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CHRONIC OSTEOMYELITIS OF THE ULNA OCCURRING IN SYRINGOMYELIA

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Although, in this day and age, acute and chronic osteomyelitis are seen much less than formerly, the conditions are by no means rare. However, chronic osteomyelitis of the ulna in the absence of a compound fracture is uncommon.

Trueta and Morgan (1954) reporting a series of cases of acute osteomyelitis, found the ulna to be the site of infection in four of one hundred patients. Blanche (1952) in his series, found two in 50 patients. Green, Nyhan and Fousek (1956) found the ulna involved in nine, in their series of 99 patients. However, in these authors' series, the lesion was acute and occurred in infancy or childhood.

Acute and chronic pyogenic osteomyelitis, in adults, are even less common conditions. Reviewing the literature of the past 35 years, a direct reference to the conditions could be found on three occasions only. Variava (1935) described the acute condition in a patient with a compound fracture of both forearm bones. The radius, however, was involved maximally. Dufour (1933) demonstrated the radiological history of one patient with chronic...
ulnar infection. A further chronic case was reported by Hendriock (1939) who regarded the ulna as a rare site for localization of bone infection.

Case History
Mr. R. A., aged 49, was admitted to hospital on September 24, 1960. His history was that in June, 1960, while pulling a heavy weight at work he experienced a sudden jerking sensation in the left forearm. Within a few hours his forearm had begun to swell, although it was seven days before this swelling was maximal. At the time of the accident he had noticed two septic abrasions on the left forearm. At this point he reported to his own practitioner for he had become generally unwell. His forearm was painful, swollen, red and stiff. Cellulitis was diagnosed and oral penicillin prescribed with resolution of signs and symptoms.

On September 14, 1960, the patient noticed a painless lump on the posterior aspect of the left forearm, associated with stiffness in the fingers. There was no systemic upset.

His previous history was significant in that in 1955 he had reported to another hospital with a rupture of the right biceps brachii. At that time there was noticed to be a dissociated loss of sensation in the right forearm and hand with obvious muscle wasting. A diagnosis of syringomyelia was made.

On examination, the patient was a stocky little man of healthy appearance. The conjunctive were well injected. There was no lymphadenopathy. On the posterior surface of the left ulna, at the juncture of its proximal and middle thirds, there was a diffuse, non-tender, bony swelling showing none of the characteristics of acute inflammation. Temperature 97° F., pulse 60/min., W.B.C. 6,500/cu. mm., E.S.R. 6 mm. hr.

Examination of the locomotor and peripheral nervous systems revealed the following:

**Upper Limbs.** Right: Extreme wasting of thenar, hypothenar and interossei groups of muscles. There was no voluntary movement in these muscles. The wrist flexors showed normal power but only the flexors to the ulnar three fingers were functioning. Extension of the wrist was normal. Elbow and shoulder movements were normal. Left: The thenar, hypothenar and interossei groups of muscles were wasted but opposition of thumb and little finger was still possible. The third dorsal and palmar interossei had weak function. Wrist, elbow and shoulder movements were normal.

**Lower Limbs.** Normal motor power and sensation were noted. Reflexes were brisk and equal. Plantar reflexes were flexor on both sides. There was, in addition, a well-marked Horner’s syndrome on the right side.

X-ray films of the left forearm, taken on admission, showed patchy decalcification with periostitis and the presence of a sequestrum in the shaft of the ulna.

A diagnosis of chronic, active osteomyelitis of the shaft of the ulna was made, and the presence of syringomyelia was confirmed.

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<td>Supinator</td>
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<td>Pain and temperature</td>
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<td>Vibration</td>
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**Treatment and Progress**
On September 29, 1960, under a general anesthetic, the ulna was approached by a longitudinal incision over the bone. The lesion was exposed and thoroughly saurized. The bone fragments and sequestrum, with some thick creamy pus, were sent to the bacteriologist. After operation, the limb was immobilized in an above-elbow plaster. Crystalline penicillin, 0.5 mega units b.d., was given intramuscularly for seven days.

**Bacteriology:** a profuse growth of coagulase-positive *Staphylococcus aureus* highly sensitive to penicillin.

The patient was discharged home on January 5, 1960, and three weeks later the plaster was removed because of the patient’s inability to use the opposite limb. Radiographs taken at intervals showed progressive regeneration of the ulna. Those on December 7, 1960, showed an almost complete return to normal contour. The wound healed rapidly, although a sero-sanguinous, bacteriologically sterile discharge was apparent for a few days.

**Discussion**
There seems to be little likelihood that this patient’s ulnar lesion was due to hematogenous spread from a distant focus of infection, but rather to be consequent upon an acute cellulitis in the forearm. It is unlikely that spread of infection to bone occurred directly, in the absence of abscess formation in association with cellulitis. A superficial cellulitis may destroy, in fibrosis, the lymphatics coursing proximally as a subcutaneous plexus, with the result that inflammatory products and viable organisms are carried into a deeper plane, and by retrograde flow into bone.

It may be suggested that the coincident syringomyelia may have predisposed to the superficial skin infection and the resultant osteomyelitis. That this may be so in the hand and fingers is undoubted, but in this patient’s case the left upper limb was less involved than the right, and the forearm itself had normal sensation. Indeed there is no reference in the literature of the past 10 years to osteomyelitis of the ulna occurring as a complication of, or in association with, syringomyelia.

I am grateful to Mr. J. H. S. Scott for his permission to publish the details of this case.

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