Clinical misdiagnosis of Morton’s neuroma: a case of early rheumatoid arthritis

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ABSTRACT
A fit and apparently healthy male patient presents with symptoms and clinical signs consistent with a Morton’s neuroma. Following excisional surgery, histopathology confirms the lesion as a rheumatoid nodule; this proves to be the presenting feature of rheumatoid arthritis in this patient.

This is a very unusual differential diagnosis, which should be considered during the assessment process and is, therefore, highly pertinent to clinicians.

CASE REPORT
A 56-year-old Caucasian male presented with moderate-to-severe bilateral forefoot pain that had been present for the past two years. The past medical history was unremarkable except for diffuse musculoskeletal pain in the cervical spine region for which the non-steroidal anti-inflammatory drug meloxicam (500mg bd) was prescribed.

Clinically, both on visual examination and palpation, there was a marked soft-tissue swelling affecting the left fourth metatarsal interspace. The swelling was particularly noticeable on weight bearing and caused the fourth and fifth toes to splay. Palpation of the swelling produced discomfort, and direct retrograde pressure applied from a plantar direction, just distal to the interspace, elicited sharp pain.

The history was one of pain developing during walking and becoming progressively worse with more activity. This pain was localised to the area of swelling and was also reported as radiating distally into the fourth and fifth toes. The size of the swelling and associated pain had increased gradually over the past two years, although the patient described both features as fluctuating. Examination of the right foot revealed similar clinical features at the same site, but these were much less obvious and of less concern to the patient. The history and clinical findings strongly suggested that the diagnosis was an intermetatarsal neuroma with associated bursa formation.1,2

MANAGEMENT
Previous treatment with foot orthoses utilising both valgus fillers and metatarsal pads to redistribute pressure had failed to provide relief. Considering the clinical history and symptomology a diagnosis of Morton’s neuroma was made. The patient was not keen to undergo local injection of glucocorticoid to the neuroma and elected to undergo neurectomy under local anaesthesia.

Pre-operative blood tests were unremarkable and excision of the suspected neuroma was undertaken as a day case under ankle-block anaesthesia, via a dorsal incision to the fourth metatarsal interspace. Recovery was uneventful. The soft tissue mass excised measured 3cm by 1cm, with a central cyst-like cavity containing a yellow-coloured viscous fluid with outer lobular fatty tissue. The macroscopic appearance of the lesion was not consistent with that of a neuroma and the specimen was sent for histological examination. The histopathology report revealed ‘the latter changes are reminiscent of rheumatoid nodule formation’. This finding was rather surprising, as at the time of surgery the only noteworthy reported medical history was chronic musculoskeletal pain in the region of the cervical spine.

During the post-operative period the patient rapidly developed many symptoms classically associated with Rheumatoid arthritis (RA).3 These included continued neck and shoulder pain weakness affecting his arms and hands (worse on the right side) swelling of the right hand and wrist area, and a soft, nodular swelling overlying the left elbow. Haematological analysis (outlined in Table 1) suggested a gross inflammatory state. Subsequent referral to a local consultant rheumatologist was arranged.

DISCUSSION
Clinical examination of both feet suggested that the soft tissue swelling in the fourth inter-metatarsal space was most likely to be a neuroma, with associated bursa formation. Morton’s neuroma (synonymously referred to as Morton’s metatarsalgia, Morton’s neuralgia or plantar digital neuritis) was named after Thomas G Morton, who described this pathology affecting the 4th intermetatarsal space.5

Classically this pathology is described as a fusiform swelling of the third and fourth plantar digital nerves.6 However, there remains debate as to which intermetatarsal space is most commonly affected, with some authors concluding that Morton’s neuroma does not occur in the 4th-5th metatarsal space.7 Histologically endoneural oedema, exceptional fibrosis and demyelination are said to be diagnostic characteristics of an intermetatarsal neuroma.8

Histopathology in this case reported findings reminiscent of a rheumatoid nodule. Excluding anaemia and constitutional symptoms, rheumatoid nodules are the most common extra-articular
manifestations of RA. Nodules usually develop with an insidious onset over areas of pressure or in tendons or tendon sheaths. Although not usually painful, nodules can become symptomatic if subjected to pressure and stress.

