

PEDIATRIC SURGERY Update © Vol. 36 No. 02 FEBRUARY 2011

Ethanol Sclerotherapy for Lymphangiomas

Lymphatic malformations are very difficult lesions to manage and eradicate in children. They constitute 6% of all benign lesions in infants and children. Head and neck are the more commonly affected regions. Most lesions will appear within the first two years of life of the child. The cystic lymphatic lesions are classified as macrocystic (> 1 cm), microcystic (< 1 cm) and mixed variety. Lymphatic malformations do not have a familial tendency and they do not become malignant with time. Surgical excision is the standard method of choice to manage lymphatic lesions that are localized and do not involve vital surrounding structures. Mutilating surgery is avoided at all costs. Diagnosis of the anatomic arrangements that these lymphatic malformations have with vital structure is obtained with the help of CT-Scan and MRI imaging. For unresectable or residual lymphatic cystic lesions the use of sclerotherapy has been advocated. Sclerosing agents include OK-432, Bleomycin, Fibrin glue, Doxycycline, 50% dextrose, Ethibloc (alcoholic solution of corn protein) and recently 98% Ethanol. Some agents are very difficult to obtained due to FDA regulation. Others cause toxic, allergic and irritant effects to the child. Sterile 98% ethanol produces minimal side effect, is easy to purchase and has a low cost. Ethanol sclerotherapy can be affected using CT-guided technology with the child under sedation or general anesthesia. Th volume of ethanol used for sclerotherapy should be no more than 1.0 ml/kg of weight.

References:

- 1- Saba C, Bossi MC, Barletta A: Therapy of subcutaneous cystic lymphangioma with ultrasound-guided puncture and alcoholization. Radiol Med. 83(3):270-2, 1992
- 2- Puig S, Aref H, Brunelle: Double-needle sclerotherapy of lymphangiomas and venous angiomas in children: a simple technique to prevent complications. AJR Am J Roentgenol. 180(5):1399-401, 2003
- 3- Alomari AI, Karian VE, Lord DJ, Padua HM, Burrows PE: Percutaneous sclerotherapy for lymphatic malformations: a retrospective analysis of patient-evaluated improvement. J Vasc Interv Radiol. 17(10):1639-48, 2006
- 4- Shiels WE 2nd, Kenney BD, Caniano DA, Besner GE: Definitive percutaneous treatment of lymphatic malformations of the trunk and extremities. J Pediatr Surg. 43(1):136-9, 2008
- 5- Shiels WE 2nd, Kang DR, Murakami JW, Hogan MJ, Wiet GJ: Percutaneous treatment of lymphatic malformations. Otolaryngol Head Neck Surg. 141(2):219-24, 2009
- 6- Impellizzeri P, Romeo C, Astra Borrut F, et al: Sclerotherapy for cervical cystic lymphatic malformations in children. Our experience with computed tomography-guided 98% sterile ethanol insertion and review of the literature. J Pediatr Surg. 45(12): 2473-2478, 2010

Ogilvie Syndrome

Acute colonic pseudo-obstruction also known as Ogilvie's syndrome is a massive colonic dilatation associated with signs and symptoms of colonic obstruction without an

evident mechanical cause. Ogive syndrome is observed predominantly in the elderly population with few cases reported in children. Predisposing factors for Ogilvie's syndrome in children includes postoperative state, trauma, infections, Sickle cell disease, cardiac diseases and chemotherapy for malignancy. Symptoms include constipation, abdominal pain, nausea, vomiting and abdominal distension. Diagnosis is suggested in flat simple abdominal films. Findings on CT-Scan are diagnostic showing massive colonic dilatation with diameters of eight to 12 cm and without evidence of overt mechanical obstruction. If left untreated, this dilatation can lead to colonic perforation and peritonitis in 10% of children with high mortality rates. Initial management consists of nasogastric decompression, bowel rest, hydration, electrolyte correction, along with discontinuation of drugs affecting bowel motility. If symptoms fail to improve with initial management then rectal tube or colonoscopy decompression is utilized. Neostigmine, an acethycholinesterase inhibitor which increases parasympathetic tone, has been found to be very effective in managing patients with Ogilvie syndrome. Neostigmine is slowly titrated in increments up to a total of 0.05mg/kg of weight. Also, oral erythromycin therapy has been used to manage this condition. Surgery will be needed if the child develops perforation or signs of bowel ischemia.

