

PEDIATRIC SURGERY Update © Vol. 30 No. 02 FEBRUARY 2008

Mesh Repair Congenital Diaphragmatic Hernia

Congenital diaphragmatic hernia continues to carry a high mortality associated with the presence of pulmonary hypoplasia. Repair of the diaphragmatic defect is usually carried on when the child obtains sufficient hemodynamic stability to tolerate a major surgical procedure. Surgical repair or closure of the defect is either carried out primarily or in case that most of the hemidiaphragm is lacking, using a piece of prosthetic mesh. Primary repair is performed when there is sufficient diaphragm to approximate with low tension, carries a low rate of recurrence and avoids the mechanical and infectious complications associated with implanted prostheses. When the size of the defect can be known preoperatively a split abdominal wall muscle flap through a low abdominal incision can be planned. Mesh repair utilizes several prosthetic materials such as polytetrafluoroethylene (Gore-Tex), polypropylene (Marlex), lyophilized dura, Surgisis or even small intestinal submucosa. Composite patch repair using Gore-Tex/Marlex has also been reported. Overall, recurrence rates after mesh repair is significant greater (almost 50%) than after primary repair. Polypropylene mesh in contact with small bowel carries a high risk of fistula formation reason why Gore-Tex is preferred. Vycryl mesh is not a suitable material for repairing these defects.

References:

1- Saltzman DA, Ennis JS, Mehall JR, Jackson RJ, Smith SD, Wagner CW: Recurrent congenital diaphragmatic hernia: A novel repair. J Pediatr Surg. 36(12):1768-9, 2001

2- Sydorak RM, Hoffman W, Lee H, Yingling CD, Longaker M, Chang J, Smith B, Harrison MR, Albanese CT: Reversed latissimus dorsi muscle flap for repair of recurrent congenital diaphragmatic hernia. J Pediatr Surg. 38(3):296-300, 2003

3- Scaife ER, Johnson DG, Meyers RL, Johnson SM, Matlak ME: The split abdominal wall muscle flap--a simple, mesh-free approach to repair large diaphragmatic hernia. J Pediatr Surg. 38(12):1748-51, 2003

4- Smith MJ, Paran TS, Quinn F, Corbally MT: The SIS extracellular matrix scaffold-preliminary results of use in congenital diaphragmatic hernia (CDH) repair. Pediatr Surg Int. 20(11-12):859-62, 2004

5- Grethel EJ, Cortes RA, Wagner AJ, Clifton MS, Lee H, Farmer DL, Harrison MR, Keller RL, Nobuhara KK: Prosthetic patches for congenital diaphragmatic hernia repair: Surgisis vs Gore-Tex. J Pediatr Surg. 41(1):29-33, 2006

6- Riehle KJ, Magnuson DK, Waldhausen JHT: Low recurrence rate after Gore-Tex/Marlex composite patch repair for posterolateral congenital diaphragmatic hernia. J Pediatr Surg 42(11): 1841-1844, 2007

Solitary Intestinal Fibromatosis

Congenital solitary intestinal fibromatosis (SIF) is a very rare tumor that occurs in the newborn and infant period. It belongs to the group of pediatric fibromatosis. Solitary Intestinal fibromatosis involves the intestinal wall, can produce bowel obstruction or perforation and usually involves most frequently the jejunum and ileum. SIF presents most

commonly as a solitary or less commonly as multiple lesions usually confined to soft tissue and bone. Lesions in the duodenum can cause gastric outlet obstruction indistinguishable from pyloric stenosis. Most cases consist of solitary tumors affecting the small or large bowel. Histologic examination in each case shows a transmural infiltrative spindle cell lesion having the morphologic features of fibromatosis. The spindle cells have no atypia, stain positively for Vimentin and CD34 while negative for muscle cell markers. Ultrastructural studies reveals the tumor to be composed of myofibroblasts. The tumor most often presents as a polypoid mass protruding into the intestinal lumen causing obstruction. Symptoms are usually bilious vomiting, abdominal distension, malabsorption and obstipation. Management consists of wide local resection of the tumor along with the segment of affected bowel. Some cases have demonstrated spontaneous regression. Solitary lesions carry an excellent prognosis after resection, while multiple lesions carry a worse prognosis.

