Case Report:

Intramuscular Haemangioma of triceps muscle

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Abstract:
A 14 years boy presented with a swelling in the left elbow on his back measuring 6cm X 4cm X 3cm extending from tip of elbow to axilla on his back, with pain and restriction of movements. Range of motion is flexion 90 degree to 150 degree, supination pronation okay; with a mild history of trauma (?) trauma) at his childhood and it is very slowly increasing. On examination, it is doughy feeling, no definite mass is felt. On raising the arm above heart it does not reduce. There is a vague history of trauma at the age of 6 years. Blood parameters are normal. X-ray revealed that in the vicinity of soft tissue swellings there are multiple pearl like swellings starting from 5mm to 8mm around the elbow. Sonogram guided study showed it may be haematoma. CT scan study was same as X rays. On incision, sprouting of blood through the intramuscular tissue of triceps muscles. No involvement of ulnar nerve or periosteal or bone and synovia or joint. Histopathology revealed intramuscular haemangioma

Key words : Triceps, Intramuscular Haemangioma

INTRODUCTION:
Intramuscular Haemangioma are rare entity. Only 0.8% of the tumours in Watson and McCarthys’ study were intramuscular haemangioma. The rate of correct diagnosis is only 8% to 19% in reported literature. Of the reported cases, approximately 45% is located in the lower extremities. The disease occurs predominantly (90%) within first three decades. Benign tumour making up to 0.8% (1,5,6) of the cases were intramuscular haemangioma and 8% of all benign tumours (6,7). It is more common in the lower limb especially in the quadriceps. The etiology of intramuscular haemangioma is unknown or controversial. Trauma or congenital theories (3,8,9,10). Triceps muscle intramuscular haemangioma are uncommon that’s why it is presented. Vascular malformation is often confused with haemangioma. Females are more common than male (1) Haemangioma are common in the skin, subcutaneous tissue followed by the deep tissue. Occasionally are intra articular synovial haemangioma may occur(12) and rarely in the bone, females are more than males(1,12). They are characterized by rapid endothelial growth. and 80% are located in head neck region. Rare in the extremity (13), especially in the upper extremity

CASE REPORT:
A 14 years male presented with a diffuse swelling on his back extending from tip of elbow to axilla measuring 6cm X 4cm X 3cm. No tethering of skin or skin involvement. The swelling is tender on movements of elbow joint. Doughy feeling. No thrill no bruit no reduction of swelling when the arm is raised above heart, no involvement of ulnar nerve.
nerve. Neurovascular examination was normal no palpable axillary lymph node. Range of motion of elbow Flexion/Extension 90 degree to 150 degree. Supination Pronation is normal no bony tenderness or no valgus or rotational disability. There is a vague history of trauma at the age of 6 year. Very it is increasing slowly increasing . Blood parameters are normal. X-rays showed (phlebolith) pearl like swellings around elbow in the vicinity of soft tissue swellings .FNAC guided by sonogram revealed Hematoma, CT Scan reports are as X-rays.Fig 1 During operation :Under tourniquet and supraclavicular block Per operatively it was revealed that Bloods are sprouting through intermuscular fibres of Triceps muscles( Fig -2) on the affected side, Pearl bits are coming out through muscle fibres No definite capsule is found no involvement of bone and capsule or periosteal or joint .Ulnar nerve is not involved in the lesions, soft tissue is removed for histopathology along with normal triceps muscle (12)

Sections examined from the specimen shows dilated and communicating large thin walled blood vessels (many containing thrombus, focally calcified) interspersed with adipose tissue fibro collagenous tissue and skeletal muscles. Histological features are suggestive of intramuscular haemangioma. No malignancy is reported.

**DISCUSSION:**

Intramuscular haemangioma are rare benign tumours making upto 0.8%of all haemangiomas (1,12),before age 30 years(4,12,); recurrence of intramuscular haemangioma is not uncommon due to incomplete excision of the lesion of the infiltrative growth pattern(12) Pain is the cardinal feature of presentation along with swelling. Lower limb is more affected than upper limb .Quadriceps is most frequently affected.(7)

Imaging strategies in the evaluation of soft tissue haemangioma
Plain X-ray of intramuscular haemangioma shows soft tissue swelling. It shows soft tissue swelling and only 25%-30% of cases shows phlebolith (5, 11). Periosteal reaction adjacent to haemangioma may mimic osteomyelitis or bone tumour (12). Periosteal reaction is not always seen.