References:

- 1- Hyman PE: Chronic intestinal pseudo-obstruction in childhood: progress in diagnosis and treatment. Scand J Gastroenterol Suppl. 213:39-46, 1995
- 2- Gmora S, Poenaru D, Tsai E: Neostigmine for the treatment of pediatric acute colonic pseudo-obstruction. J Pediatr Surg. 37(10):E28, 2002
- 3- Jiang DP, Li ZZ, Guan SY, Zhang YB: Treatment of pediatric Ogilvie's syndrome with low-dose erythromycin: a case report. World J Gastroenterol. 13(13):2002-3, 2007
- 4- Kim TS, Lee JW, Kim MJ, Park YS, Lee DH, Chung NG, Cho B, Lee S, Kim HK: Acute colonic pseudo-obstruction in postchemotherapy complication of brain tumor treated with neostigmine. J Pediatr Hematol Oncol. 29(6):420-2, 2007
- 5- Khosla A, Ponsky TA: Acute colonic pseudoobstruction in a child with sickle cell disease treated with neostigmine. J Pediatr Surg. 43(12):2281-4, 2008
- 6- Lee JW, Bang KW, Jang PS, Chung NG, Cho B, Jeong DC, Kim HK, Im SA, Lim GY: Neostigmine for the treatment of acute colonic pseudo-obstruction (ACPO) in pediatric hematologic malignancies. Korean J Hematol. 45(1):62-5, 2010

Colostomy Closure

Colostomy closure is a common and important surgical procedure performed in children which carries a significant risk of morbidity and mortality. Some complication associated with closure of a colostomy includes wound infection, anastomotic dehiscence, bleeding, anastomotic stricture, incisional hernia and death. The periostoma lymphatics are colonized with bacteria, reason why surgery site infection rises during this procedure. For colostomy closure most children should be admitted the day before surgery for mechanical cleansing of the proximal and distal bowel. Systemic antibiotics during anesthesia induction are necessary. Infection rate is not affected by the use of oral antibiotics. Meticulous surgical technique including packing of proximal stoma, use of plastic drapes for surgical field immobilization, correct dissection, careful hemostasis avoiding contamination and performing an anastomosis in well vascularized limbs are

essential to reduce complications. Peritoneal irrigation, fascial closure in layers, adequate hemostasis and avoidance of dead spaces are also essential issues to watch for. Postoperative nasogastric tubes are not necessary. Early feeding is encouraged.

References:

- 1- Weber TR, Tracy TF Jr, Silen ML, Powell MA: Enterostomy and its closure in newborns. Arch Surg. 130(5):534-7, 1995
- 2- Sangkhathat S, Patrapinyokul S, Tadyathikom K: Early enteral feeding after closure of colostomy in pediatric patients. J Pediatr Surg. 38(10):1516-9, 2003
- 3- Chandramouli B, Srinivasan K, Jagdish S, Ananthakrishnan N: Morbidity and mortality of colostomy and its closure in children. J Pediatr Surg. 39(4):596-9, 2004
- 4- Breckler FD, Rescorla FJ, Billmire DF: Wound infection after colostomy closure for imperforate anus in children: utility of preoperative oral antibiotics. J Pediatr Surg. 45(7):1509-13, 2010
- 5- Bischoff A, Levitt MA, Lawal TA, Pena A: Colostomy closure: how to avoid complications. Pediatr Surg Int. 26(11):1087-92, 2010

* 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. Chief – Pediatric Surgery, San Jorge Childrens Hospital.

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 1993-2011 ISSN 1089-7739