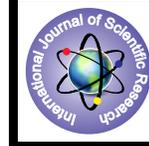


Cystic Hygroma(Hydrocele of Neck)



Medical Science

KEYWORDS :-

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ABSTRACT

Although cystic hygroma is well recognized in pediatric practice, it seldom presents de novo in adulthood. Cystic hygroma is very rare in adults but it should be considered in the differential diagnosis of neck swelling.

Patients presenting with a painless, soft, enlarging neck mass should have a careful history and physical examination along with the appropriate radiological imaging to assist the diagnosis with deeper extensions. Surgical intervention is the treatment of choice for this rare condition. Very few cases of cystic hygroma have been reported in adults. Here we are reporting a case of cystic hygroma in a 18-year-old female patient.

Case report

A 18-year-old female patient presented in the Surgery Department of kamineni institute of medical sciences, Narketpally with the complaint of a painless, progressively enlarging mass in the neck which was noticed by a family member. The mass was initially of the size of a lemon situated in the root of the neck on the right side. Over a period of five years, the swelling gradually enlarged to occupy the whole neck on the right side. Her complaints were fullness on right side of the neck since 5 years and unable to abduct the arm (pain develops on abduction >300).pain over swelling science 20 days. There was no history of trauma in the neck region, no history of recent or past upper respiratory infections, or any previous history of neck mass as an adult or child. She has no complaints of dysphasia, dyspnoea, hoarseness of voice, fever or loss of weight. No history of tuberculosis. Her past medical and family history was unremarkable.



Figure :-1 cystic hydroma over right side of neck

On examination, a soft, fluctuant, non-tender, transilluminant, well-defined, regular mass measuring approximately 15x12cm could be palpated on the right side of the neck extending from the submandibular region to the supraclavicular fossa in the vertical plane, and from the paratracheal region to the posterior triangle of the neck in the horizontal plane. There was no thrill, bruit or cervical lymphadenopathy and the remaining neck and systemic examination was normal.

Ultrasonographic evaluation revealed an irregular, multilocular, cystic anechoic lesion in the posteriolateral region of the neck, anterior to major neck vessels. CECT scan of the neck and thorax showed a large multi loculated cystic

mass lesion of near-water density with few enhancing septa within. It occupied the posterior cervical space on the right side of the neck lateral to the carotid artery. It displaced the right internal jugular vein anteriomedially and the sternocleidomastoid muscle anteriolaterally. In its cranio-caudal extent, it extended from C2 to the level of vertebra C7. A preliminary diagnosis of cystichygroma was made and the patient was scheduled for surgery.

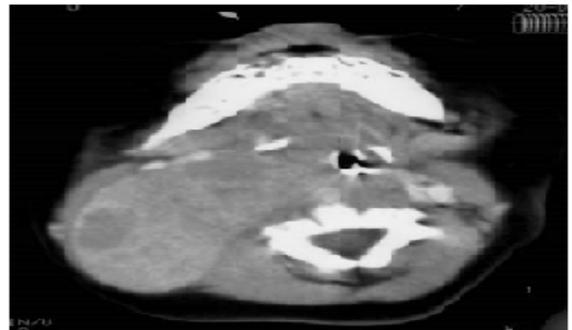


Figure 2:-CECT scan of cystic hydroma

Under general anaesthesia, curvilinear incision was taken, subplatysmal flaps were raised and the sternocleidomastoid muscle was displaced laterally to expose the multiloculated cystic lesion throughout its extent. The spinal accessory nerve was separated anteriorly all across over the cystic mass. The spinal accessory nerve lying on the anterior aspect of the lesion and the internal jugular vein were carefully preserved with meticulous dissection, and the cystic lesion was excised. Hemostasis secured.



Figure 3:- cystic hydroma with intact cyst wall

Histopathology revealed thin connective-tissue stroma separating the cystically dilated spaces lined by a single layer of benign endothelial cells which was consistent with cystic hygroma. The postoperative course was uneventful and patient was discharged in stable condition.

Review of literature:

Cystic hygroma (lymphangioma) is a benign congenital malformation of the lymphatic system that occurs as a result of sequestration or obstruction of developing lymph vessels. Lymphangiomas usually present in infancy or early childhood. Approximately 50 to 60 percent of cystic hygromas appear before the end of the first year of life. Almost 80 to 90% of cystic hygromas present before the end of the second year of life. Cystic hygroma is an uncommon differential diagnosis of a progressively enlarging neck mass in adulthood.

