A 73 year old white woman was admitted to our hospital in May 2003 complaining of lower abdominal pain for 1 month. On admission, the gynaecological examination disclosed a large, non-tender lower abdominal mass of 7 cm in diameter, which was highly suspicious of ovarian cancer with peritoneal infiltration in the computed tomography scan. In accordance with these findings, serum levels of CA 12-5 were about 3000 U/ml (normal range <35 U/ml).

Therefore, we performed an exploratory laparotomy. The frozen section of the adnexal mass showed a serous papillary carcinoma of fallopian tube origin. Consecutively, a complete staging with hysterectomy, bilateral salpingo-ophorectomy, omentectomy, and resection of the descending part of the colon due to tumour infiltration was performed.

Recovery was complicated owing to prolonged ileus-like symptoms. First line chemotherapy with topotecan in combination with carboplatin under study conditions (prospective, open label, phase II study) was started 3 weeks after the operation.

Before the diagnosis of carcinoma of the fallopian tube, the patient reported a progressive painful swelling and stiffness of both hands, especially the palms, which were noted by the patient to have thickened progressively since December 2002, with no improvement after initial corticosteroid treatment. Apart from generalised arthrosis, she denied any previous joint diseases, Raynaud’s phenomenon, skin tightening, or trauma.

Physical examination on admission was remarkable for symmetrical swollen hands and fingers, painful on active and passive movement, and palmar fascial thickening with erythema (fig 1). The fingers of the patient showed flexion contractures; making a fist was impossible. No evidence of skin sclerosis or arthritis in other locations was found.

The patient’s symptoms were orally treated with a cyclo-oxygenase(COX)-2 inhibitor, a transdermal delivery system for buprenorphine, and local ointments with diclofenac. Beside an increased acute phase reaction (postoperatively), the immunological laboratory investigation failed to show any specific abnormalities (rheumatoid factor, antinuclear antibodies, antistreptolysin titre, antineutrophil cytoplasmic antibodies, complement CH 50 and C3d). Radiography of both hands showed a mild arthrosis in the joints, but no signs of acute arthritis.

After two cycles of antineoplastic treatment the patient’s paraneoplastic symptoms currently show a good response: the fasciitis and acute arthritis have gradually improved, but the contractures in both hands persist despite extensive physiotherapy.

DISCUSSION

Paraneoplastic syndromes affect a variety of organ systems and are often an initial sign of occult malignancies. About 30 cases have been reported describing palmar fasciitis and polyarthritis (PFPA)-like changes of the hands in association with malignant tumours. PFPA has been reported to be associated with different neoplasms—for example, ovarian cancer, carcinoma of the breast, carcinoma of the prostate, and gastric tumours, preceding or accompanying the diagnosis of malignancy.

In our case we present a patient demonstrating a severe progressive deforming rheumatic disease affecting the palms and fingers of both hands, with the onset of the corresponding symptoms 5 months before the diagnosis of a malignant neoplasm was made.

To our knowledge this is the first report of the association of PFPA as a paraneoplastic syndrome with a carcinoma of the fallopian tube.

The underlying immunological mechanisms have not been defined so far and may result from activation of certain factors with profibrotic activities—for example, transforming growth factor β or connective tissue growth factor. As long as the pathomechanisms of PFPA are unclear, the only effective therapy is cancer treatment, which may lead to arrest or even complete resolution of the rheumatic disease.

The characteristic hand deformities of PFPA, recently described with the illustrative term of woody hands, should alert the clinician to search for an underlying malignant disease.

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Mycobacterium chelonae infections are uncommon and have not been reported as a complication of Raynaud’s phenomenon previously. We describe a patient who responded well to treatment.

CASE REPORT
A 21 year old female telephonist was referred with a 1 week history of swollen right index and middle fingers. There was no history of trauma but she had had Raynaud’s phenomenon since the age of 12. There was no history of recent travel abroad or of unusual hobbies.

On examination she was apyrexial, she had diffuse swelling but no discoloration of the right index and middle fingers. She had no blisters or ulceration and the fingers were not tender to touch. Nine days later, despite flucloxacillin orally and a non-steroidal anti-inflammatory agent, all her fingers were cold and bluish, and the right index and right middle fingers remained diffusely swollen. She was admitted for intravenous prostacyclin because of worsening of her Raynaud’s phenomenon.

Her baseline investigations were normal, including erythrocyte sedimentation rate and C reactive protein. A hand x ray examination showed no bony abnormality. Blood cultures were negative. Her immunoglobulin levels were normal, and cryoglobulins were not detected. Her autoantibody profile was negative except for raised IgG cardiolipin antibodies at 46 GPLU (reference range 0–13).

As she finished the 72 hour intravenous prostacyclin infusion, a pustule appeared on the pulp of the right index finger, and intravenous ceftriaxone was started. Two days later, a second pustule appeared on the other finger (fig 1). Microscopy of a needle aspiration showed acid fast bacilli. Antibiotics were changed to clarithromycin 500 mg twice daily and ciprofloxacin 750 mg twice daily. Both lesions were incised and drained, and histology of the material showed an abscess wall with no granulomata.

