HLA-B27 positive juvenile arthritis with cardiac involvement preceding sacroiliac joint changes

S J Lee, H Y Im, W Schueller

Abstract
While cardiovascular disease develops in up to 50% of adult patients with ankylosing spondylitis, it is very uncommon in its juvenile counterpart. Regarding the early stage of the disease, before onset of sacroiliac joint changes, only two cases with aortic incompetence have been published while reports of mitral valve involvement are not available. A 13 year old boy is described with an HLA-B27 positive asymmetric oligoarthritis and enthesitis, without back pain or radiographic evidence of sacroilitis. Echocardiography showed an echogenic structure measuring 8 × 11 × 20 mm at the fibrous continuity between the aortic and mitral valves extending through a false tendon into an echogenic thickened posterior papillary muscle, causing a grade II aortic and grade I/II mitral regurgitation. Short term corticosteroid and long term non-steroidal anti-inflammatory drug and disease modifying antirheumatic drug treatments efficiently controlled the symptoms. The cardiac findings remained unchanged during a follow up of 20 months. Careful cardiac evaluation appears to be mandatory even in these young patients.

Case report
We present the case of a 13 year old boy with a painful swelling of the left knee and pain of the left heel. He had a one year history of sustained pain and recurrent swelling of the left ankle and intermittent pain in the right hip. The upper extremities and the back were not involved and fever was never observed. There was no family history of arthritis or other connective tissue diseases.

Physical examination
The left knee was swollen with tenderness and limitation of motion. The left heel was tender at the insertion of the Achilles tendon into the calcaneus. There was no tenderness, flattening, or rigidity of the lumbosacral spine. Chest expansion was normal and neurological symptoms or systemic signs such as fever, diarrhoea, urethritis, and iritis were absent.

Cardiac auscultation detected a grade 2/6 systolic murmur that was mesosystolic and with low frequency (vibrating) at the cardiac apex. Furthermore, there was a grade 1/6 holodiastolic decrescendo murmur at the lower left sternal border.

Laboratory findings
Signs of active inflammation were an increased erythrocyte sedimentation rate of up to 114 mm/h and a maximal C reactive protein concentration of 32.5 mg/l but no leucocytosis. Mild microcytic anaemia (haemoglobin 1.70 mmol/l, haematocrit 0.326, mean corpuscular volume 77.9 fl) was associated with thrombocytosis (platelet count 571 × 10^9/l). The HLA-B27 marker tested positive, and the immunological markers rheumatoid factor and antinuclear antibodies were negative. As positive antistreptolysin O titres are very common in the Korean population, its occurrence in our patient was considered unspecific.

Signs of inflammation were also found in the synovial fluid (red cell count 864 × 10^9/l, white cell count 1.764 × 10^9/l, 60% polymorphonuclear neutrophil leucocytes, 35% lymphocytes) without evidence of a bacterial genesis (Gram stain and cultures were negative).
RADIOGRAPHIC EVALUATION
Radiographical (posteroanterior view) findings of both knees and the pelvis including the sacroiliac joints were normal. Active inflammation of the right hip joint was seen in a scintigraphic bone scan. Magnetic resonance imaging documented a pronounced joint effusion in the left knee with synovial enhancement and an abnormally enhanced bone marrow signal in the proximal tibia.

ECHOCARDIOGRAPHY
In the absence of pericardial or myocardial involvement, we found an echogenic structure measuring 8 × 11 × 20 mm at the site of the fibrous continuity between the aortic and mitral valve (fig 1). This structure served as the origin of a left ventricular false tendon connecting to the posterior papillary muscle, which itself had a gross thickening with a diameter of 16 mm (fig 1). The false tendon explained the child’s typical vibrating systolic murmur. Both the aortic and the mitral valve were partially included in this echogenic structure, causing a grade II aortic and a grade I/II mitral regurgitation (fig 2).

TREATMENT
Short term low dose oral administration of a corticosteroid (prednisolone 0.5 mg/kg body weight, tapered over 12 weeks) led to quick relief of the symptoms. A parallel long term treatment with a combination of a disease modifying antirheumatic drug (sulfasalazine) and a non-steroidal anti-inflammatory drug (naproxen) was initiated. Single intra-articular injections of a corticosteroid (triamcinolone) effectively controlled recurrences of joint inflammation. Physical therapy included daily active bending and deep breathing exercises directed towards preventing loss of range and poor positioning in the spine and chest.

FOLLOW UP
The effectiveness of the treatment was shown by diminishing signs of inflammation in

Figure 1  Echogenic 8 × 11 × 20 mm structure between the aortic and mitral valves (arrows, parasternal long axis (upper left), parasternal short axis (lower left)) connected by a false tendon to the echogenic posterior papillary muscle (arrows, parasternal long axis (upper right), parasternal short axis (lower right)).

Figure 2  Colour Doppler echocardiography showing valve dysfunction: grade II aortic regurgitation (arrows, parasternal long axis (left)) and grade I/II mitral regurgitation (arrows, parasternal long axis (right)).
scintigraphy of the right pelvis. Two recurrences of painful swelling and inflammation of the left knee 7 and 15 months, respectively, after initiation of the oral treatment prompted acute local corticosteroid injections. The cardiac findings remained unchanged over the follow-up of 20 months.

Discussion
The combination of pauciarticular arthritis of the lower extremities, typical enthesitis, positivity for HLA-B27, and negativity for rheumatoid factor and antinuclear antibodies in an adolescent boy is very suggestive of early juvenile ankylosing spondylitis, even without sacroiliitis.

Such patients, although most of them eventually develop sacroiliitis, do not meet the accepted diagnostic criteria for ankylosing spondylitis or juvenile ankylosing spondylitis at the onset of the disease. Therefore, descriptive terms such as “HLA-B27 positive juvenile arthritis” or “syndrome of seronegativity, enthesopathy, and arthropathy” (SEA syndrome) have been proposed and may be more appropriate.

Cardiovascular involvement is a common finding in adult patients with ankylosing spondylitis. However, it is considered rare in juvenile ankylosing spondylitis and only a small number of cases have been published with aortitis preceding the onset of sacroiliitis. Inflammation of the mitral valve is less common in adults, rare in juvenile ankylosing spondylitis, and not yet reported to precede the onset of sacroiliitis. The cardiac involvement in this case might also have been missed had a notable vibrating murmur of an aberrant left ventricular false tendon not alarmed the involved clinician. In the present patient the false tendon might have promoted the development of inflammation and fibrous transformation of the involved structures by means of unusual shear and tension at both ends—the fibrous continuity between the aortic and mitral valves and the posterior papillary muscle.

Careful cardiac evaluation appears to be mandatory in all cases of HLA-B27 positive juvenile arthritis, even in young patients. Although the long term outcome is not yet clear in our case, control of the inflammation by systemic treatment with non-steroidal anti-inflammatory and disease modifying antirheumatic drugs seems to stop or at least slow down progression of the valvulitis.

HLA-B27 positive juvenile arthritis with cardiac involvement preceding sacroiliac joint changes

S J Lee, H Y Im and W C Schueller

*Heart* 2001 86: e19
doi: 10.1136/heart.86.6.e19

Updated information and services can be found at:
http://heart.bmj.com/content/86/6/e19

These include:

**References**
This article cites 9 articles, 3 of which you can access for free at:
http://heart.bmj.com/content/86/6/e19#BIBL

**Email alerting service**
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

**Notes**

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/