CT Scan:
III defined mass of similar attenuation to muscle may be identified. Phlebolith if present may be seen.

MRI Scan:
Lobulated, and heterogeneous soft tissue mass with no features of local invasion. Phlebolith if present may be seen.

Signal characteristics:
T1 overall signal is often intermediate to slightly high signal (relative to skeletal muscle) (14).

Some focal high signals may be present in a large proportion of lesions (upto70%)

T2 high signal intensity tends to dominate T2 weighted images

Correlation of the findings of plain radiography, angiography, CT, MRI, Ultrasonograph, in 12 histologically proven cases of soft tissue haemangioma. By Adam Greenspan M D, Jhon Pet al in the Journal of Skeletal Radiology January 1992 Volume 21 issue 1:

- In 8 out of 9 intramuscular haemangioma plain radiograph demonstrated phleboliths suggested the diagnosis while 3 had normal radiograph. (in Springer Link)

- Angiogram showed pathognomonic features of soft tissue haemangioma in six patients which are deeply situated because of its invasiveness. MRI was characterised in all three patients. The lesion demonstrated intermediate signal intensity on T1 weighted spin echo images and extremely bright signal on T2 weighting. Ultrasonogram showed hypoechoic soft tissue mass in one case and mixed echo pattern in other. Doppler signal CT showed a nonspecific mass in one of four cases and a mass with phlebolith in three.” (15).

Patients with intramuscular haemangiomas may present with pain and swellings for years. The gross and microscopic appearance: capillary type is nonvascular and spongy whereas cavernous type is composed of large thin walled dilated vessels lined by flattened endothelial cells (6).

Histologically haemangioma are classified based on the predominant type vascular channel as follows (11):

1. Capillary haemangioma which are composed of small vessels lined by flattened endothelium. It is commonest type and subdivided into Juvenile, verrucae’s and senile type.

2. Cavernous haemangioma: which are composed of dilated blood filled spaces, lined by flattened endothelium.

3. Venous haemangioma which is composed of thick walled vessels containing muscles.

Intramuscular haemangioma may have 9% recurrences’ rate (5) if incomplete surgical excision is done, but never metastasize or undergo malignant transformation. (5, 12).

Hamartomas’ and malformations: Hamartomas’ are result from an error in embryological development and are present at birth. In the case of vascular haemartomas and malformations, they may become more obvious clinically sometime after birth as a consequence of progressive ectasies. The constituent vessels may be capillaries, veins, arteries, lymphatics, or a combinations of these vessels.

Recent (Vascular malformations: Classification (16))

Recent Hypothyses and Observations Regarding the Pathogenesis of haemangiomas:

Vaasoproliferative tumours result from vasculogenesis (formations of primitive blood vessels from angio blast) rather than from...
angiogenesis (The growth of vessels from pre-existing vessels) within the anomaly that may originate from the
placenta(18).

As previously thought By(17)Immature cells or progenitor stem/Progenitor cells are identified

[ Comparison of Previous Terminology and New ISSVA(International Society for the study of Vascular Anomalies) ]

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<thead>
<tr>
<th>Previous</th>
<th>ISSVA</th>
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<tbody>
<tr>
<td>Capillary or cavernous haemangioma of any organ(s)</td>
<td>Infantile haemangioma(s)</td>
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<td>Infantile haemangioendothelioma(s)of the liver</td>
<td>Hepatic or infantile haemangioma(s)</td>
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<td>Adult vertebral haemangioma</td>
<td>Venous malformations</td>
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<td>Adult hepatic haemangioma</td>
<td>Venous malformation</td>
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<td>Adult orbital cavernous haemangioma</td>
<td>Venous malformation</td>
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REFERENCES:

5. Allen PW,Enzingerangioma of skeletal muscle analysis of 89 casesesCancer 1972 ;29: 8 22,
6. A T Wild ,PRaab, RKrauspe; Skeletal haemangioma, Archive of orthopaedic and Trauma Surgery Feburary 2000 VOLUME 120 Issue 3-4 pp139-143


16. Lisa H. Lowe M D., Tracy C., Marchant, DO., Douglas C. Rivard, DO and Amanda J., Scherbel, BS; - Vascular Malformations: Classification and Terminology the Radiologist needs to Know. ELSEVIER Seminars in Roentgenology pp106-117