In patients with RA, the general incidence of subcutaneous nodules is 20–40%, and rheumatoid nodules are reported to be relatively common in the foot. Rheumatoid nodules may increase or decrease in size, resolve, reappear or persist indefinitely. With repeated trauma rheumatoid nodules may ulcerate and form a portal for infection. The treatment of a symptomatic subcutaneous nodule, therefore, is usually excision, though recurrence is a possibility.

Rheumatoid nodules are said to form the hallmark of seropositive rheumatoid disease. They do form part of the diagnostic criteria for RA, and, as such, nodules constitute a useful diagnostic or prognostic disease marker. However, it is more usual for extra-articular pathologies such as nodules to develop much later in the course of the disease and only rarely are they the first presenting feature. Their usefulness as disease markers is thus limited. It is more commonly reported in RA that inflammation affecting the inter-metatarsal bursa is sufficient to cause stretching or compression of the inter-digital nerves, producing symptoms similar to those of Morton’s neuroma.

The more usual differential diagnoses for Morton’s neuroma include a range of conditions including synovitis, bursitis and Freiberg’s disease, as well as neurological causes. However, consideration of rheumatoid nodules as a differential diagnosis does not receive wide attention in the contemporary podiatric literature. Recently Emery et al have developed guidelines for the referral of patients in the event of clinical suspicion of RA. The typical presentation in this case did not lead to the suspicion that the patient may have RA.

The diagnosis of Morton’s neuroma is generally made through history and clinical examination. More recently ultrasonography has been employed to aid the diagnosis of neuromas. Recent reports have indicated sensitivity for the diagnosis of neuromas using ultrasonography of 94–100% and a specificity of 80–100%, although the authors point out that there is little consistency in the methodologies employed.

In retrospect, the use of ultrasound imaging in this case would almost certainly have confirmed the presence of a soft tissue mass, although whether a clear differentiation (using this investigative method) between a rheumatoid nodule or Morton’s neuroma can be accurately identified is more difficult to determine. Whilst there are guidelines for ultrasound imaging in rheumatology, there remains a paucity of literature outlining the nature and/or prevalence of foot pathologies in early RA.

X-ray remains an important part of the diagnostic criteria for patients with RA. It has been repeatedly reported that patients with RA demonstrate periarticular erosions in the feet earlier than in the hand. However, all patients in these data series already had a positive diagnosis of RA.

In this case the clinical signs and symptoms were strongly suggestive of Morton’s neuroma. In the absence of other features suggestive of early RA, reported by Emery et al, it was felt that plain film radiography was not appropriate. In cases where there are inconclusive clinical findings, ultrasound or magnetic resonance imaging of suspected neuromas might be considered to help confirm the diagnosis.

CONCLUSION

This case was noteworthy, as the first presentation of RA was a subcutaneous nodule in the foot. Symptoms arising from the nodule mimicked those of an inter-metatarsal neuroma and in the absence of other clinical features was incorrectly diagnosed as this lesion. Imaging modalities such as ultrasonography could be considered as a useful aid to diagnosis.

Pain and inflammation of the small joints in the hands and feet more commonly typify early RA. The case also highlights the importance of recognising the early manifestations of systemic inflammatory arthropathies that may present within the foot.

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REFERENCES


Table 1. Haematological analysis.

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<thead>
<tr>
<th>Results of Haematological Analysis</th>
<th>Normal values</th>
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<tbody>
<tr>
<td>ESR 67 mm/hour</td>
<td>1-3 mm/hour</td>
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<tr>
<td>CRP 80 µg/ml</td>
<td>&lt;0.8 mg/dL</td>
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<tr>
<td>WBC 13.5 x 10⁹/L</td>
<td>4.0 - 10.0 x 10⁹/L</td>
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<tr>
<td>Neutrophil 9.2 x 10⁹/L</td>
<td>2.5 - 7.5 x 10⁹/L</td>
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