CASE REPORTS

Case 1. In March 1998, a 2.5-year-old Caucasian girl presented with an 8 month history of swelling of one knee following minor trauma. She had mild pain, morning stiffness, and swelling that waxed and waned but never resolved completely. She limped continuously and developed a fixed flexion deformity. There was no history of fever, cough, or weight loss. There was no history of travel to an endemic TB area and no contact with anyone known to have TB. She had not received bacillus Calmette-Guérin (BCG). When seen in our clinic, she was well grown (height 50th percentile and weight 25th percentile) and the general physical examination was entirely normal. Her left knee showed a large effusion and 10° fixed flexion deformity with pain at the end range. There was no skin erythema or bony tenderness. Laboratory investigations including a test for antinuclear antibodies (ANA) were normal apart from an elevated erythrocyte sedimentation rate (ESR) (53 mm/h). Radiographs of the knee showed a large effusion with soft tissue swelling, but no osteopenia or joint space loss. She was given naproxen for the presumptive diagnosis of oligoarticular onset JRA.

One month later, the naproxen was discontinued because she developed a rash on her legs and there was no improvement in her symptoms. She had considerable pain and stiffness, and she was prescribed prednisone 5 mg bid. Because of the history of minor trauma prior to onset and because of the poor response to initial treatment, a magnetic resonance image (MRI) was performed to look for other causes for the knee swelling. The MRI showed nodular synovial thickening with large effusions and numerous “loose bodies.” Gram stain of synovial fluid (SF) showed no organisms and the culture was negative after 48 h of incubation. In spite of intraarticular triamcinolone hexacetonide therapy on 2 occasions, 2 months apart, and the addition of prednisone and tolmetin sodium, there was no improvement of her symptoms. Arthroscopy and synovectomy yielded straw colored fluid with a large number of yellowish “rice bodies” and villous synovitis. The histologic examination showed noncaseating granulomatous synovitis. The culture of synovial tissue grew M. tuberculosis sensitive to all the usual agents. A PPD skin test was reactive with 12 mm induration. Her chest radiograph was normal. Both parents and her 3 siblings had nonreactive PPD skin test.

She was started on triple chemotherapy (INH, rifampin and pyrazinamide). Pyrazinamide was discontinued after 3 months and the plan was to continue the other anti-TB medications for one year. She improved significantly. Three months after starting the TB treatment there were no joint effusions, and radiograph of the knee showed no bony lesions.

Case 2. In June 1998 a 6-year-old boy who had recently immigrated with his family from Somalia presented with an 11 month history of pain and swelling of his left knee that was preceded by minor trauma. He complained of knee pain that was worse at night and morning stiffness lasting 30 min daily. He was seen by several physicians including the family doctor and an orthopedic surgeon, and had started taking tolmetin sodium, and although this improved his symptoms, it was discontinued because of abdominal pains. Although TB...
is common in Somalia, he was not known to have had contact with anyone with the disease, and he had had BCG.

On evaluation in the pediatric rheumatology clinic, he was small (height 5th percentile, weight 3rd percentile). His chest was clear and he had no lymphadenopathy. The left knee showed a large effusion and 15° flexion deformity. The left leg was 1 cm longer than the right. His laboratory investigations including ANA were within normal limits apart from an elevated ESR (45 mm/h). Radiographs of the left knee showed a large effusion and soft tissue swelling with no osteopenia or joint space loss.

He was given naproxen for the presumptive diagnosis of monoarticular onset JRA. One month later, he had an arthrocentesis, with removal of 30 cc of clear yellow fluid from left knee, followed by intraarticular injection of 20 mg triamcinolone hexacetonide and serial casting. There was minimal improvement and MRI showed lobulated thickened synovium with large effusions and normal bone. Despite a second intraarticular dose of triamcinolone hexacetonide 3 months after the first one, his knee effusion and deformity persisted. A second MRI showed similar findings to the first study. Subsequently, he had an arthroscopy and the synovial pathology showed caseating granulomatous synovitis and \textit{M. tuberculosis} was isolated. He had a strong PPD reaction with 17 mm induration and normal chest radiograph.

No one in the family had clinical evidence of TB, but all had a strong PPD reaction. The patient was started on combination anti-TB chemotherapy to be continued for one year. His parents and siblings were started on INH. Followup radiograph of the knee one month after anti-TB therapy was started showed no evidence of bony lesion. Significant improvement in pain, swelling, and range of motion of the affected knee were noted 2 months after commencing TB therapy.

