sively in the dorsal columns of the spinal cord.

Results of right phrenic nerve conduction (n = 25) and transcortical magnetic stimulation with recording of the right diaphragm (n = 35) in healthy subjects, and in the patient with cervical infarction and critical illness neuropathy

<table>
<thead>
<tr>
<th>Measurement</th>
<th>Healthy subjects (Mean SD)</th>
<th>Patient</th>
</tr>
</thead>
<tbody>
<tr>
<td>Phrenic nerve conduction:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Latency (ms)</td>
<td>6.5 (0.8)</td>
<td>8.3 (5)</td>
</tr>
<tr>
<td>Amplitude (µV)</td>
<td>669 (159)</td>
<td>330 (5)</td>
</tr>
<tr>
<td>Cortical magnetic stimulation:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Latency (ms)</td>
<td>13.5 (1.4)</td>
<td>No response (5)</td>
</tr>
</tbody>
</table>

For the patient, the day of examination after onset of illness is given in parentheses.

Electrophysiological monitoring in neurological respiratory insufficiency

In patients with neurological respiratory insufficiency it may be difficult to determine whether there is a neuromuscular problem, impaired central drive, or both. Phrenic nerve conduction studies and needle EMG of the diaphragm are well established methods in the diagnosis and prognosis of neuromuscular causes of diaphragmatic weakness. Transcortical and cervical magnetic stimulation with recording from the diaphragm and somatosensory evoked potentials of the phrenic nerve can assess the motor and sensory pathways of the central respiratory drive. We report the value of peripheral and central respiratory electrophysiological studies in a patient with acute high cervical spinal cord infarction who developed secondary critical illness polyneuropathy.

This 76 year old previously healthy man had an acute onset of sharp, knife-like chest pain, and could not move his limbs a short time later. The pain lasted about an hour. On admission to hospital, three hours after onset of symptoms, his mental status and cranial nerves were intact. He was flaccidly quadriplegic with no voluntary movement except of elevation of both shoulders. Initially, he had diaphragmatic breathing clinically. Plantar responses were extensor. Pinprick, temperature, and light touch sensation were lost below C4 bilaterally. Vibratory sensation was lost in the limbs, but position sense was incompletely affected: passive movements were perceived in the toes on the left side, and at the knee on the right side. Within a few hours breathing became rapid and shallow. He was intubated and placed on a ventilator. The patient transiently worsened for about a week: he lost his ability to shrug his shoulders and the sensory level rose to below C3 bilaterally.

Spinal cord MRI showed abnormally increased signal anteriorly within the spinal cord signal from C4–T1 vertebral levels, and oedema ascending to C2. Examination of CSF was unremarkable. Serial electrophysiological studies for monitoring diaphragmatic muscle function were done (table). Phrenic nerve conduction studies were performed with electrical stimulations at the distal neck just posterior to the sternomastoid muscle. Transcortical magnetic stimulations were performed at the beginning of inspiration with a 90 mm circular coil positioned over Cz (determined by the 10–20 EEG system). No diaphragmatic compound muscle action potentials (CMAPs) were recorded after transcortical magnetic stimulation, and phrenic nerve conduction studies showed borderline onset latency and diaphragmatic CMAP amplitude at day 5 (figure). Transcortical magnetic stimulation on day 21 showed a delayed but recordable diaphragmatic CMAP, and normal phrenic nerve conduction. At this time he developed a septic syndrome due to pneumonia. The patient could not be weaned from the ventilator within the following weeks, and the necessity of a phrenic nerve pacemaker was raised. Subsequently, the sensory level descended distal to C6, and some movement was noted in the proximal left upper limb muscles (MRC 3). Electrophysiological follow up study on day 51 showed complete

Values of latencies and amplitudes of the right diaphragmatic compound muscle action potential.

Cortical magnetic stimulations were performed with a 90 mm circular coil positioned flat over the vertex (Cz). The phrenic nerve was stimulated electrically at the posterior border of the sternomastoid muscle in the supraclavicular fossa. The diaphragmatic compound muscle action potential was recorded bilaterally with surface electrodes from the xiphoid (active electrode) and the costal margin (reference electrode) for both magnetic and phrenic nerve stimulation. The table shows that there was no significant right-left differences.
We are venous thromboses associated with the ported by for possibly, lower in increased potentials evoked potentials the site amplitude after phrenic relation. CMAPs on respiratory electrophysiological extubated. Respiratory electrophysiological study, of the somatosensory spectral analysis of the time of critical illness was transferred to the site. Needle EMG of the hand was transferred to the site. Neurophysiology was performed on the day. EMG of the hand was transferred to the site. The patient remained in hospital for five months after onset of the disease. The study demonstrates the value of respiratory electrophysiological studies in localising the site of neurological causes of respiratory failure. An unusual feature occurred in the time of critical illness. The diaphragm CMAP amplitude after phrenic nerve stimulation was lower than the CMAP amplitude evoked by central magnetic stimulation. Submaximal stimulation was excluded as a cause. Central enhancement of somatosensory evoked potentials is known to occur in patients with peripheral neuropathy and may be a cause of the explanation in our patient. Moreover, the amplitude of magnetically evoked potentials increases in contracted muscles. In healthy subjects magnetic stimulations during forced inspiration increase the diaphragmatic CMAP by about 100%. Possibly, the increased muscle tone leads to facilitation of the recording muscle resulting in increased CMAP amplitude. Further studies will be necessary to clarify the pathophysiological mechanisms of this phenomenon.

