

Investigations showed: erythrocyte sedimentation rate 55 mm in the first hour; C reactive protein concentration normal; haemoglobin concentration 12.1 g/dl; and white cell count $5.1 \times 10^9/l$; IgM rheumatoid factor was absent and HLA B27 antigen was present. Radiology of the spine and pelvis showed changes typical of ankylosing spondylitis. Detailed views of the atlantoaxial region showed rotational subluxation of C1 relative to C2 and the occiput. There was apparent collapse of the lateral mass of the atlas on the right side, which was confirmed by tomography and computed tomography. Injected white blood cells labelled with isotopes were not taken up at the site of the lesion.

Conservative management by immobilising the cervical spine in a moulded collar gave rapid symptomatic relief.

Comment

Ankylosing spondylitis and rheumatoid arthritis differ in the prevalence and distribution of disease of the spinal joints. Atlantoaxial displacement has been reported in both diseases but less commonly in ankylosing spondylitis.^{2,3} A recent report of non-reducible rotational head tilt in patients with rheumatoid arthritis attributed this deformity to collapse of the lateral masses of the atlas and axis as a result of erosive arthropathy of the facet joints.¹ We report the first case of non-reducible rotational head tilt in ankylosing spondylitis. Special radiological views of the upper cervical spine are required in rheumatoid arthritis. In ankylosing spondylitis radiographs of the cervical spine may be more difficult to interpret. The appearance on the lateral view may be misinterpreted as a fracture of a vertebral facet. Views through the mouth are difficult to obtain in patients with abnormal neck posture. A slight degree of rotation of the head may result in radiographic artefacts mimicking rotatory and lateral subluxations. Tomography of the upper cervical spine was helpful in our case, and computed tomography permitted more precise diagnosis of the subluxations.

The development of non-reducible rotational head tilt appears to have two components. Firstly, there is destruction of part of the lateral mass of the atlas. In rheumatoid arthritis and ankylosing spondylitis this is secondary to chronic synovitis with erosion of cartilage and subchondral bone and eventual destruction and collapse of the lateral mass. Secondly, although in rheumatoid arthritis ligamentous laxity may contribute to the rotational element of the deformity, this is unlikely to be an important feature of ankylosing spondylitis, in which fibrosis and even bony ankylosis are the usual outcome. It might be argued that in ankylosing spondylitis conservative treatment of non-reducible rotational head tilt with immobilisation in a support collar is more likely to succeed because subsequent ankylosis of the inter-facet joints would tend to stabilise the deformity and relieve the pain.

¹ Halla JT, Fallahi S, Hardin JG. Non reducible rotational head tilt and lateral mass collapse. *Arthritis Rheum* 1982;**25**:1316-23.

² Martel W. The occipito-atlanto-axial joints in rheumatoid arthritis and ankylosing spondylitis. *Am J Roentgenol* 1961;**86**:223-40.

³ Sharp J, Purser DW. Spontaneous atlanto-axial dislocation in ankylosing spondylitis and rheumatoid arthritis. *Ann Rheum Dis* 1961;**20**:47-76.

⁴ Weinstein PR, Karpman RR, Gall EP, Pitt M. Spinal cord injury, spinal fracture, and spinal stenosis in ankylosing spondylitis. *J Neurosurg* 1982;**57**:609-16.

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Twelfth rib syndrome: a differential diagnosis of loin pain

The slipping or clicking rib syndrome affecting the eighth, ninth, or 10th rib is recognised as a cause of upper abdominal pain.¹⁻³ Pain associated with the 11th or 12th rib is less well known and has been reported only twice.^{4,5} We report on four consecutive patients whose pain was in the loin region and was due to intercostal neuralgia

associated with the 11th or 12th rib, or both, who were cured by resection of the relevant rib or ribs. The patients were followed up for between six and 30 months (mean 17 months).

Case reports

Case 1—A 34 year old woman complained of intermittent right loin pain over the previous five years. There were no associated symptoms and no history of trauma. The pain was sometimes preceded by certain activities, and, interestingly, she found that using a vacuum cleaner was especially likely to produce discomfort. She had consulted practitioners in various specialties about her pain and had been extensively investigated on several occasions with no result. On examination we found that manipulation of the tip of the right 12th rib exactly and dramatically reproduced her pain. Subsequent excision of the rib resulted in an immediate cessation of the pain, and she remained pain free 30 months after the operation.

Case 2—A 21 year old girl was referred because of persistent left loin pain. She had been extensively investigated previously, when gastrointestinal and repeated urological x ray films had been normal. Urine culture had consistently yielded negative results, and a gynaecological examination had shown no abnormality. Most recently a therapeutic trial with antibiotics had failed to cure her "chronic pyelonephritis." She had complained of severe intermittent pain in her left loin over the previous two years. Initially the pain was described as sharp and radiating towards the corresponding groin, but subsequently, during the attacks, the pain became a dull ache lasting for some 24 hours at a time. She had noticed that she could gain relief by flexion of her spine. Manipulation of the left 11th and 12th ribs resulted in exquisite tenderness, exactly reproducing her pain. The ribs were resected with immediate relief of her pain, which did not recur during 18 months of follow up.

Case 3—A 37 year old woman complained of an ache in her left loin over the previous 10 months. It tended to be worse at night, when it disturbed her sleep, and was relieved by sitting forwards. There was no history of trauma, and no associated symptoms were present, but she recalled a milder form of the same discomfort during her two pregnancies 17 and eight years previously. Manipulation of the left 11th and 12th ribs exactly reproduced her pain, and after excision of the ribs the pain disappeared. She remained pain free in the 15 months after the operation.

Case 4—A 19 year old girl presented with acute right loin pain and a history of intermittent right loin pain for 18 months. The pains were initially sharp but then subsided into a dull ache lasting up to two days. The pain radiated to her right groin and subcostal region and was exacerbated by rotation of the trunk. The symptoms had been attributed to urinary tract infections, but intravenous urography had yielded normal results and infection had been proved on only one occasion. Examination elicited extreme tenderness over the 11th rib, and manipulation of that rib exactly reproduced her symptoms. The rib was resected and she remained pain free six months after the operation.

Comment

The hallmark of the syndrome we describe is pain in the loin. This may be intermittent or continuous and may be described as sharp pain or a dull ache or a combination of the two. None of our patients gave a history of trauma to the ribs or relevant associated symptoms. The condition appears to be quite common, and one of us (DGM) has seen four further cases referred for investigation of "renal pain."

The diagnosis is a clinical one and is made when the patient's pain is exactly reproduced by manipulation of the affected rib or ribs. Patients may be referred for investigation of presumed renal pain, and failure to recognise the true cause of the symptoms may lead to unnecessary investigations, possible inappropriate treatment, and needlessly prolonged pain for want of a simple surgical procedure.

¹ Davies-Colley R. Slipping rib. *Br Med J* 1922;**i**:432.

² Holmes JF. Slipping rib cartilage. *Am J Surg* 1941;**54**:326-38.

³ Heinz GJ, Zavala DC. Slipping rib syndrome. *JAMA* 1977;**237**:794-5.

⁴ Rawlings MS. The "rib syndrome." *Diseases of the chest* 1962;**41**:432-41.

⁵ Soltau HKV. Memoranda. *Br Med J* 1922;**i**:516.

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