CASE REPORTS

Systemic lupus erythematosus presenting as pneumococcal septicaemia and septic arthritis

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Abstract
A 50 year old woman presented with pneumococcal septicaemia, septic arthritis, and a lobar pneumonia and was subsequently diagnosed as having systemic lupus erythematosus. The blood film and splenic $^{99m}$Tc sulphur colloid uptake were normal, although selective functional hyposplenism was shown by the impaired clearance of immunoglobulin coated erythrocytes. Systemic lupus erythematosus presenting with fulminating pneumococcal sepsis in the presence of selective defects in spleen function is previously unreported.

Hyposplenism, with its characteristic blood film appearances and absence of splenic activity on $^{99m}$Tc sulphur colloid scan, has occasionally been reported in patients with systemic lupus erythematosus (SLE). The spleen plays a crucial part in the elimination of certain microorganisms from the blood, and patients who have undergone splenectomy are more susceptible to overwhelming infection, particularly by the pneumococcus. Hyposplenism in patients with SLE also increases their susceptibility to overwhelming infection with this organism.

We report a case in which the patient presented acutely with a septic arthritis, a cavitating lobar pneumonia, and pneumococcal septicaemia. She was subsequently diagnosed as having SLE. Both the blood film and $^{99m}$Tc colloid scan were normal, but more selective studies of splenic function showed that she had a significant splenic macrophage Fc receptor defect.

Case report
A 50 year old white woman presented initially with an eight month history of epigastric discomfort and increasing exertional dyspnoea. In the preceding 12 months she had been taking piroxicam for pain in the hands and shoulders. During the preceding five years she had experienced three episodes of respiratory tract infection, which had required treatment with oral antibiotics. She had been a smoker of 20 cigarettes a day for many years until five months previously.

On examination she had synovitis of the metacarpophalangeal joints and the extensor tendon sheath of the left wrist, bilateral tender-ness of the metatarsophalangeal joints, and slight restriction in the movement of both shoulders. Laboratory investigations showed a normochromic normocytic anaemia (haemoglobin 95 g/l), a white cell count of 4.2 x 10⁹/l, a normal blood film, and an erythrocyte sedimentation rate of 65 mm/h. The rheumatoid arthritis latex test was weakly positive. Lung function tests showed a restrictive defect with decreased static lung volumes (forced expiratory volume in one second 1.22 litre and forced vital capacity 1.65 litre—both 40% of predicted values, a total lung capacity 3.4 litre (84% predicted), and a carbon monoxide transfer factor of 17 ml/min/mmHg (normal range 18–32 ml/min/mmHg)).

Four weeks later she was admitted following the acute onset of swelling and pain in the right knee and right calf. She had become increasingly breathless over the previous two weeks and had developed a cough productive of purulent sputum. On examination, despite the clinical signs of a left upper lobe pneumonia, she had no fever. There was a large effusion in the left knee and the right calf was swollen, red, and tender. The chest radiograph confirmed the lobar pneumonia. Investigations showed a haemoglobin of 85 g/l, a white cell count of 3.8 x 10⁹/l, and an erythrocyte sedimentation rate of 125 mm/h. All three blood cultures grew Streptococcus pneumoniae and this organism was also present in the fluid aspirated from the right knee. Gram positive cocci were identified in the sputum, but as this sample was obtained after starting treatment with antibiotics (benzylpenicillin 3 MU intravenously six hourly) no organism was grown. During the next 48 hours the right calf became increasingly painful and swollen and an ultrasound scan showed a large but localised collection of fluid in the medial compartment of the calf. This was aspirated, yielding 6 ml of seropurulent fluid, but no organisms were grown.

During the next seven days she improved clinically, but cavity developed within the left upper lobe. Metronidazole was added to her treatment to cover possible anaerobic superinfection. Over the next three weeks her joint and respiratory infections resolved completely. She had no fever throughout, and the white cell count never rose above 10 x 10⁹/l and was often below 4 x 10⁹/l.

The persistent arthralgia and synovitis, and the lung function abnormalities in association with overwhelming infection with an encap-
sulated organism, prompted further investigations. The antinuclear factor (homogeneous nuclear staining) was positive at a titre of 1/128, high levels of antibodies to double stranded DNA were also identified in her sera on five separate occasions (>120 units/ml (normal range <30 units/ml)), and antcardiolipin antibodies (IgG and IgM classes) were present in high titre. High levels of circulating immune complexes were detected using a monoclonal rheumatoid factor binding assay (90% binding (normal range <15% binding)) and the complement concentrations were normal. There was a polyclonal increase in the immunoglobulin concentrations.