In lesions diagnosed pre-natally, before 30 weeks, chromosomal abnormalities are common. When diagnosed in the prenatal period, this disease may be associated with Turner syndrome, Noonan syndrome, cardiac anomalies, trisomy syndromes and fetal hydrops. [2, 4, 7]

Approximately 75-80% of cystic hygromas appear in the neck and the lower portion of the face. Other sites are axilla [20%], mediastinum [5%], abdomen [liver, spleen and colon], pelvis, retroperitoneum [kidney], groin, scrotum and skeleton. In adults, cystic hygromas are commonly seen in the sublingual, submandibular and parotid spaces. [3]

Grossly, cystic hygromas are multiloculated masses composed of many cysts. Microscopically, the cyst wall consists of a single layer of flattened epithelium and the spaces may or may not have blood-containing capillaries suggesting that there may be a combined vascular and lymphatic defect. These poorly supported blood vessels in cystic hygromas may bleed and produce rapid enlargement and discoloration of the hygroma.

Infection within the cysts (usually caused by streptococcus or staphylococcus species) may occur. This complication can also cause rapid enlargement which may result in airway obstruction. Malignant transformation has not been reported in cystic hygromas, and there have been no cases of spontaneous regression.

Clinically depending on size it is divided into 3 types;

- lymphangiomacircumscripta(5cm),
- lymphangiomadiffusum (> 5cm),
- lymphangioma ab agne (reticulate pattern).

Three types of lymphatic malformations are described: capillary, cavernous and cystic.

Pre-operative imaging is essential to look for intrathoracic extension, which is present in 10% of cases. Ultrasonography or magnetic resonance imaging (MRI) is recommended. Lymphangiomas are best visualized by magnetic resonance imaging (MRI). The high water content allows lymphangiomas to appear hyperintense on T2-weighted images.

According to de serres adapted classification for lymphatic malformation of head and neck staging is as follows :-

Stage	Location of lesion
I	Unilateral infrahyoid
II	Unilateral suprahyoid

III	Unilateral suprahyoid and infrahyoid
IV	Bilateral suprahyoid
V	Bilateral infrahyoid or suprahyoid

Doppler studies are done to evaluate the vascularity of the mass and see the relation with adjacent vessels. On CT scan, cystic hygromas appear as non-enhancing thin-walled multiloculated masses with near-water attenuation values [3].

The differential diagnosis of a neck mass in adults may include lipoma, branchial cleft cyst, haemangioma, cold abscess, lymphoma, hamartoma, thyroid mass, thyroglossal duct cyst, dermoid cyst, metastatic disease, or other tumors.

The preferred treatment for cystic hygroma is surgical excision because of the dangers of infection and fistula formation. Sometimes, this may be impossible due to the infiltrating nature of the hygroma within and around neurovascular structures, muscles and blood vessels. In this condition, unroofing, partial cystectomy and drainage of the cystic content should be performed and all adjacent crucial structures should be preserved. In these situations, a recurrence rate of 10-15 % is reported [1].

Complications following resection are postoperative muscle weakness, nerve injuries and neural weakness, bleeding and wound infection.

Laparoscopic techniques have been used in both diagnosis and treatment of retroperitoneal cystic hygromas. VATS has been performed in some cases for mediastinal cystic hygromas. Laser therapy for oropharyngeal extension has been used with good effect.

Percutaneous aspiration is not preferred because of the risk of bleeding, infection and recurrence. Injection of sclerosing agents like alcohol, bleomycin [5] and OK-432 (a lyophilized mixture of Streptococcus pyogenes and Penicillin G potassium) [6], PICIBANAL , have been reported with favorable results.

Discussion:

In our patient the cyst was located on right side of neck occupying both anterior and posterior triangle of 15 x 12 cms fitting Intode serresstage II classification. The cause for the swelling in the adulthood could not be sort out. After preanesthetic checkup patient was taken for surgery with curvilinear incision over the swelling. Spinal accessory nerve which was coursing anteriorly all across the cystic mass which was carefully isolated and spared. On the medial aspect, the mass could be carefully separated from the internal jugular vein, carotid vessels, vagus nerve and hypoglossal nerve and the mass was dissected into and sent for biopsy which revealed to be showing lymphatic pattern consistent with cystic hygroma.

Conclusion

Cystic hygromas are a rare differential diagnosis in adult neck masses and should be remembered for the adult patients who have neck, axillary, mediastinal, groin, or retroperitoneal masses. Preoperative imaging for diagnosis and to look for extension of lesion is essential along with its relation to important adjacent structures. Surgical excision is the treatment of choice though multiple techniques like aspiration, sclerocent injection etc was tried.

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