SONOGRAPHY OF CONGENITAL NECK MASSES IN CHILDREN

Sladjana Petrović¹, Dragan Petrović², Zoran Pešić², Predrag Kovačević³

¹Institute of Radiology, Clinical Centre, Niš, Serbia and Montenegro
²Clinic of Stomatology, Department of Maxillofacial Surgery, Niš, Serbia and Montenegro
³Clinic of Plastic and Reconstructive Surgery, Clinical Centre, Niš, Serbia and Montenegro
E-mail: spetrovic@bankerinter.net

Summary. Congenital neck masses in children are a relatively frequent finding. To the clinicians, they may pose various diagnostic and therapeutic dilemmas and high-resolution sonography is therefore a method of choice. Differential diagnosis of paediatric congenital neck masses primarily includes the presence of hemangioma, lymphangioma, thyroglossal and branchial cleft cysts. The aim of this paper is to examine the ability of echosonography in differential diagnosis of various congenital neck masses in children and to determine the efficacy and significance of echosonography in preoperative patient preparation. Patients and methods: The investigation enrolled 53 paediatric patients with palpable masses in the neck, who have been examined by high-resolution sonography and color doppler sonography before surgical treatment. CT and MRI, as intra-operative and histopathological findings, confirmed echosonographic diagnosis. Results: Our experience showed that sonography is a sensitive method in differential diagnosis of congenital neck masses in children. Conclusion: We recommend this method as an accurate, cost-effective, noninvasive imaging modality in the preoperative evaluation.

Key words: Congenital, neck masses, Doppler, sonography

Introduction

Congenital neck masses in children are a relatively frequent finding. To the clinicians, they may pose various diagnostic and therapeutic dilemmas and high-resolution sonography is, therefore, a method of choice. Differential diagnosis of paediatric congenital neck masses primarily includes the presence of hemangioma, lymphangioma, thyroglossal and branchial cleft cysts. High-resolution sonography with colour and power Doppler sonography is the method which can provide reliable differentiation of these changes.

Aim

The aim of the paper is to examine the ability of echosonography in differential diagnosis of various congenital neck masses in children and to determine the efficacy and significance of echosonography in preoperative patient preparation.

Patients and Methods

The investigation enrolled 53 paediatric patients with palpable masses in the neck, referred to echosonography by clinicians in the period from January 2000 till the end of December 2005. The age of the patients ranged from 3 months to 15 years. All examinations were performed at the Institute of Radiology, Clinical Centre, Niš, using conventional machines for echosonography with high-resolution multi-frequency probes (7.5-10 MHz) with colour and power Doppler option. The examinations were targeted and consisted of longitudinal, transversal and oblique grey scale sonograms, and after that colour and power Doppler options were applied and flow curves were registered. The results were postoperatively compared with intra-operative and histopathological findings, and, in some cases, ultrasound findings were preoperatively compared with CT and MRI findings.

Results

Out of the total proportion, 58% of lesions were located at the medial line and 42% in the lateral portions of the neck.

Based on the echosonographic picture, all lesions were categorised as cystic, semisolid without vascularization and solid vascularized lesions. In 21 patients solid structures were found with irregular transonic spaces demonstrating vascularization at colour Doppler sonography. Power Doppler shows an increased high vessel density, whereas pulsed Doppler demonstrates high-flow velocity and low-resistance index – these changes were categorised as hemangiomas (Fig. 1). They presented as well-vascularized solid masses, which were hypoechoegenic in 50%, hyperechoegenic in 17% and heteroechoegenic in
33% of the cases. All hypoechoic and mixed-type lesions demonstrated a higher degree of vascularization, compared to hyperechogenic ones.

In 27 patients we found cystic changes and cystic changes filled with dense fluid, without evidence of vascularization. Based on their echosonographic picture and their medial or lateral position these changes were classified as medial and lateral cysts, i.e., thyroglossal or branchial cleft cysts. Thyroglossal cysts had a variable sonographic appearance but in 48% they presented as pseudo-solid formations, homogeneously hypoechoic with internal debris in 19%; heterogeneous in 9%; and in only 24% as anechogenic, typically cystic (Fig. 2). Colour and power Doppler signals were not detected. A fistulous channel was detected in one patient. Branchial cleft cysts also had a pseudo-solid appearance in 73% of the cases and hypoechoic with internal debris in 27% (Fig. 3).
Fig. 3. Branchial cleft cyst.
   a, b) Transverse and longitudinal sonograms of the neck show a hypoechoic mass with pseudosolid echo texture and without flow on power doppler sonography.

   In 3 patients unilocular cystic changes were found, with dense contents and calcifications – they were classified as possible dermoid cysts. Dermoid cysts had a heterogeneous sonographic appearance.