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Fulminant encephalopathy due to the cerebral hemispheric primary antiphospholipid syndrome

The antiphospholipid antibody syndrome is characterised by recurrent arterial and venous thromboses associated with the presence of antiphospholipid antibodies. Neurological features can include cerebral ischaemia, multi-infarct dementia, migraine, epilepsy, transient ischaemic attacks and chorea. We describe a patient with catastrophic primary antiphospholipid syndrome who presented with right hemispheric dysfunction which rapidly evolved to a fulminant encephalopathy associated with multiorgan failure.

A fifty-eight-year-old woman was admitted with a general malaise, headache, anaemia, and weight loss of a few months. On examination, the patient was alert and cooperative but had several focal neurological features, including left sided facial and limb weakness. She had a history of classic migraine for 35 years, had had several lower limb deep venous thromboses, and had been investigated for pleuritic chest pain for which no cause was established. She developed Graves’ disease at 45 years of age and was treated with radioiodine followed by thyroxine. She had no relevant family medical history.

On admission there were no significant abnormalities in the cardiovascular, respiratory, or gastrointestinal systems. She had a 5 cm skin lesion over the right lateral malleolus. She was disoriented, restless, and unable to give a coherent history. There was no meningism and the optic fundi and eye movements were normal. She had a mild dysarthria and mild upper motor neuron facial weakness and a mild left hemiparesis with brisk tendon reflexes and an extensor plantar response. She also had bilateral papillary dilatation and pout reflexes, and left sided sensory inattention.

She had thrombocytopenic (platelets 125 × 10^9/L) but her full blood count and peripheral blood film were otherwise normal. Her erythrocyte sedimentation rate was 89 mm/h and C reactive protein concentration was 386 mg/L. The prothrombin time was normal, but the activated partial thromboplastin time (aPTT) was prolonged at 37.2 s (normal range 23.5–36.4 s). The D dimer assay was normal but her fibrinogen concentration was low (1:15 g/L, normal 1–4.0 g/L). Her plasma urea was slightly increased at 5.4 mmol/L (normal range 2.8–7.1 mmol/L). Her plasma electrolytes, creatinine, liver function tests, bone chemistry, thyroid function, and blood glucose were normal and a venereal disease research laboratory test was negative. Urine, blood, and CSF cultures were sterile and only skin commensal organisms were isolated from the foot ulcer. A chest radiograph, ECG, echocardiogram, and brain CT were unremarkable. Lumbar puncture disclosed clear CSF, with a normal opening pressure, 6 white blood cells/μL, and normal protein and glucose concentrations.

She was treated with broad spectrum antibiotics for an unusual infection of unknown origin with a provisional diagnosis of right hemispheric ischaemic stroke. However, she became increasingly obtunded, developing signs of bilateral hemispheric dysfunction including pyramidal distribution weakness in all four limbs and bilateral extensor plantar responses. She developed livedo reticularis over her back and lower limbs (fig 1). A systolic murmur was noted at the cardiac apex. Her respiratory function deteriorated, she became hypotensive and oliguric, and required assisted ventilation and inotropic support.

Thyroid microsomal antibodies were present at a titre of 1 in 400, but other autoantibody titres were negative (including dsDNA-Ab, ANCA, and Anti-GBM-Ab). Her anticardiolipin antibody concentrations were raised (serum IgG aCL 41·5 GPL, IgM aCL 25·9 MPL, normal <10 GPL/MPL). Biopsies of the skin ulcer and an area of livedo reticularis disclosed widespread thrombosis within the small vessels but no vessel wall inflammation. Echocardiography now showed a vegetation on the mitral valve. Abdominal CT was normal and a repeat brain CT showed no abnormality. An EEG showed diffuse generalised slow wave activity.

After a diagnosis of catastrophic antiphospholipid syndrome was confirmed, she was managed aggressively with intravenous heparin and immunosuppressed with methylprednisolone (1 g intravenously/day), cyclophosphamide (500 mg, single dose), and plasmapheresis. Her urinary output increased over the next week and cardiorespiratory support was no longer needed. Her neurological condition did not improve and she remained unaware and unresponsive with no clinical evidence of cortical function but intact brainstem reflexes. She died 24 days after admission to hospital.

The most striking pathological finding postmortem was of widespread small vessel thrombosis; the tissues involved included the brain, myocardium, lung, kidney, skin, bone marrow, uterus, ovary, bladder, pancreas, stomach, and small intestine. Examination of the heart showed marantic vegetations of the mitral valve which were Gram stain and culture negative. The larger veins and arteries, including the coronary arteries, were patent with only mild atheroma seen.

Macroscopic examination of the cerebral hemispheres showed extensive confluent areas of laminar cortical infarction in the watershed zones. The midbrain, brainstem, cerebellum, and spinal cord were macroscopically unremarkable. Microvascular changes included small arteriolar and venular thrombotic occlusion in the brain, brainstem, spinal cord, and meninges. There was no associated inflammatory reaction (fig 2).

In the absence of convincing evidence for infection, the evolution of multiorgan failure, together with livedo reticularis, suggested the possibility of a systemic vasculitis or a coagulopathy. Disseminated intravascular coagulation and the haemolytic uraemic syndrome were excluded by the absence of

![Oclusive vascular lesion in the corpus striatum showing an amorphous protein plug with peripheral endothelial and macrophage proliferation. The vessel seems to be dilated at the site of the lesion. The surrounding neuropil shows a pronounced gliosis. There is no evidence of vasculitis. Magnification bar = 100 μm.](image-url)
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