**DISCUSSION**

It is estimated that 1.3 million children in the world develop disease caused by \textit{M. tuberculosis} every year and the annual risk of infection with this organism is 1-2% in the developing world\textsuperscript{5}. In some areas in the United States the number of children who developed tuberculosis during the period 1987-1993 increased by 40%\textsuperscript{6}. The number of children in the United States infected with tubercle bacillus is unknown because the infection is reported in only 4 states\textsuperscript{7}. Over recent years, TB has not declined in Canada as expected and the increasing incidence among children is of concern. The proportion of nonrespiratory cases increased from 18% in 1980 to 25% in 1995\textsuperscript{8}, and 20% of these involved the musculoskeletal system\textsuperscript{9}. The resurgence of TB is attributed to several factors: immigration from endemic TB areas, the rise in the number of people with immune deficiency secondary to human immune deficiency virus infection or chemotherapy, and the increase in the aging population\textsuperscript{2,6}. Generally, children acquired the infection usually from an infectious adult family member. One of our patients was a recent immigrant from an endemic TB area, but the other patient had no identifiable risk factors.

Tuberculous arthritis most frequently causes a monoarthritis, with predilection for weight bearing joints\textsuperscript{1,10}, as in our patients. Delayed diagnosis of TB arthritis is not unusual\textsuperscript{1}, particularly in areas where TB is not endemic. Clinically, TB arthritis may cause pain, stiffness, effusion, and low grade fever. The usual insidious chronic course of TB arthritis and the SF characteristics may be indistinguishable from other forms of chronic arthritis\textsuperscript{10,11}. SF is inflammatory; white cell count is variable but usually ranges between 10,000 and 20,000 cells/mm\textsuperscript{3}. Polymorphonuclear cells predominate, but the count may vary from 10 to 99%. Protein is > 3.5 g/dl and the glucose is normal\textsuperscript{12}. Radiologically, soft tissue swelling and periarticular osteopenia are the early abnormalities, but progression to blurring of the subchondral bone surface and marginal erosions with joint space narrowing and frank destruction of bone may occur in the late stages of the infection\textsuperscript{12,13}. Extrapulmonary infections usually result from lymphohematogenous dissemination at the time of primary infection\textsuperscript{10,13,14}. In most children with TB arthritis (including both our cases) there were no pulmonary symptoms and the chest radiographs were normal\textsuperscript{10}. Prior BCG vaccination is not a contraindication to tuberculin testing and, in general, a reactive area of > 10 mm in a BCG vaccinated child probably indicates infection and necessitates further diagnostic evaluation and preventive chemotherapy\textsuperscript{2}.

Two other cases of TB arthritis in children were seen in our hospital: a 15-year-old Asian girl who is a recent immigrant presented with a 5 month history of left hip pain. A radiograph of her hip showed loss of articular space and changes at proximal femur suggestive of osteomyelitis. Large hilar lymph nodes with calcification were seen on her chest radiograph. Her PPD skin test was strongly positive. She was started on therapy for the presumptive diagnosis of TB. She is not living in the country now and no further followup is possible. The 4th patient was a 2-year-old North American Indian boy with Poncelet’s disease who presented with increasing back pain and sterile effusions of the knee and ankle, who was found to have active vertebral tuberculosis and a perivertebral abscess\textsuperscript{15}.

Treatment of bone and joint tuberculosis requires prolonged combination therapy, initially with isoniazide, rifampin, and pyrazinamide for the first 2 months, followed by isoniazide and rifampin administered once a day or twice a week under direct observation for a period of 9–12 months\textsuperscript{5,16}. Shorter courses of chemotherapy (6 month regimens) are associated with higher failure rate, especially if surgical intervention has not been undertaken\textsuperscript{17}.

Arthritis with TB may be a result of direct infection of the joint or a “reactive” arthritis secondary to visceral disease (i.e., Poncelet’s disease). Arguably much of the joint damage in the former situation is mediated by activated lymphocytes, and this may be modified by steroids. Adjuvant treatment with corticosteroids is controversial, except in the treatment of tuberculous meningitis. It may also be considered in the treatment of miliary disease, endobronchial disease, and pleural and pericardial effusions\textsuperscript{16}. There are no studies that evaluate the role of systemic or intraarticular corticosteroid in tuberculous arthritis. In both our patients, intraarticular corticosteroid was used before the diagnosis of TB was established. Despite the considerable delay in the diagnosis (18 months), there was no clinical or radiological evidence suggestive of disease extension or complication and the outcome in both cases, once chemotherapy was started, was good. The role of intraarticular corticosteroids in preventing or exacerbating damage in TB arthritis remains unclear.
In an era of increasing incidence of tuberculosis in children, especially the extrapulmonary form, tuberculous arthritis should be considered in the differential diagnosis of monoarthritis. A history of TB exposure should be sought; and it may be that even in the absence of risk factors PPD application should be considered. In a patient with presumed JRA with a monoarticular course, TB should be considered if there is an inadequate response to usual treatment. Our 2 patients were fortunate not to have destructive joint disease, in spite of a delayed diagnosis.

REFERENCES