   Lymphangioma presented as anechoic, multilocular cystic mass containing septa of variable thickness and solid components, without vascularization on colour Doppler sonography (Fig. 4).

Fig. 4. Macrocystic lymphangioma (cystic hygroma) of the neck.
   a,b) Ultrasound axial scans show a subcutaneous cystic mass (arrows) deeply extending, behind the carotid space, on the right of the thyroid.
   c,d) Axial T2-weighted MR images demonstrate a high signal lesion with the suggestion of a septation (arrows) that extends from the posterior triangle to the retrolaryngeal space.
In 5 patients computerised tomography was also applied, confirming echosonographic diagnosis: in 2 patients branchial cleft cysts, in 1 patient thryeoglossal cysts, in 1 hemangioma, and in 1 patient lymphangioma. In 2 patients magnetic resonance imaging was done, confirming the presence of branchial cleft cyst and lymphangioma.

After surgical treatment and histopathological verification it was established that 17 (32.0%) patients had branchial cleft cysts, 10 (18.9%) thryeoglossal cysts, 3 (5.7%) dermoid cysts and 2 (3.8%) lymphangiomas (cystic hygromas). Out of 21 patients (39.6%) with hemangiomas, 5 patients were treated surgically and other patients conservatively.

The intra-operative and histopathological findings completely correlated with the echosonographic findings.

Discussion

Hemangiomas are the most common tumours occurring in infancy (1,2). They usually appear in the first week of life and are located in the head and neck in almost 60% of cases. Haemangiomas are characterised by three phases: rapid postnatal endothelial proliferation (3-9 months); stable period of variable length; and spontaneous slow involution (approximately 18 months to 10 years). Histologically, in the proliferative phase, they are formed by well-delimited lobular masses of endothelial cells with an increased number of mast cells. Later, capillary-size lumina are often seen. During the involutive phase there is progressive perivascular deposition of fibrofatty tissue, enlargement of the vascular lumen and thinning of the endothelial lining. Diagnosis is usually made on the basis of clinical findings, and, in most instances, further investigations are not needed. Imaging is indicated either in the diagnosis of deep haemangiomas with normal overlying skin to evaluate their extent or in cases of “alarming haemangiomas”, i.e. lesions which are dangerous to vital structures (e.g. obstruction of the airway, impairment of vision, heart failure or thrombocytopenic coagulopathy) (3,4). According to Dubois and Garel, echosonography is the best imaging modality for defining haemangiomas (3). At echosonography, haemangiomas may appear homogeneously hyperechoic (multiple tiny vascular channel interfaces, proteinaceous matrix and areas of thrombosis and fibrosis), or with a typical hypoechoic lobular pattern (Fig. 1), or even like a complex mass containing vascular spaces (5). The colour and/or power Doppler show an increased high vessel density (defined as more than five structures per square centimetre), whereas the pulsed Doppler demonstrates high flow velocity (up to 90 cm/s) and a low-resistance index (RI: 0.4-0.7) with broadening of the spectrum (Fig. 1) (6). During the fibrolipomatous involution, the number of vessels decreases and RI progressively increases. Both CT and MRI provide a better tissue differentiation and, as a consequence, a better evaluation of the extent of the lesions if they need to be surgically treated. In a majority of cases, however, no treatment is required because all haemangiomas undergo spontaneous involution.

Lymphangioma is believed to develop from sequestered lymphatic sacs that fail to communicate with peripheral draining channels (7). Approximately 75% of all lymphangiomas occur in the neck, generally located in the posterior compartment (Fig. 4), and 3-10% may extend into the mediastinum (8). The presence of loose fatty tissue in the neck allows the formation of cystic hygroma, which consists of hugely dilated cystic lymphatic spaces, but a combination of the four histological types of lymphangioma (cystic hygroma, cavernous and capillary lymphangioma, vascular-lymphatic malformations) can often be seen in a single lesion (8,9). Actually, at echosonography, they appear as multinodular, predominantly cystic, masses (Fig. 4) containing septa of variable thickness and solid components. A correlation between the sonogram and pathological specimen demonstrates that the echogenic component corresponds to a cluster of abnormal lymphatic channels, too small to be resolved with echosonography (10). Haemorrhagic or infected cystic spaces are also more echogenic (11). Large lesions had ill-defined boundaries, with cystic components dissecting between normal tissue planes. Lymphangiomas are treated by surgery and MRI is the most accurate technique for evaluating the extent of the tumour and its relatedness to surgical planning (12,13,14). Sclerosing therapy with echosonography guidance may be an alternative for macro-cystic lymphangiomas (15).

Sonographically, these tumours can usually be differentiated from other cervical masses, especially soft tissue haemangiomas and venous malformations. The differential diagnosis of a cystic neck mass includes thryeoglossal duct cyst, branchial cleft cysts and dermoid tumours.

