cardiac failure. Atrial fibrillation was present, but there was no cardiac enlargement and no murmurs. Chest radiography confirmed that the cardiac contour and size and lung fields were normal. Echocardiography showed no abnormality of either the mitral or aortic valve. Cardiac enzyme activities, electrolyte concentrations, and full blood count were all normal, and glandular fever screening test was negative. Complement fixation tests were negative for Coxiella burnetti, Mycoplasma pneumoniae, influenza A and B, adenovirus, parainfluenza, herpes simplex, respiratory syncytial virus, and cytomegalovirus. There was no psittacosis, however, as was follows: on 22 December 80, 512 on 14 January 81, 512 on 18 February 81, 64 on 16 April 81, and 32 on 17 June 81.

Treatment with tetracycline was started and continued for two months. By late January 1981 his symptoms had disappeared and when the titre to Chlamydia B had fallen from 512 in May 1981 he was admitted to hospital for cardioversion because of persistence of atrial fibrillation. After successful cardioversion, electrocardiography showed no evidence of pulmonary mitrale, ventricular hypertrophy, or myocardial ischaemia. Ten days later, after a mild flu-like illness, the heart lapsed into atrial fibrillation again despite maintenance quinidine treatment. Repeat cardioversion was not required as sinus rhythm occurred spontaneously after several days.

Closer questioning failed to elicit any exposure to or contact with birds by the patient or any member of the family.

Comment

Chlamydia B infection (psittacosis) is probably more common in this country than the reported cases indicate. In 1972 the total number of cases reported was 156\(^3\) and in 1975 it was 177.\(^3\) The benign course of the illness in most patients is undoubtedly responsible for many cases going undetected. There may not necessarily be a history of contact with infected birds, this being established in only 40 (26\%) of the 156 cases reported in 1972.

Complications are rare, but may include meningitis,\(^5\) encephalitis,\(^5\) and occasionally cardiac disease. Endocarditis necessitating valve replacement has been reported,\(^6\) and myocarditis may rarely occur.\(^6\)

We think that in the absence of any history of rheumatic fever and previous cardiac disorder, and with a normal echocardiogram, the atrial fibrillation in our patient occurring at the height of his psittacosis infection was directly related to it, possibly owing to mild accompanying myocarditis.


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Extensor digitorum brevis—a predictor of neuropathy in the leg?

The measurement of conduction velocity in the deep peroneal nerve is a common investigation in patients with suspected peripheral neuropathy. An impression was formed that the extensor digitorum brevis muscle was difficult to palpate in patients who had slowed conduction in the peroneal nerve. Muscle wasting is common in peripheral nerve lesions,\(^1\) so a formal study was performed to see whether wasting of this particular muscle was a reliable clinical predictor of electrophysiologically demonstrable nerve disease.

Patients, methods, and results

One hundred consecutive unselected patients, referred for routine measurement of nerve conduction velocity in the leg, were included in the study. No investigation was performed which was not a part of routine nerve testing. The results were collected from March 1979 to May 1981. Most patients were referred with suspected peripheral neuropathy, injury, or compression neuropathy.

An attempt was made to palpate the extensor digitorum brevis and a decision on its palpability or otherwise was made before any measurements of conduction velocity were made. Motor conduction velocity in the deep peroneal nerve was measured using a pair of Ag/AgCl surface electrodes to record the compound action potential of the extensor digitorum brevis. Supramaximal rectangular isolated electrical stimuli (duration 0.1 ms) were applied to the deep peroneal nerve just proximal to the ankle, and to the common peroneal nerve at the neck of the fibula. The latency between the stimulus and the onset of the muscle action potential was measured with a Medelec MS6 oscilloscope; the conduction velocity was calculated by dividing the distance between stimulus sites by the difference in the two latencies. Where conduction velocity was measured in both legs, data from one or other side were rejected according to the toss of a coin. All observations and measurements were made by the author, and therefore the study incorporated no element of “blindness.”\(^2\)

Of the 100 patients, 67 had a palpable extensor digitorum brevis, and in 33 the muscle was impalpable. There were 10 patients in whom myoelectrical activity could not be recorded with surface electrodes during nerve stimulation. In three of these patients, the dorsum of the foot was subsequently explored with a concentric needle electrode, but still no myoelectrical activity could be recorded during nerve stimulation. Even if these 10 patients were excluded from the calculations, there was a highly significant reduction in the mean conduction velocity for the “non-palpable” group (table), with a significant increase in the distal latency from ankle to muscle. The value for the “palpable” group were close to those reported for normal subjects.\(^3\)

The values for conduction velocity were separated into a high group (>40 m/s) and a low group (<39 m/s); the values for latency were divided into a long-latency group (>6.0 ms) and a short-latency group (<5.9 ms). The patients for whom myoelectrical data could not be obtained were placed in the low-velocity and long-latency groups. Of the non-palpable group, 27 had low conduction velocity while 57 of the palpable group had high conduction velocity (x\(^2\)=39.6; p<0.001). Of the non-palpable group, 19 had long latency while 60 of the palpable group had short latency (x\(^2\)=23; p<0.001).

Comment

The attempt to palpate the extensor digitorum brevis is a very simple clinical test to apply, and these results suggest that failure to palpate it is an excellent predictor of disease in the peroneal nerve.


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