CASE PRESENTATION
A 54-year-old woman presented with a large mass on her right hand that was progressively enlarging over a period of a few years. It was located on the palm in the thenar eminence. She did not have any neurological symptoms or muscular dysfunction, but had cosmetic concerns about the enlarging mass and reported that it interfered with her routine activity.

Physical examination revealed a soft, mobile, non-tender lobulated mass with well-defined margins. The overlying skin was intact, with no evidence of redness or increased temperature. The patient had no clinical evidence of compressive neuropathy or muscle dysfunction. The provisional diagnosis was a soft-tissue mass of unknown aetiology, and further evaluation with radiography and magnetic resonance (MR) imaging was suggested. What do the radiographs and MR images show? Is there cause for concern?
Frontal and oblique radiographs of the right hand (Figs. 1a & b) demonstrated a soft-tissue swelling between the first and second metacarpals. It had a central lucent component, suggesting the presence of a lipomatous lesion. No intralesional calcification was seen. MR images of the right hand (Figs. 2a–d) revealed a lobulated soft-tissue mass in the thenar space between the adductor pollicis brevis and opponens pollicis muscles, superficially extending to the dorsal aspect of the first web space. The mass was hyperintense on T1-weighted images (Figs. 2a & b), with multiple thin internal septations, and showed uniform suppression of signal on fat-saturated images (Figs. 2c & d), which was consistent with a lipoma. No enhancing component was seen within the lesion on the post-intravenous, contrast-enhanced, T1-weighted, fat-saturated image (Fig. 2d). The lesion measured approximately 17 cm × 10 cm. There was no involvement of adjacent muscles or encasement of the tendon sheaths and adjacent neurovascular structures.

Diagnosis
Intermuscular lipoma in the thenar eminence of the right hand.

Clinical Course
Surgical excision was performed, as the lipomatous mass was interfering with the patient’s routine activity and for cosmetic concerns. During surgery, a soft-tissue mass was seen; it was deep to the thenar and adductor pollicis muscles, and superficial to the first dorsal interosseus muscle in the region of the deep arterial arch of the right hand. Histopathology revealed benign mature adipose tissue with occasional fibrous septations, in keeping with benign lipoma. No evidence of malignancy was noted.

Discussion
Lipomas are the most common tumour throughout the body. They are more frequently seen in obese individuals between the fifth and seventh decades of life. Lipomas are often associated with other conditions such as diabetes mellitus, hyperlipidaemia, goitre and Cushing’s syndrome. More rarely, they are associated with syndromes such as adiposis dolorosa, Gardner syndrome, Madelung’s disease and familial multiple lipomatosis. Although fat is abundant in the deep palm, lipomas are rarely encountered in this location. The reported incidence of lipomas in the hands and fingers is about 1% of all lipomas that occur in the body. Such lipomas are more commonly found in the subcutaneous plane. Less common locations include the subfascial plane, within Guyon’s canal, the carpal tunnel or deep palmar space, where they are usually of larger size. Thickness of the deep central palmar fascia may account for a more peripheral location in the thenar and hypothenar eminence. This thickness also makes it clinically difficult to correctly assess the exact size and extent of the lipoma.

The clinical course of lipomas is often prolonged, with initial insidious growth followed by a prolonged and latent maintenance phase. Deep palmar lipomas are usually asymptomatic; symptoms arise mainly due to mass effect. Pain and paraesthesia secondary to nerve compression are more likely to occur if the lipoma is found in a closed anatomical space such as the carpal tunnel.
(Fig. 3) and Guyon’s canal, compressing the median nerve and the ulnar nerve, respectively. Some patients may present to the clinician either for cosmetic reasons or abnormal grasping due to the large size of the lipoma. Muscular atrophy and muscle paralysis are very rarely encountered. Another rare presentation is acute neuropathy due to nerve compression by interstitial haemorrhage.

Lipomas larger than 5 cm are called giant lipomas, such as in the case presented. Lipomas usually grow at a very slow rate and the aetiology of rapid growth into giant lipomas is debatable. It has been postulated that blunt trauma can cause disruption of fibrous septa between the skin and deep fascia, facilitating growth of fatty tissue. No apparent association has been established between tumour volume and patient symptomatology. Rydholm and Berg, in a retrospective study of 428 cases of musculoskeletal lipomas, found that the incidence of solitary lipoma to sarcoma was 150/1 for lesions ≤5 cm and 20/1 for lesions > 5 cm. Hence, patients should be informed that giant lipomas have an increased risk of malignancy.