The thryeoglossal duct is the most frequent midline cyst and the most common congenital neck mass, accounting for 70% of congenital neck masses (16). The echosonography diagnosis is readily made by following the base of the tongue in the midline towards the sternal notch. On echosonography, thryeoglossal duct cysts in children are not simple cysts but have a complex pattern ranging from a typical anechoic cyst to a pseudo-solid appearance (most common; Fig. 2) (17). The origin, as well as a sinus to the base of the tongue, is sometimes identified (thryeoglossal duct cysts move with movement of the tongue, whereas other masses generally do not) (18). Thryeoglossal-duct remnant should be differentiated from ectopic thyroid prior to surgical excision. When the thyroid gland can be identified in the normal position, coexistent ectopic thyroid is seldom found, even if it is possible (19); however, when ectopic thyroid is suspected, Tc-99m and thyroid function tests are valuable (20).

The branchial cleft cysts are the most common non-inflammatory paediatric lateral neck masses; almost all arise from the second branchial cleft

SONOGRAPHY OF CONGENITAL NECK MASSES IN CHILDREN  167
Unilocular cystic midline neck mass at the suprasternal glands, sweat glands and dermal appendages. A contain only ectodermal and mesodermal elements true teratomas, which contain tissue elements derived cyst, which has only a squamous epithelium, and to cholesterol crystals. A fluid-filled cyst usually can be distinguished from a solid mass by the enhancement of the US beam posterior to the cyst.

Dermoid cysts are benign, usually unilocular, dermalized structures. In contrast to the epidermoid cyst, which has only a squamous epithelium, and to true teratomas, which contain tissue elements derived from all three germinal layers, the dermoid cysts contain only ectodermal and mesodermal elements. They include hair follicles, hair, sebaceous glands, sweat glands and dermal appendages. A unilocular cystic midline neck mass at the suprasternal notch in a child should suggest a dermoid cyst.

All the cysts have internal echoes, with a solid appearance with only slight or no posterior echo enhancement. Amorphous keratinous debris from keratinizing stratified squamous epithelium fills the lumen of each cyst, producing the internal echoes.

Epidermoid cysts are rare lesions in the head and neck. Most often they are located in the submental region. Commonly they have the same echosonography appearance as dermoid cysts. A cystic tumour with the unusual echosonography aspect of multiple smaller spherical formations caused by multiple spherical keratin formations has been described.

**Conclusion**

The sonographic appearance of congenital cystic masses of the neck in children is variable; to make correct preoperative assessment, the sonologist must be familiar with these characteristics. Cystic cervical masses do not seem to be simple cysts, as previously suggested, but instead they have a complex cystic pattern ranging from an anechoic to a pseudo-solid appearance. Solid and mixed cervical masses, like hemangiommas and cystic hygromas, had a typical ultrasound and colour Doppler appearance. Our experience has shown that sonography is a sensitive method in differential diagnosis of congenital neck masses in children. We recommended this method as an accurate, cost-effective, noninvasive imaging modality in the preoperative evaluation.

**References**


SONOGRAFIJA KONGENITALNIH MASA NA VRATU DECE

Sladjana Petrović¹, Dragan Petrović², Zoran Pešić², Predrag Kovačević³

¹Institute za radiologiju, Klinički centar Niš
²Stomatološka klinika, Odeljenje maksilofacijalne hirurgije, Niš
³Klinika za plastičnu i rekonstruktivnu hirurgiju, Klinički centar, Niš
E-mail: spetrovic@bankerinter.net

Kratak sadržaj: Kongenitalne mase na vratu u dece su relativno čest nalaz. Mogu zadavati kliničarima dijagnostičke i terapeutske dileme, zbog čega se visokorezolutivna sonografija preporučuje kao metoda izbora. Diferencijalna dijagnoza kongenitalnih masa na vratu u dece primarno uključuje prisustvo hemangioma, limfangioma, tireoglosalnih i branhijalnih cističnih promena. Cilj ovog rada je da se utvrdi mogućnost ehosonografije u diferencijalnoj dijagnostici različitih kongenitalnih masa na vratu u pedijatrijskim pacijenata, i odrediti efikasnost i značaj ehosonografije u preoperativnoj pripremi pacijenata. Ovo istraživanje obuhvata 53 pedijatrijska pacijenta sa palpabilnim masama na vratu, koji su bili podvrgnuti visokorezolutivnoj i color doppler sonografiji pre hirurškog tretmana. CT, MRI kao intraoperativni i patohistološki nalazi potvrdili su ehosonografske dijagnoze. Preporučujemo ovu metodu kao senzitivan, jeftin, neinvazivan modalitet imaging-a u preopertivnoj pripremi pacijenata.

Ključne reči: Kongenitalne mase vrata, Doppler, sonografija