Charcot joint in a case of syringomyelia with spinal dysraphism

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DESCRIPTION
A 35-year-old man with a swelling over the right shoulder and right anterior chest wall was referred to our hospital. The swelling had been present for over 2 months and was mildly painful. No history of trauma was present. On examination, there was limitation of shoulder movements. Swelling was firm in consistency, non-tender, warm on palpation, fixed to underlying muscles and with reduced sensation. There was no significant history of diabetes mellitus. No prior investigation had been done.

Shoulder radiograph revealed joint destruction and extensive bone resorption of the head of the right humerus with increased shoulder joint space. Also noted was sclerosis of the proximal right humerus and glenoid cavity. Multiple areas of periarticular soft tissue calcifications were seen around the right shoulder joint with displaced adjacent fat planes (figure 1).

CT of the chest was done, which was suggestive of a large, thick-walled peripherally enhancing collection in the region of the right shoulder joint and right anterior and lateral chest wall with multiple osseous fragments/loose bodies within and multiple air foci (figure 2). There was complete resorption of the head and neck of the right humerus, glenoid fossa and lateral aspect of acromion process as well as the coracoid process with resultant disorganisation and dislocation of the right shoulder joint. Radiological features were suggestive of neuropathic joint. In view of presence of air foci, possibility of infection was suggested. A spine MRI was further advised to look for syrinx.

MRI showed bony destruction involving the humeral head and glenoid with complete tear of the tendons of the rotator cuff. Also seen was destruction of the glenoid labrum, glenohumeral ligaments and biceps-labral anchor. A large, thick-walled peripherally enhancing collection with patchy areas of diffusion restriction was seen with extension into the acromioclavicular joint superiorly and into the right arm into biceps muscles and posterior compartment of the right arm (figure 3).

MRI of the whole spine showed a low-lying cord with syrinx from C1 to L3 vertebral level. Diastematomyelia of the terminal portion of the spinal cord at L3 level was noted. Anterior meningocele was seen from S3 vertebral level to the coccyx causing scoloping of the posterior cortex of the vertebral bodies with spina bifida from S2 to S5 vertebral levels (figure 3).
The patient was treated conservatively with anti-inflammatory drugs and antibiotics for infection. He was advised protective motion exercises and a sling.

**Learning points**

- Neuropathic osteoarthropathy is described as the osseous and joint changes that occur secondary to loss of sensation, accompanying a variety of disorders. Location of the bony and joint findings, age of the patient and clinical history may provide clues to the underlying disease.\(^1\)
- The six Ds of neuropathic joint include distension of joint, increased density, debris, destruction of articular surface, dislocation and disorganisation of joint.
- In syringomyelia, neuropathic changes are relatively common in the shoulder and elbow.\(^2\)

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**REFERENCES**


**Figure 3** Sagittal T2-weighted images of the cervicothoracic (A) and lumbosacral (E) spine show syrinx within the spinal cord (white arrow in A) axial stir (B) and coronal stir (C). MRIs show resorption of the right humeral head and glenoid and collection in the right shoulder joint and proximal right arm. Complete tear of the tendons of the rotator cuff muscles and destruction of the glenoid labrum, glenohumeral ligaments and biceps–labral anchor is seen. T2-weighted images of the spine show focal splitting of the terminal portion of the spinal cord at L3 level suggesting diastematomyelia (white arrow in F). A CSF-filled sac is noted in the spinal canal suggesting a meningocele (arrow in E and H). Spina bifida is noted in the sacral vertebrae. CSF, Cerebrospinal Fluid.