M chelonae was cultured from pus and was susceptible in vitro to azithromycin and clarithromycin, but resistant to ciprofloxacin. As her lesions were healing at the time these results became available (14 days later) her antibiotics were not changed.

DISCUSSION
M chelonae is associated with a variety of infections. Cutaneous lesions occur secondary to wound infections after surgery, accidental trauma, or needle injections.1 2 M chelonae isolates are susceptible in vitro to clarithromycin, but generally resistant to ciprofloxacin. The use of at least two drugs (one of which should be clarithromycin) is recommended for treatment of M chelonae infection, as the emergence of resistance is a risk associated with monotherapy.1 3 Treatment is usually given for 4–6 months, but our patient responded to combination therapy within weeks. A case has been reported of postoperative infection at a donor vein graft site that healed after 2 months of clarithromycin treatment, combined with heat treatment.1 Thus shorter courses of treatment may be appropriate in non-immunosuppressed patients.

Isolates of M chelonae have optimal growth at 28–30°C,6 unlike the standard 35°C for most organisms. Our patient had severe Raynaud’s disease before the infection, so the coldness of her hands may have promoted the growth of this unusual organism. We are not aware of any previous reports of...
Progression of lupus nephritis during treatment with mycophenolate mofetil

K Ahmadi-Simab, P Lamprecht, W L Gross

M chelonae infection in a patient with Raynaud’s phenomenon. A significant portion of the 100 patients with M chelonae infections reported by Wallace et al. were receiving corticosteroids or immunosuppressant drugs, or both. In their series 35 patients had a localised cutaneous infection and only five had no history of trauma, of whom three had an underlying immunosuppressive condition.

Our patient had no history of trauma, she was immunocompetent, but had a history of Raynaud’s phenomenon. Because there were no systemic features during the acute phase of her illness, an atypical infection was suspected and acid fast bacilli were specifically looked for. In conclusion, unusual presentations have unusual explanations and should raise suspicion of atypical infections.

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Lupus nephritis (LN) determines the prognosis of systemic lupus erythematosus. The standard treatment for the proliferative forms (focal and diffuse proliferative LN, WHO classes III and IV) is intravenous pulse cyclophosphamide in combination with oral prednisone (Austin scheme).1 2 Recently mycophenolate mofetil (MMF) has been shown to be as effective as standard pulse cyclophosphamide for the induction of remission in proliferative LN (n = 21 in each treatment group). Side effects of MMF were fewer than with pulse cyclophosphamide.3 We report on a patient with deteriorating renal function and switch of LN class despite MMF treatment.

CASE REPORT
Systemic lupus erythematosus was diagnosed in a 63 year old white patient with non-erosive polyarthritis, photosensitivity, malar rash, nephritic sediment (5000 white cells/µl, dysmorphic erythrocytes, non-selective glomerular proteinuria 2.26 g/day), antinuclear antibodies, and anti dsDNA antibodies. A renal biopsy disclosed non-proliferative LN with mesangial hypercellularity (WHO class type IIb). The patient had experienced adverse effects with azathioprine and methotrexate previously. Thus, MMF (2.0 g/day p.o., CellCept, Roche Pharmaceuticals) and prednisolone (1.0 mg/ day p.o. with subsequent tapering) were given. However, renal function deteriorated within 3 months, with declining creatinine clearance (65 ml/min to 49.1 ml/min), increasing serum creatinine (112 µmol/l to 155 µmol/l), and persistent nephritic sediment and proteinuria. A second renal biopsy disclosed a switch to focal proliferative LN (WHO class III). Treatment with MMF was discontinued and intravenous cyclophosphamide pulses were given, resulting in stabilised renal function and reduced proteinuria (0.6 g/day) after six cyclophosphamide pulses.

DISCUSSION
In this case MMF was given in non-proliferative mesangial LN (WHO class IIb), characterised by a better prognosis than proliferative LN. However, renal function deteriorated owing to progression of the LN to focal proliferative LN (WHO class III). Although the presence of both LN types at onset and late effects of MMF on renal function after switching to pulse cyclophosphamide cannot be ruled out, the course with deteriorating renal function during MMF treatment suggests
inefficacy of MMF and a concomitant switch of LN class in this case. A personal communication on a higher relapse rate with MMF than with cyclophosphamide pulse for proliferative LN during follow up of Chan’s study\(^4\) (46% vs 17%; \(p = 0.019\)) has been reported in a review recently.\(^4\)

We conclude that further studies on larger patient groups are needed to establish the place of MMF in the treatment of LN and to develop therapeutic algorithms, including early intervention in non-responders.

