

# PEDIATRIC SURGERY Update © Vol. 22 No. 01 JANUARY 2004

## Hashimoto Thyroiditis

Hashimoto thyroiditis (HT) is a chronic lymphocytic autoimmune thyroiditis seen with some frequency in adolescent females and children. Most common cause of asymptomatic enlargement of the thyroid gland in children in iodine-sufficient geographic regions. Thyroid cell damage in HT is caused by antithyroid antibody-dependent cell-mediated direct toxicity linked to deficiency in antigen-specific suppressor T lymphocytes. The gland shows lymphocyte infiltration with follicular cell hyperplasia. Thyroid antibodies are elevated. Radionuclear scans show absent uptake. Initially the child develops elevated thyroid hormones (T3 and T4) followed by symptomatic hypothyroidism. Following the hypothyroid phase there is final recovery in most patients. Indications for surgery in HT include: 1- firm enlargement of the gland causing tracheal compression with dyspnea, hoarseness or swallowing difficulties, 2- failure to respond to suppressive therapy and development of symptomatic hyperthyroid goiter, and 3- development and enlargement of a solitary thyroid nodule. The incidence of malignancy in HT is low. Differentiating a hyperplastic follicular cell nodule from a follicular neoplasm is very difficult using fine needle aspiration biopsy. Patient with malignant nodules in Hashimoto glands are most commonly papillary, females, low frequency of extrathyroidal invasion and nodal metastasis with absent distal metastasis. It is believed the lymphocytic infiltration of HT causes a form of immune reaction to control tumor growth and proliferation.

### **References:**

1- Okayasu I, Fujiwara M, Hara Y, Tanaka Y, Rose NR: Association of chronic lymphocytic thyroiditis and thyroid papillary carcinoma. A study of surgical cases among Japanese, and white and African Americans. Cancer 76(11):2312-8, 1995

2- Webb AJ, Brewster S, Newington D: Problems in diagnosis and management of goitre in childhood and adolescence. Br J Surg 83(11):1586-90, 1996

3- Nguyen GK, Ginsberg J, Crockford PM, Villanueva RR: Hashimoto's thyroiditis: cytodiagnostic accuracy and pitfalls. Diagn Cytopathol 16(6):531-6, 1997

4- Loh KC, Greenspan FS, Dong F, Miller TR, Yeo PP: Influence of lymphocytic thyroiditis on the prognostic outcome of patients with papillary thyroid carcinoma. J Clin Endocrinol Metab 84(2):458-63, 1999

5- Hopwood NJ, Kelch RP: Thyroid masses: approach to diagnosis and management in childhood and adolescence. Pediatr Rev 14(12):481-7, 1993

6- Lafranchi S: Thyroiditis and acquired hypothyroidism. Pediatr Ann 21(1):29, 32-9, 1992

7- Strakosch CR: Thyroiditis. Aust N Z J Med 16(1):91-100, 1986

## **Bile Reflux Gastritis**

Alkaline reflux gastritis develops in patients with previous operations that destroy the integrity of the pylorus as a true sphincter by removing (antrectomy), bypassing

(gastrojejunostomy) or obliterating (pyloroplasty) the pylorus. It can be seen in children after repair of duodenal atresia. Symptoms of bile reflux gastritis consist of epigastric pain, bilious vomiting, anemia, gastrointestinal bleeding and weight loss. Eating increases the discomfort. Endoscopy with biopsy in the presence of achloridia is diagnostic. Mainstay of treatment for bile reflux gastritis consists of histamine 2-receptor blockers, aluminum-containing antacids (to absorb bile salts) and metoclopramide (improve gastric emptying). Medical management should be tried for many months. The operation of choice is a Roux-en-y diversion. If the original operation was vagotomy with pyloroplasty, the gastric antrum should be removed (to eliminate cephalic and humoral phase of gastric secretion) and a Roux-en-y gastrojejunostomy constructed.