History and clinical examination are usually sufficient to diagnose superficial lipomas. To diagnose deeper lesions, imaging is required to map the surrounding anatomic structures and detect any neurovascular structure involvement. Clinically, deeper lesions are soft, mobile, non-tender and doughy on palpation. However, the surgeon may not be able to palpate the entire extent of the deeper lesions due to overlying thick palmar fascia. In patients with compressive neuropathy, Tinel’s sign (defined as distal paraesthesia secondary to tapping on the tumour) may be elicited. Electrophysiological studies can be useful, but are not definitively diagnostic of lipomas. On radiography, lipomas are nonspecific, often appearing simply as focal soft-tissue swelling. Lucency indicative of a fatty component (Fig. 4) or internal calcification may sometimes be present. Large lesions can cause extrinsic erosion of the underlying bones. On ultrasonography, lipomas commonly appear as elongated isoechoic to hyperechoic lesions, often with well-defined margins (Fig. 5a & b). No internal or increased vascularity and posterior acoustic enhancement are seen. The relationship of the lesion with adjacent neurovascular and musculotendinous structures can be evaluated. However, definite tissue characterisation and determination of extension and relationship with adjacent structures are inferior to the findings of MR imaging.

MR imaging, with its multiplanar capability and superior tissue characterisation, is the imaging modality of choice in the assessment of lipomas. The use of MR imaging is recommended before biopsy attempts, as it may help in the selection of suspicious areas for histological evaluation. Non-image-guided biopsies can result in non-representative biopsy samples, potentially leading to

Fig. 5 Deep palmar lipoma in the thenar eminence of the left hand in a 61-year-old woman who presented with a lump of several months’ duration. Extended field-of-view (a) longitudinal and (b) transverse US images of the palm show a heterogeneously echogenic solid lump (arrows) in the deep thenar eminence superficial to the thenar muscles (arrowheads). The margins are slightly ill-defined.

Fig. 6 Deep supramuscular palmar lipoma of the left hand in a 61-year-old woman who presented with a lump of several months’ duration associated with radial side numbness of the fourth finger. (a) Axial T1-W, turbo spin-echo (TSE) and (b) T1-W, TSE, fat-saturated with contrast MR images of the left hand show a thin-walled, septated lesion (arrow) that is hyperintense on T1-W MR imaging, with uniform suppression of fat signal (arrowhead in b) on the fat-saturated image suggestive of lipoma. The lesion is located superficial to the underlying flexor digitorum superficialis tendons and thenar muscles. No obvious enhancing soft-tissue component is seen.
misdiagnosis. In contrast, MR imaging clearly defines the deeper extension, adjacent tendons and neurovascular structures, which is vital if surgery is being contemplated; the degree of compression of the underlying nerves is often well-demonstrated. Classically, a lipoma appears as a slightly lobulated homogeneous mass with well-defined borders, and demonstrates hyperintense signal on both T1- and T2-weighted sequences. Small, thin, internal hypointense septa may be present. Complete suppression of the fatty signal on fat-saturated sequences would show the benign nature of the mass. It can be located above muscle (i.e. supramuscular; Fig. 6a & b), below muscle (i.e. submuscular), between muscle (i.e. intermuscular; Fig. 2) and within muscle (i.e. intramuscular). Intramuscular lipomas usually demonstrate intermingled muscle fibres and are rare among the described subtypes. Areas of incomplete fat suppression with nodular and thick enhancing septa as well as a solid enhancing soft-tissue component are suspicious of malignancy, and liposarcoma should be excluded. Recurrent lipomas should also be viewed with suspicion, since the risk of malignancy is higher. In a review of 134 cases of hand and wrist tumours, Capelastegui et al found a positive predictive value of up to 94% when comparing MR images with histological reports.

Conservative management is advocated for asymptomatic patients with palmar lipoma. For symptomatic patients, surgery is the treatment of choice, ideally performed by a hand surgeon. All lipomas in the upper extremities that are > 5 cm (i.e. giant lipomas) should ideally be surgically removed due to the potential increased risk of malignancy. Monobloc resection with careful dissection of neurovascular components is performed to avoid iatrogenic injury. Local recurrence is very rare and the incidence of recurrence correlates with the heterogeneity detected on MR imaging.

Other common, benign mass lesions in the palm include ganglion cysts, nerve sheath tumours, giant cell tumours (GCTs) of the tendon sheath, fibromas of the tendon sheath, arteriovenous malformations/haemangiomas and tenosynovitis. Ganglion cysts (Figs. 7a–c), one of the most common tumours, arise from adjacent tendons or joints and contain gelatinous fluid rich in hyaluronic acid and mucopolysaccharides. Their cause is not well-understood, although some patients may have a history of trauma. Patients may present with swelling or symptoms, depending on the mass effect on surrounding structures. Ganglion cysts occur more commonly in young women, especially around the dorsum of the wrist. On MR imaging, they appear as lobulated, septated lesions that are hypointense on T1-weighted and hyperintense on T2-weighted.
images. Often, there is an identifiable communicating tract with the joint or tendon sheath.

