and ureter, Biliary atresia, Congenital diaphragmatic hernia, Duodenal or small bowel stenosis or atresia, Duodenal web, Gastrochisis, Hirschsprung disease, Imperforate anus, Meckel diverticulum, Omphalocele (7). Some of the syndromes associated with malrotation include Apple-peel intestinal atresia, Cornelia de Lange syndrome, Cantrell syndrome, Cat-eye syndrome, Chromosomal abnormalities (trisomies 13, 18, and 21) (8). The presented case did not have evidence of these abnormalities.

Other rare types of rotation include a. Reverse rotation of the duodenojejunal limb resulting in a duodenum that rests anterior to the superior mesenteric artery. b. Reverse rotation of the cecocolic limb resulting in a transverse colon that is posterior to the superior mesenteric artery. c. Reverse rotation of the duodenojejunal limb with the normal rotation of the cecocolic limb resulting in a para duodenal hernia. In reverse rotation, the duodenum is located anterior to the superior mesenteric artery. Anterior to the duodenum, the cecocolic limb rotates normally, and the mesentery of the right colon creates a pouch into which the small bowel can herniate (9).

Although the clinical history of the cadaver was not available, her age at death and absence of a large scar on the anterior abdominal wall signify that she may not have had any major surgical complication. The classic clinical manifestation of malrotation in newborns is biliocolic obstruction with or without abdominal distention associated with either duodenal or midgut volvulus. A delay in diagnosis and treatment may result in small bowel necrosis, short gut syndrome, and dependence on total parenteral nutrition (1). The clinical presentation associated with nonrotation of gut ranges from vague intermittent pain to symptoms of acute bowel obstruction, although it may be asymptomatic too. The abnormally located appendix may lead to misdiagnosis of appendicitis. The peritoneal bands may lead to intestinal compression or volvulus. Nonrotation is not as dangerous for the patient as malrotation because, in general, the base of the mesentery is wider than in malrotation, and the risk of volvulus is less compared to that in malrotation. However, nonrotation can be a difficult diagnosis radiologically, symptomatic patients may warrant laparoscopic or open exploration to confirm the diagnosis. Asymptomatic patients with radiologic findings suggesting nonrotation can be observed (9).

CONCLUSION

The altered anatomical location of the abdominal organs in nonrotation of intestine may not always result in clinical presentation. However, misunderstanding of incidental clinical or diagnostic findings may compromise the health of the patient. Adequate knowledge of Embryology and such resultant variations aid in understanding the abnormal findings during diagnostic interventions and also prevent surgical complications.

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REFERENCES


