A suspicious haematoma

On Bon Chan

Case

A man, 75 years of age, presented with an enlarging lump in his left thigh after he fell from the roof six months ago. He saw a general practitioner (GP) several weeks after the initial injury. An ultrasound scan showed a large intramuscular collection of 20 cm x 8 cm x 14 cm. The diagnosis of a large intramuscular haematoma was made. The GP advised him to manage the swelling with a warm compress and simple analgesics.

Five months later, the patient presented again because the lump had been growing and was affecting his ability to walk. The lump caused minimal pain; however, he reported weight loss of 10 kg and feeling fatigued. A physical examination revealed a large, firm, immobile mass with significant oedema in the medial aspect of the left thigh.

Urgent magnetic resonance imaging (MRI) was performed. The MRI showed

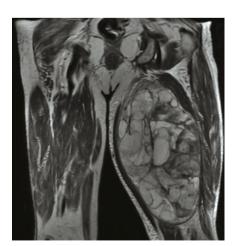


Figure 1. Magnetic resonance imaging of patient's limbs

a large lobulated heterogeneous solid mass measuring approximately 15.4 cm x 9.7 cm x 25.6 cm (Figure 1) situated between the adductor magnus, the semitendinosus and semimembranosus.

Question 1

What are the differential diagnoses?

Answer 1

The size, growth rate of the soft tissue mass and the patient's systemic symptoms raise suspicion of a malignant tumour. The differential diagnoses include sarcoma, metastatic tumour or lymphoma.

Case continued

The patient lived three hours away from the tertiary referral centre, so his case was discussed over the phone with a sarcoma surgeon. An urgent consultation was arranged and, subsequently, a biopsy was performed. The biopsy confirmed the diagnosis of a myxofibrosarcoma.

Question 2

What is a myxofibrosarcoma?

Question 3

What are the clinical features of a soft tissue sarcoma?

Question 4

How can the diagnostic accuracy for sarcomas be improved?

Answer 2

Myxofibrosarcoma is a subtype of soft tissue sarcoma,1 a rare malignant tumour of connective tissues. Sarcomas account for <1% of adult malignancy

and there are more than 100 histological subtypes.1 Myxofibrosarcoma is an aggressive subtype that usually occurs in elderly patients, predominantly in the extremities.1 It tends to recur locally and metastasise to the lungs and bones.2

Answer 3

Soft tissue sarcoma usually presents as a gradually enlarging, painless mass.3 In some cases, such as this one, patients present shortly after an injury. To date there is no evidence for a causal relationship between trauma and sarcoma. It is likely that the injury brings the patient's attention to the tumour site, rather than causing the tumour.4 When the tumour grows to a certain size, it can compress the surrounding neurovascular structures, causing oedema and discomfort. There can also be associated systemic features, such as weight loss and fever.3

Sarcomas tend to favour the extremities. Figure 2 shows the anatomical distribution of soft tissue sarcomas in 4550 adults collected by the American College of Surgeons.⁵ Clinical features that have been identified as suggestive of a sarcoma are listed in Box 1.3,6,7

Box 1. Clinical features suggestive of sarcoma^{3,6,7}

Size >5 cm in any diameter Deep or inherent to fascia Painful or tender Rapidly increasing in size Recurrence of a lump after previous excision

Answer 4

Diagnosis of soft tissue sarcoma is commonly delayed. Given that sarcomas

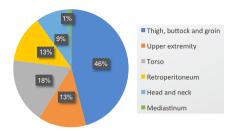


Figure 2. Anatomical distribution of soft tissue sarcomas in 4550 adults⁵

are rare, and it is much more likely for GPs to encounter a benign lump than a sarcoma, deciding which lump to investigate further can be difficult.3

Patients with an initial diagnosis of an intramuscular haematoma should be re-assessed in four weeks to ensure improvement or resolution. If the lesion does not improve or resolve, the diagnosis of a sarcoma should be suspected.

The current recommendation for a suspected sarcoma is an urgent referral to a sarcoma unit.8 However, many remote communities in Australia do not have immediate access to a specialist centre. In such situations, the diagnosis of a sarcoma will rely heavily on the GP's

clinical skills and awareness of sarcoma. When a sarcoma diagnosis is suspected, a phone consultation with a sarcoma specialist may help to provide guidance on further investigation and management.

Case continued

Further imaging at the specialist unit found metastases in the patient's lungs. The patient underwent radiotherapy and subsequently had surgical excision of the soft tissue sarcoma on his left thigh. The surgical excision was of palliative intent because of the metastatic lung disease. At the time of writing, he had returned home and had recovered well from his surgery.

Key points

- Consider malignancy in the differential diagnosis of a soft tissue mass.
- Be familiar with the clinical features of sarcoma.

Author

On Bon Chan MBBS, FRACGP, BAppSc (Physiotherapy), DCH, Certificate of Dermoscopy, General Practitioner, Sale Medical Centre, Sale, Vic. louischan451@gmail.com

Competing interests: None.

Provenance and peer review: Not commissioned, externally peer reviewed.

Acknowledgements

I would like to thank Dr Wing Ka Vicki Chan and Dr lain Nicholson for reviewing this article.

References

- 1. Fletcher C, Unni K, Mertens F. World Health Organization classification of tumours: Pathology and genetics of tumours of soft tissue and bone. Lyon, France: IARC Press, 2002; p. 102. Available at www.iarc.fr/en/publications/pdfs-online/ pat-gen/bb5/BB5.pdf [Accessed 25 August 2017].
- 2. Morrison AS. Soft tissue: Fibroblastic/ myofibroblastic tumors - Myxofibrosarcoma. Michigan: Pathologyoutlines.com, 2002-16. Available at www.pathologyoutlines.com/topic/ softtissuemyxofibrosarcoma.html [Accessed 11 August 2017].
- 3. Pike J, Clarkson P, Masri B. Soft tissue sarcomas of the extremities: How to stay out of trouble. BCMJ 2008:50(6):310-17.
- 4. Morrison BA. Soft tissue sarcomas of the extremities. Proc (Bayl Univ Med Cent) 2003;16(3):285-90.
- Lawrence W Jr, Donegan WL, Natarajan N, Mettlin C, Beart R, Winchester D. Adult soft tissue sarcomas. A pattern of care survey of the American College of Surgeons. Ann Surg 1987;205(4):349-59.
- 6. Sinha S, Peach AH. Diagnosis and management of soft tissue sarcoma. BMJ 2010;341:c7170.
- Hussein R, Smith MA. Soft tissue sarcomas: Are current referral guidelines sufficient? Ann R Coll Surg Engl 2005;87(3):171-73.
- Cancer Council Australia, Sarcoma Guidelines Working Party. Clinical practice guidelines for the management of adult onset sarcoma. Sydney: Cancer Council Australia, 2017. Available at http:// wiki.cancer.org.au/australia/Guidelines:Sarcoma [Accessed 11 August 2017].