Six weeks after her acute admission to hospital, and before any specific treatment for her SLE had been started, splenic function was assessed by a $^{99m}$Tc colloid scan. Both splenic uptake and size were normal. Further tests of splenic function were carried out by determining the clearance of N-ethylmaleimide treated autologous red blood cells and IgG coated erythrocytes from her circulation. These results showed normal N-ethylmaleimide clearance but a significant delay in the clearance of the IgG coated cells, the $t_1$ being 163 minutes (figure).

Subsequently she remained breathless despite complete clinical and radiological resolution of the pneumonia, and repeated lung function tests showed further deterioration in the forced vital capacity and total lung capacity coupled with a 30% decrease in carbon monoxide transfer. A transbronchial biopsy showed interstitial pulmonary fibrosis with very little cellular infiltration, and a bronchopulmonary lavage produced fluid with a very low white cell count. She was treated with 40 mg of prednisolone and within four weeks there had been a marked clinical improvement with a corresponding increase in static lung volumes and carbon monoxide transfer. Six weeks after the introduction of prednisolone, cyclophosphamide 100 mg daily was introduced, and over the next two years both drugs were gradually withdrawn without any significant change in lung function. Three years after presentation the patient remains well and all treatment has been discontinued.

Discussion

Our patient, with a non-erosive arthropathy, haematological abnormalities, a positive antinuclear factor, and high levels of antibodies to double stranded DNA, satisfied four of the revised American Rheumatism Association criteria for the diagnosis of SLE. Although infection occurs quite often in patients with SLE, it is rarely the presenting feature of the disease. The presentation of our patient with pneumococcal septicemia, pneumococcal joint infection, and a cavitating lobar pneumonia is very unusual. A number of conditions, which include chronic chest disease, alcoholism, haematological malignancy, cirrhosis, sickle cell disease, and splenectomy, are known to predispose to pneumococcal bacteraemia, but we were unable to identify any of these conditions in our patient. Septic arthritis has been well reported in SLE with a variety of organisms, but the joint is a very uncommon site for pneumococcal infection. In a study of over 500 episodes of pneumococcal infection in 494 patients pneumococci were isolated from the synovial fluid in only three. Twelve of the 494 patients in that study had undergone a surgical splenectomy, while SLE was considered to be the underlying condition in only five patients. The site of the infection and splenic status of these individuals were not described.

Hypersplenism, characterised by typical blood film appearances, including Howell-Jolly bodies, spherocytes, target cells, and Pappenheimer bodies, and the absence of splenic activity on a $^{99m}$Tc sulphur colloid scan, occurs rarely in SLE. The association was first described in 1980 by Dillon et al in a young North American Indian woman with SLE. Eight years after the demonstration of hypersplenism in the above patient she died from overwhelming pneumococcal infection. At necropsy the spleen was small and fibrotic. Dillon and coworkers have subsequently described two more patients, out of 70 with SLE, with blood film and scan appearances which were indicative of hypersplenism. The mechanisms which lead to the development of hypersplenism are controversial and not clearly understood.

Our patient had a normal blood film and a normal splenic uptake of $^{99m}$Tc sulphur colloid. We were able to show, however, a marked defect in the rate of clearance of IgG coated erythrocytes from her circulation, a process which is dependent on the recognition of
spleen plays a crucial part in the induction of the antibody response to pneumococcal capsular polysaccharide and other thymus independent antigens. The antibody response to thymus independent antigens in patients who have undergone splenectomy is profoundly impaired, whereas the secondary responses in splenectomised individuals, primed with antigen before the removal of the spleen, are normal.\(^{21}\) Although we cannot exclude the possibility that the pneumococcal antibody response in this patient was impaired, we think it unlikely given the normal clearance function of the spleen and the likelihood of natural exposure and priming of our patient to pneumococcal polysaccharides.

In conclusion, our patient presented with overwhelming pneumococcal infection and was subsequently diagnosed as having SLE. Hypopplenism, assessed on the blood film appearance and the uptake of sulphur colloid, was not present, though there was a delay in the clearance of IgG coated erythrocytes by the spleen. This defect is unlikely to be responsible for her increased susceptibility, but an associated C3b receptor defect might have impaired the clearance of pneumococci from her circulation and facilitated the development of overwhelming infection.

REFERENCES