References:

1- Canioni D, Fekete C, Nezelof C: Solitary intestinal fibromatosis: a rare cause of neonatal obstruction. Pediatr Pathol. 9(6):719-24, 1989

2- Srigley JR, Mancer K: Solitary intestinal fibromatosis with perinatal bowel obstruction. Pediatr Pathol. 2(3):249-58, 1984

3- Türken A, Senocak ME, Kotiloglu E, Kale G, Hiçsönmez A: Solitary intestinal fibromatosis mimicking malabsorption syndromes. J Pediatr Surg. 30(9):1387-9, 1995

3- Arets HG, Blanco C, Thunnissen FB, Heineman E: Solitary intestinal fibromatosis as a cause of bile vomiting in a neonate. J Pediatr Surg. 35(4):643-5, 2000

4- Numanoglu A, Davies J, Millar AJ, Rode H: Congenital solitary intestinal fibromatosis. Eur J Pediatr Surg. 12(5):337-40, 2002

5- Coulon A, McHeik J, Milin S, Levard G, Levillain P, Fromont G: Solitary intestinal fibromatosis associated with congenital ileal atresia. J Pediatr Surg. 42(11):1942-5, 2007

Natural Orifice Transluminal Endoscopic Surgery

Natural orifice transluminal endoscopic surgery (NOTES) is the next frontier in minimal invasive surgical procedures. NOTES takes advantage of doing a laparoscopic procedure through natural orifices of our body such as mouth (trans-gastric), anus (trans-colonic) or vagina totally eliminating a scar in the abdomen. Theoretically, this approach could reduce postoperative abdominal wall pain, wound infection, hernia formation, and adhesions. So far studies in animal models have demonstrated the feasibility of performing such procedure to remove the gallbladder through either per-oral transgastric or per-anal transcolonic by perforating such viscera and introducing a multichannel locking endoscope to introduce the laparoscopic instruments utilized during actual transperitoneal procedures. The incision done in the perforated viscera is subsequently closed with endoscopic clips, endoloops, or a prototype closure device. In contrast to the transgastric method, a transcolonic approach provides more consistent identification of structures in the upper abdomen and provides better en face orientation and scope stability. There is no bleeding or laceration of adjacent organs. Animal models have been used to demonstrate the possible applications of NOTES, including transgastric peritoneoscopy, tubal ligation, gastrojejunostomy, partial hysterectomy, oophorectomy, nephrectomy and transcolonic

exploration, liver biopsy, distal pancreatectomy and cholecystectomy. The first human report in 2007 was a successful trans-vaginal cholecystectomy.

References:

1- Ko CW, Kalloo AN: Per-oral transgastric abdominal surgery. Chin J Dig Dis. 2006;7(2):67-70

2- Pai RD, Fong DG, Bundga ME, Odze RD, Rattner DW, Thompson CC: Transcolonic endoscopic cholecystectomy: a NOTES survival study in a porcine model (with video). Gastrointest Endosc. 64(3):428-34, 2006

3- Fong DG, Pai RD, Thompson CC: Transcolonic endoscopic abdominal exploration: a NOTES survival study in a porcine model. Gastrointest Endosc. 65(2):312-8, 2007

4- Wagh MS, Thompson CC: Surgery insight: natural orifice transluminal endoscopic surgery--an analysis of work to date. Nat Clin Pract Gastroenterol Hepatol. 4(7):386-92, 2007

5- Marescaux J, Dallemagne B, Perretta S, Wattiez A, Mutter D, Coumaros D: Surgery without scars: report of transluminal cholecystectomy in a human being. Arch Surg. 142(9):823-6, 2007

6- Fritscher-Ravens A, Ghanbari A, Thompson S, Patel K, Kahle E, Fritscher T, Niemann H, Koehler P, Milla P: Which parameters might predict complications after natural orifice endoluminal

surgery (NOTES)? Results from a randomized comparison with open surgical access in pigs. Endoscopy. 39(10):888-92, 2007

* Edited by: Humberto Lugo-Vicente, MD, FACS, FAAP

Professor /Academic Director of Pediatric Surgery, University of Puerto Rico - School of Medicine, Rio Piedras, Puerto Rico.

Address: P.O. Box 10426, Caparra Heights Station, San Juan, Puerto Rico USA 00922-0426. Tel (787)-786-3495 Fax (787)-720-6103 E-mail: *titolugo@coqui.net* Internet: http://home.coqui.net/titolugo

> © PSU 1993-2007 ISSN 1089-7739