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**Assessment of the clinically relevant change in pain for patients with sciatica**

B Giraudieu, S Rozenberg, J-P Valat

The visual analogue scale (VAS) is widely used for pain assessment. However, the minimum clinically relevant change remains a debated question, even with regard to the method of assessment. On the one hand, considering intra-individual change in acute pain, Todd et al suggested a crude change of 13 mm (for a 100 mm VAS) to discriminate between reporting a little less or a little more pain.\(^1\) In the study of Farrar et al the best cut off points were estimated at −20 mm (for a crude change) or −33% (for a relative change) to discriminate between patients who require a rescue opioid dose or not.\(^2\) Similar results were also derived when considering patient self appreciation of improvement in chronic pain.\(^3\) On the other hand, Redelmeier et al suggested focusing on interindividual change to assess a minimal important difference.\(^4\) Wells et al thus observed that when a patient feels “somewhat worse” than another patient who faces him, then there is a 16% relative difference in their mean answer to a pain questionnaire.\(^5\)

We performed an ancillary study to a clinical trial in order to estimate the minimum clinically relevant change in pain for patients with sciatica. The main study was a randomised double blind trial conducted to assess the effectiveness of epidural corticosteroid injections in patients with sciatica, presumably due to a herniated nucleus pulposus, for 15–180 days.\(^6\) We thus administered three epidural injections at 48 hour intervals with 2 ml prednisolone acetate (50 mg) or 2 ml isotonic saline. Self evaluation was the primary outcome, measured at day 20 on a four item scale (recovery, important improvement, poor improvement, or worse). The first two items were then pooled, thus defining a treatment success, while the last two items defined a treatment failure. As a secondary outcome we assessed the severity of pain on a VAS both at day 0 and day 20. Eighty five patients were included in the main study. However, in the present work, analyses were performed on a subsample of 75 patients for whom we had both the self evaluation at day 20 and the two pain assessments. The subsample comprised 45 men and 30 women with a mean (SD) age of 41 (11) years. Those patients had had pain for a mean (SD) duration of 49 (41) days and their mean baseline pain was 58 (16). For 39 of those 75 patients, treatment was a success. We performed analyses without taking into account the randomisation arm: there is indeed no foundation for a different relationship between self evaluation and pain assessment according to a patient’s treatment. For each patient, we calculated both the crude and relative changes in pain assessment and plotted receiver operating characteristic curves (fig 1).

**Figure 1** Receiver operating characteristic curves for crude and relative changes in pain assessment v self evaluation. Self evaluation is defined as “treatment success” or “treatment failure” at day 20. Change in pain assessment is defined as the crude or relative change in pain VAS assessment between day 0 and day 20.

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Optimal cut off points, which thus offer the best compromise between sensitivity and specificity, are estimated at $30\text{ mm}$ for the crude change and $50\%$ for the relative change. These values are higher than those estimated by Todd et al.\textsuperscript{1} and Farrar et al.\textsuperscript{2,3} Such discrepancies may be due to a difference in the “gold standard”. They may also be related to the length of the period of observation because we focused on a 3 week period.

In conclusion, the values we estimated may be seen as the clinically relevant long term changes in pain assessment for patients with sciatica. Such findings, however, have to be confirmed by further studies planned with this specific aim because patients included in a randomised trial may not be representative of the general population of patients with sciatica.

Table 1 Performance indexes associated with different cut off values of (A) crude and (B) relative change in pain VAS assessment

| (A) Cut off value for crude change in pain VAS between day 0 and day 20 |
|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|
|                            | -40 mm                      | -30 mm                      | -20 mm                      | -10 mm                      |
| Sensitivity                | 56.4 (39.6 to 72.2)         | 76.9 (60.7 to 88.9)         | 87.2 (72.6 to 95.7)         | 92.3 (79.1 to 98.4)         |
| Specificity                | 91.7 (77.5 to 98.2)         | 80.6 (64.0 to 91.8)         | 69.4 (51.9 to 83.7)         | 58.3 (40.8 to 74.5)         |
| Accuracy*                  | 73.3 (61.9 to 82.9)         | 78.7 (67.7 to 87.3)         | 78.7 (67.7 to 87.3)         | 76.0 (64.7 to 85.1)         |

| (B) Cut off value for relative change in pain VAS between day 0 and day 20 |
|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|
|                            | -60%                        | -50%                        | -40%                        | -33%                        |
| Sensitivity                | 76.9 (60.7 to 88.9)         | 87.2 (72.6 to 95.7)         | 87.2 (72.6 to 95.7)         | 89.7 (75.8 to 97.1)         |
| Specificity                | 88.9 (73.9 to 96.9)         | 80.6 (64.0 to 91.8)         | 69.4 (51.9 to 83.7)         | 69.4 (40.8 to 74.5)         |
| Accuracy*                  | 82.7 (72.2 to 90.4)         | 84.0 (73.7 to 91.4)         | 78.7 (67.7 to 87.3)         | 80.0 (69.2 to 88.4)         |

Results are expressed as point estimates (95% confidence intervals).

*Accuracy is defined as the proportion of patients for whom treatment was correctly classified as success or failure.

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