

Intramuscular hemangioma in neck: A case report

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Abstract

Hemangiomas of skeletal muscle represent 0.8% of all benign vascular neoplasm¹. Of these 13.8% occur in the head and neck region², with the masseter muscle being the most common site, followed by the trapezius and sternocleidomastoid muscles respectively^{2,3}. Intramuscular Hemangiomas [IMH] generally occur in the first three decades of life⁴.

Key words: Intramuscular hemangioma, USG, CECT, CT angiography.

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INTRODUCTION

A 24 year old girl presented with a swelling in the right supraclavicular region of 11 months duration. Swelling gradually increased in size, becoming more pronounced. Over the last one month she developed pain over the swelling. She gave no history of trauma or oral contraceptive pill usage. On physical examination there was a swelling in the right supraclavicular region 3 x 2 cm which was non-tender. On rotating the neck to the

opposite side the swelling became comparatively taught than normal. Imaging include X ray and USG, followed by CECT scan and CT angiography. On plain X ray film (PA view) chest showed multiple small rounded calcified opacities with central lucency in right supraclavicular region extending into the soft tissue planes on right side itself.(Figure 1) An ultrasound scan of the abdomen and pelvis was done on GE Voluson S6 using 3.5-5MHz frequency transducer, it revealed a multiseptated irregular cystic lesion measuring 26 x 15 mm in the right supraclavicular region,(Figure 2) the cystic lesion was anechoic with multiple septas within with no evidence of internal echoes. The lesion extending into the right lateral neck region, it appeared to be in communication with the Sternocleidomastoid muscle though it was unclear on USG. On Doppler study, the lesion did not show any vascularity. (figure 3)The right common carotid artery appeared to be normal. (figure 4) and other major vascular structures appeared to be normal.



Figure 1



Figure 2



Figure 3



Figure 4

Figure 1:

Figure 2: A multiseptated cystic lesion anechoic lesion in right supraclavicular region

Figure 3: No evidence of colour flow uptake, the lesion extending into right lateral neck region

Figure 4: Right common carotid artery appears normal, while the lesion is far lateral from right common carotid artery

CECT was done, it revealed a soft tissue density lobulated poorly defined lesion with multiple calcific foci in right side of neck, extending inferiorly to involve upper part of posterior chest, superiorly to involve the posterior wall of the pharynx and posterior triangle of neck with mild compression and displacement of adjacent muscles and internal jugular vein. (figure5). It also showed involvement of right paraspinal muscles, the lesion also show focal areas of pooling of contrast with in the lesion on post contrast study. (figure 6 and 7). CT angiography

was done which revealed an aberrant branch arising from right subclavian artery lateral to the origin of vertebral artery (figure), which appears to be the feeder artery, the aberrant artery further giving multiple branches to the mass (figure 8 and 9) FNAC revealed greenish color aspirate, cytology of which did not reveal any cellular material, this probably was extravasated blood and the greenish tinge, due to breakdown products of hemoglobin. This was evident as areas of hemorrhage on histology.

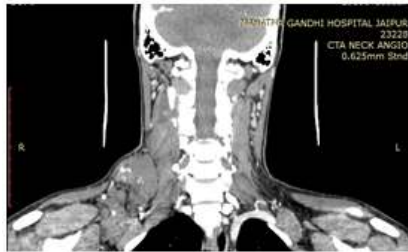


Figure 5

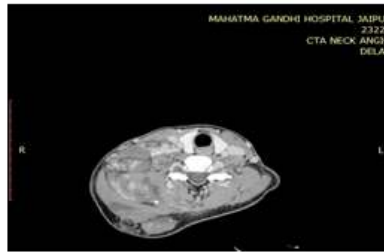


Figure 6



Figure 7



Figure 8



Figure 9

Figure 5 Figure 6: Axial CECT image showing patchy heterogenous enhancement of the mass lesion on right side, its also showing involvement of the right paraspinal muscles.

Figure 7: Sagittal CECT image showing pooling of the contrast in seen in multiple dilated sinusoidal channels.

Figure 8: CT angiography showing an aberrant vessel arising from right subclavian artery, lateral to the origin of the right vertebral artery, multiple rounded phleboliths also noted.

Figure 9:

DISCUSSION

Hemangiomas are the most frequent benign vascular neoplasm, being four times more frequent in females. Usually they appear as a single mass, but in a small percentage, they can be multiple. Hemangiomas develop in the head and neck, followed by the trunk and extremities^{5,6}. The most common localization of intramuscular hemangiomas is the lower extremities, followed by the upper extremities. These soft tumors are often placed subcutaneously, and intramuscular extension is rare⁷. Cavernous hemangiomas are larger in size and deeper, with a high tendency in thrombosis and calcification⁸. The case we presented enclosed the previously described characteristics, the lesion being discovered in a young female and localized in the upper limb, but unlike the other studies, the tumor extended to the brachial muscle fibers. Sinusoidal hemangioma is a

rare variant of cavernous hemangiomas⁸. Imaging diagnosis of these lesions is usually made by MRI and angiography. X-ray can reveal a nonspecific mass with calcification and bone changes if the tumor is located adjacent to the osseous structures⁹. The CT may identify a soft tissue mass with densities similar to muscles and also the phleboliths, even the small ones. The tumors enhance after contrast media administration, with depiction of serpentine vascular areas¹⁰. The imaging evaluation of choice is MRI, this imaging technique being important in the assessment of these tumors. MRI is able to determine the extent in the surrounding tissue and accurately differentiate between vascular lesions with low-flow and high-flow^{11,12}. Haemangiomas are readily distinguished from other soft-tissue tumours by computerized tomography (CT), MRI, and arteriography. CT is useful for defining the form, size and anatomic relationship of

the tumour but MRI is the method of choice in defining the vascular nature of the tumour. On T1 weighted imaging, haemangiomas are iso-intense or hypo-intense to muscle. With T2 imaging, the lesions are hyper-intense on account of the volume of stagnant blood, clearly differentiated from the normal muscle and fibro-fatty septa¹³ Intra-muscular haemangiomas usually present as a slow-growing mass with distinct margins, mobile and do not exhibit any of the vascular signs such, discoloration of overlying skin or pulsation. The differential diagnosis includes: neurofibroma, lipoma, dermoid cyst, enlarged lymph nodes, soft-tissue sarcoma, myositis ossificans and temporal arteritis^{14,15}. Arteriography is helpful in delineating major vascular feeders for pre-operative embolization.

REFERENCES

1. Watson WL, McCarthy WD. Blood and lymph vessel tumors; a report of 1056 cases. *SurgGynecol Obstet.* 1940; 71:569–588.
2. Wolf GT, Daniel F, Krause CJ, Kaufman RS. Intramuscular Hemangioma of the head and neck. *Laryngoscope.* 1985; 95:210–213. doi: 10.1288/00005537-198502000-00018. [PubMed] [Cross Ref]
3. Ingalls GK, Bonnington GJ, Sisk AL. Intramuscular hemangioma of the temporalis muscle. *Oral Surg Oral Med Oral pathol.* 1985; 60:476–481. doi: 10.1016/0030-4220(85)90232-4. [PubMed] [Cross Ref]
4. Rossiter JL, Hendrix RA, Tom L, Potsic W. Intramuscular hemangioma of the Head and neck. *Otolaryngol. Head Neck Surg.* 1993; 108:18. [PubMed]
5. Drolet B, Esterly N, Frieden I. Hemangiomas in children. *N Engl J Med.* 1999; 341:173–181. [PubMed]
6. Zhang L, Lin X, Wang W. Circulating level of vascular endothelial growth factor in differentiating hemangioma from vascular malformation patients. *PlastReconstr Surg.* 2005; 116:200–204. [PubMed]
7. Song BH, Youn SH, Park EJ. A case of sinusoidal hemangioma with lipoma. *Dermatol.* 2011; 23:250–253. [PMC free article] [PubMed]
8. Calonje E, Fletcher CD. Sinusoidal hemangioma. A distinctive benign vascular neoplasm within the group of cavernous hemangiomas. *Am J SurgPathol.* 1991; 15:1130–1135. [PubMed]
9. Wild AT, Raab P, Krauspe R. Hemangioma of skeletal muscle. *Arch Orthop Trauma Surg.* 2000; 120:139–143. [PubMed]
10. Donnelly LF, Adams DM, Bisset GS. Vascular malformations and hemangiomas: a practical approach in a multidisciplinary clinic. *AJR Am J Roentgenol.* 2000; 174:597–608. [PubMed]
11. Van Rijswijk CSP, Van der Linden E, Van der Woude HJ. Value of dynamic contrastenhanced MR imaging in diagnosis and classifying peripheral vascular malformations. *AJR Am J Roentgenol.* 2002; 178:1181–1187. [PubMed]
12. Greenspan A, McGahan JP, Vogelsang P. Imaging strategies in the evaluation of soft tissue hemangiomas of the extremities: correlation of the findings of plain radiography, angiography, CT, MRI, and ultrasonography in 12 histologically proven cases. *Skeletal Radiol.* 1992; 21:11–18. [PubMed]
13. Buetow PC, Kransdorf MJ, Moser RP Jr, Jelinek JS, Berrey BH. Radiologic appearance of intramuscular hemangioma with emphasis on MR imaging. *Am J Roentgenol* 1990; 154:563-7. [PubMed]
14. To EW, Tsang WM, Pang PC, Ahuja A. Cavernous hemangioma of the temporalis muscle: report of a case. *J Oral MaxillofacSurg* 2001; 59:1229-32. [PubMed]
15. Bui-Mansfield LT, Myers CP, Fellows D, Mesaros G. Bilateral temporal fossa hemangiomas. *Am J Roentgenol* 2002; 179:790. [PubMed]

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