GCTs of the tendon sheath (Figs. 8a–c) are benign tumours that are histologically similar to pigmented villonodular synovitis. They are slow-growing and may present with painless or painful swelling in the third to fifth decades of life, with a slight female preponderance. Commonly, GCTs of the tendon sheath arise from the palmar tendons. The characteristic imaging features include a lobulated mass with internal low-signal-intensity foci on T1- and T2-weighted imaging and variable enhancement. \(^{(17)}\) The low-signal-intensity foci are better seen on gradient-echo sequences, which are more sensitive for susceptibility artefacts.

In the hand, nerve sheath tumours usually arise from deeper, larger nerves, particularly schwannomas. Neurofibromas, however, tend to arise from smaller cutaneous nerves. Both types of tumours present in young patients and are small, solitary and slow-growing. They may be painless or present with pain and/or paraesthesia. Most of them are isolated cases without history of neurofibromatosis. On imaging, a nerve sheath tumour (Figs. 9a–d) appears as a fusiform mass with tapered ends representing the entering and exiting nerve at the expected location of the nerve. It shows isointensity or slight hyperintensity to muscle on T1-weighted MR images and marked hyperintensity on T2-weighted MR images. Classically, neurofibromas demonstrate the target sign and enhancement. The target sign refers to peripheral hyperintensity and central hypointensity on T2-weighted images.

Arteriovenous malformations/haemangiomas (Figs. 10a–d) normally appear isointense on T1-weighted images, but may have
high signal foci due to fat and haemorrhage. They are usually heterogeneously hyperintense on T2-weighted images and foci of low signal may be seen on both T1- and T2-weighted MR images, likely due to phleboliths, flow voids or haemosiderin deposition. Avid serpentine or lattice-like contrast enhancement may be present. MR angiography can be complimentary to standard MR imaging. These patients, who are usually children or young adults, seek medical help due to localised pain or soft-tissue swelling. On examination, arteriovenous malformations/haemangiomas may be tender with normal or bluish overlying skin. Patients may have combined osseous, muscular or subcutaneous infiltration.\(^{1,18}\)

Tenosynovitis is inflammation of the synovial membrane, which surrounds the tendon sheath (Figs. 11a–c). The causes of tenosynovitis are infection, mechanical irritation and inflammation. Its clinical features include swelling, pain, surface redness and reduced movement of the hand, depending on the aetiology. On MR imaging, increased fluid within the tendon sheath may be seen, appearing as high signal on T2-weighted images and low-to-intermediate signal on T1-weighted images. Peritendinous thickening and enhancement with surrounding muscle and subcutaneous enhancement can also be noted.

In summary, though lipomas are commonly encountered clinically, deep palmar lipomas are relatively rare. The extent and depth of these deep palmar lipomas can be easily detected and characterised with the use of MR imaging. Understanding neurovascular and tendon relationships through MR imaging can help in surgical planning. Meticulous surgery is curative and provides relief for associated neuropathy.

**REFERENCES**

Question 1. Regarding the anatomical aspect of lipomas in the hand:
   a) Thick, deep central palmar fascia accounts for the central location of lipomas.
   b) Lipomas in Guyon’s canal affect the radial nerve.
   c) Lipomas in the carpal tunnel affect the ulnar nerve.
   d) Tinel’s sign is distal paraesthesia secondary to tapping on the tumour.

Question 2. Regarding the clinical course of lipomas in the hand:
   a) The incidence of lipomas in the hands and fingers is about 10%.
   b) Muscular atrophy and muscle paralysis are very common.
   c) Acute neuropathy due to nerve compression by interstitial haemorrhage may be seen.
   d) Giant lipomas are larger than 3.5 cm.

Question 3. Regarding imaging of lipomas in the palm:
   a) On plain radiography, lucency within soft-tissue swelling may be seen.
   b) On ultrasonography, they commonly appear as a hypoechoic mass with increased vascularity.
   c) Magnetic resonance (MR) imaging is recommended before biopsy as it helps in the selection of suspicious areas for biopsy.
   d) Tissue characterisation is superior on ultrasoundography as compared to MR imaging.

Question 4. Regarding MR imaging of lipomas in the hand:
   a) They appear hyperintense on both T1-weighted and T2-weighted images.
   b) They appear homogenously hyperintense on fat-suppressed sequences.
   c) Intramuscular lipomas are the most common subtype of palmar lipoma.
   d) Areas of incomplete fat suppression and contrast enhancement are suspicious for malignancy.

Question 5. Regarding differential diagnosis of lipomas in the hand:
   a) Ganglion cysts can have a communicating tract with the joint or tendon sheath.
   b) Giant cell tumours of the tendon sheath usually have high-signal foci on T1-weighted and T2-weighted MR imaging sequences.
   c) On MR imaging, nerve sheath tumours can have a fusiform appearance with tapered ends.
   d) Flow voids may be seen in arteriovenous malformations on MR imaging.

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