#### **References:**

1- Davidson ED, Hersh T: The surgical treatment of bile reflux gastritis: a study of 59 patients. Ann Surg 192(2):175-8, 1980

2- Cooperman AM: Postoperative alkaline reflux gastritis. Surg Clin North Am 56(6):1445-59, 1976

3- Sorgi M, Keighley MR: Alkaline reflux gastritis: assessment and therapy. Surg Annu 14:153-79, 1982

4- Burden WR, Hodges RP, Hsu M, O'Leary JP: Alkaline reflux gastritis. Surg Clin North Am 71(1):33-44, 1991

5- Ritchie WP Jr.: Alkaline reflux gastritis. Gastroenterol Clin North Am 23(2):281-94, 1994

## **Ovarian Teratoma**

Two-third of all malignant tumors of the ovary in children are germ cell tumors. Overall, teratoma is the most common germ cell tumor. Ovarian teratomas contain tissue from the three primitive germ cell layers in an ectopic location and seldom appear before the age of five years. Ovarian teratomas are classified as mature, immature and malignant. The vast majority of ovarian teratomas in children are benign, cystic, mature tumors. Plain abdominal films may show calcifications. Degree of immaturity depends on cellular differentiation and foci of neuroepithelium. Immature teratoma can grow into large tumors presenting with ascites, peritoneal implants and liver metastasis. Also, AFP and HCG levels can be elevated. Survival in ovarian teratoma is inversely proportional to the grade of immature elements present and stage of the disease. Mature teratomas are cured with surgical resection only. Surgery is also curative for most children and adolescents with resected ovarian immature teratoma of any grade, even when elevated levels of serum AFP or microscopic foci of yolk sac tumor are present. Chemotherapy should be reserved for cases with relapse. Upon resection surgeons must collect peritoneal fluid for cytology, examine peritoneal surface and liver, perform wedge biopsy of suspicious contralateral ovarian lesions, omentectomy and lymph node sampling of enlarged retroperitoneal nodes.

#### **References:**

4- Chaung JH, Chen L: Ovarian teratoma with gliomatosis peritonei. J Pediatr Surg 27(5):662-4, 1992

5- Brown MF, Hebra A, McGeehin K, Ross AJ 3rd: Ovarian masses in children: a review of 91 cases of

<sup>1-</sup> Piver MS, Patton T: Ovarian cancer in children. Semin Surg Oncol 2(3):163-9, 1986

<sup>2-</sup> Lazar EL, Stolar CJ: Evaluation and management of pediatric solid ovarian tumors. Semin Pediatr Surg 7(1):29-34, 1998

<sup>3-</sup> Kobayashi RH, Moore TC: Ovarian teratomas in early childhood. J Pediatr Surg 13(4):419-22, 1978

malignant and benign masses. J Pediatr Surg 28(7):930-3, 1993

6- Cass DL, Hawkins E, Brandt ML, Chintagumpala M, Bloss RS, Milewicz AL, Minifee PK, Wesson DE, Nuchtern JG: Surgery for ovarian masses in infants, children, and adolescents: 102 consecutive patients treated in a 15-year period. J Pediatr Surg 36(5):693-9, 2001

7- Marina NM, Cushing B, Giller R, Cohen L, Lauer SJ, Ablin A, Weetman R, Cullen J, Rogers P, Vinocur C, Stolar C, Rescorla F, Hawkins E, Heifetz S, Rao PV, Krailo M, Castleberry RP: Complete surgical excision is effective treatment for children with immature teratomas with or without malignant elements: A Pediatric Oncology Group/Children's Cancer Group Intergroup Study. J Clin Oncol 17(7):2137-43, 1999

8- Cushing B, Giller R, Ablin A, Cohen L, Cullen J, Hawkins E, Heifetz SA, Krailo M, Lauer SJ, Marina N, Rao PV, Rescorla F, Vinocur CD, Weetman RM, Castleberry RP: Surgical resection alone is effective treatment for ovarian immature teratoma in children and adolescents: a report of the pediatric oncology group and the children's cancer group. Am J Obstet Gynecol 181(2):353-8, 1999

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