New successful one-step surgical repair for apple peel atresia

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Abstract: A new successful surgical approach in two identical twins delivered with intestinal atresia, “apple peel” type is reported. This technique consists of: (1) an end-to-end oblique primary anastomosis with single layer inverted 5/0 Vicryl® sutures (Ethicon, Inc, Somerville, NJ); (2) proximal tube jejunostomy using Foley catheter, size 10 French, inserted through a stab wound in the left upper quadrant and entering the proximal dilated loop at about 10 cm proximal from its blind end (site of anastomosis); and (3) transanastomotic stenting using feeding tube, size 6 French, exteriorized in conjunction with the Foley catheter and reaching the lumen of the distal loop for more than 20 cm. The postoperative course was uneventful and progressive oral feed became ad libitum around day 40 postoperative. This procedure is simple, performed in one stage, and responds to the most obligatory requirements of this congenital malformation.

Keywords: proximal jejunal atresia, intestinal atresia, surgical repair, anastomosis

Introduction
Intestinal atresia is a congenital malformation causing neonatal intestinal obstruction. “Apple peel” type is rare but still presents serious surgical repair challenges with high morbidity and mortality. A new successful surgical approach is presented, with the hope that it will be adopted as a first-line therapeutic approach for apple peel atresia.

Case reports
Two identical twins were delivered at full-term via cesarean section, following an uneventful pregnancy to a healthy 26-year-old gravida 1 para 1 mother. There was no consanguinity and their birth weights were 1460 g and 1995 g, respectively.

Diagnosis of neonatal intestinal obstruction was made prenatally by ultrasound at the end of the seventh month. The ultrasound showed an intraabdominal structural abnormality, suspected to be an abnormal dilated intestinal loop. No other abnormality, except for a very mild polyhydramnios, was noticed at the time of the ultrasound, nor during cesarean section. In both of the twins, postnatal assessment revealed the presence of mild systolic murmur, related to a patent ductus arteriosus.

After birth, there was a significant amount of bile-stained gastric aspirate; plain abdominal X-ray confirmed the presence of high intestinal obstruction with a short proximal dilated loop (Figure 1). After postnatal evaluation, informed parental consent was obtained and the babies were operated on during the first 36 hours.

Exploration confirmed a jejunal atresia about 15 cm from the ligament of Treitz, with complete separation of blind-end loops and extensive mesenteric defect giving
the typical appearance of apple peel atresia. The distal small intestine was short (less than 40 cm), with coils around the vascular axis. The disparity in lumen diameter between the proximal and distal tiny loops was about 15 folds (Figure 2).

The operation consisted of (Figure 3):
1. An end-to-end oblique anastomosis with a single layer inverted 5/0 Vicryl® sutures (Ethicon, Inc, Somerville, NJ).
2. Proximal tube jejunostomy constructed using Foley catheter (Foley catheter latex-free mannsield MA02048; Tyco/Healthcare Kendall, Mexico City, Mexico), size French 10, inserted through a short stab wound in the left upper quadrant and in the proximal dilated loop about 10 cm proximal to the anastomosis.
3. Transanastomotic stent using a feeding tube (Feeding Tube, Nontoxic Pyrogen Free; Kawamoto Corporation, Tokyo, Japan), size French 6, exteriorized in conjunction with the Foley catheter. The Foley catheter and stent were inserted through a jejunal stab wound, closed by 3/0 Vicryl purse-string suture, the balloon was inflated, and the intestine was fixed to the inner abdominal wall by three opposite 3/0 Vicryl sutures. Finally, the Foley catheter and transanastomotic tube were fixed to the skin by a nonabsorbable suture to prevent their dislodgment.

Postoperative course was uneventful. Initially, Foley catheter drained about 80–100 mL of bile-stained fluid per day; this drainage progressively decreased to 0 mL by day 15. Abdominal X-ray showed the presence of air in the distal small intestine on day 5 postoperative (Figure 4). Oral feeding started on day 15 with 2 mL 5% dextrose half-normal saline every 8 hours, which was increased gradually and then changed to half strength and later to full strength milk.
Intestinal atresia is the most common cause of neonatal intestinal obstruction, occurring at a rate of 2.9 per 10,000 births (including stillbirths). Sex distribution is almost equal. Apple peel atresia, which represents <10% of all types of intestinal atresia, is defined as complete separation of blind ends with an extensive mesenteric defect where the distal small intestine coils around the axis of blood supply, acquiring its blood supply from a single ileocecal artery. In these types of congenital anomalies the overall bowel length is shorter than normal and usually accompanies extreme prematurity. Hereditary predisposition is also reported.

Apple peel atresia poses challenging problems particularly for reconstruction. The most recommended surgical repair for apple peel atresia consists of a proximal jejunostomy (usually with tapering or plication of the dilated segment to enhance its return to a normal caliber and active peristalsis) and ileostomy (Mikulicz exteriorization) or distal tube ileostomy at the level of distal blind-end loop.2–4 The latter is used later on for very progressive fluid administration before scheduling a definitive anastomosis. In this condition, definitive anastomosis is usually not possible before the age of 3–6 months. This method leads to high mortality due to the major complications of proximal jejunostomy in the neonatal period and the complications of long-term total parenteral alimentation.5 Previously, this method was applied without any incident in several cases of simple jejunal atresia and in many referred multioperated cases for different causes of proximal intestinal obstruction.

Primary anastomosis is protected by proximal stomy and disruption should not occur if anastomosis was correctly done. Decompression of the proximal loop is maintained as long as needed by the Foley catheter without any related complication; it does not interrupt the intestinal transit even when it is kept open permanently. Ten days after insertion, the Foley catheter can easily be replaced in case it falls out (like gastrostomy tube). The Foley catheter should not be removed before the proximal loop returns to its adequate caliber (no more dilatation on X-ray). Plication or tapering the proximal dilated loop is unnecessary as well as partial resection (except when dilated end-loop is in a cystic shape). Foley catheter can be opened to relieve any episodes of vomiting or abdominal distention.

The transanastomotic tube (stent) enables the content of the proximal intestinal loop to reach the distal loop progressively and does not prevent stricture at the site of the anastomosis. The stent can be removed a few days after the intestinal air shadow reaches the rectum and oral feeding can start to be tolerated. This small-sized transanastomotic tube should be inserted in conjunction with the Foley catheter – rather than nasogastric – for better tolerance, and can be left in place for months without any related complication.

The most surprising issue in these case reports was that in spite of this typical aspect of apple peel atresia, and the complete disconnection from sympathetic and parasympathetic...
innervations, full early oral feeding was tolerated without any incident, except for mild chronic diarrhea with steatorrhea. There was no indication for intestinal contrast examination since the patients did not show any intestinal transit difficulties.

This method is recommended as first-line therapeutic approach for apple peel atresia as well as for different types of proximal jejunal atresia (with proximal dilated loop).

Disclosure
The author reports no conflicts of interest in this work.

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