Motor root conduction in neuralgic amyotrophy: evidence of proximal conduction block

Yew-Long Lo, Kerry R Mills

Abstract
Objective—To determine the presence and role of proximal conduction block in neuralgic amyotrophy.
Methods—Percutaneous electrical stimulation of cervical roots and brachial plexus was employed in eight patients with neuralgic amyotrophy. Root to Erb’s point compound muscle action potential amplitude ratios for abductor digiti minimi, extensor digitorum communis, biceps, and deltoid muscles were compared with results obtained from 10 healthy controls.
Results—Conduction block in the nerve to one muscle was found in three of eight patients (38%) suggesting focal proximal demyelination. Repeat studies showed axonal degeneration, resolution, and persistence of conduction block in these three patients respectively.
Conclusion—Focal conduction block plays a significant part in the pathogenesis of neuralgic amyotrophy, which is generally regarded as an axon loss process. Therapeutic intervention should be directed to patients with persistent conduction block, with the aim of eradicating the block and possibly minimising subsequent axon loss.

Keywords: neuralgic amyotrophy; proximal stimulation; conduction block

Neuralgic amyotrophy, otherwise known as the Parsonage-Turner syndrome, idiopathic brachial plexopathy, or brachial neuritis, refers to a well defined clinical syndrome having documented precipitating factors and possible aetiologies. Severe pain of rapid onset around the shoulder region is usually the initial symptom, often persisting for days to weeks. This is followed by weakness and wasting of shoulder girdle and upper limb muscles, which become prominent features as the pain begins to subside. The distribution of weakness is variable, although muscles around the shoulder girdle are most commonly involved. Sensory symptoms are not marked, but may occur in a root or nerve distribution. Lesions in a peripheral nerve distribution rather than a patchy plexopathy may occur. Cases with involvement of the sternomastoid or diaphragm have also been described, and familial and recurrent cases are encountered.

Although the clinical entity has long been recognised, little is known of its aetiology. An immune mediated or allergic process has been suggested on the basis of its association with infections, but other well documented precipitating factors such as childbirth, trauma, and surgery create difficulty in proposing a unified pathogenetic mechanism. Neurophysiologically, neuralgic amyotrophy usually has features of an axonopathy, but several reported cases, often with atypical clinical features, show evidence suggestive of nerve fibre demyelination. It has also been suggested that neuralgic amyotrophy may represent a variant of chronic inflammatory demyelinating polyneuropathy.

Most neurophysiological studies in this condition have been performed on distal nerve segments, which are not helpful in the localisation of proximally situated focal lesions. Essentially, neurophysiological investigation relies on the demonstration of neurogenic changes in a patchy distribution by EMG. In this study, we have employed proximal nerve stimulation of the cervical roots and the brachial plexus to investigate for the first time proximal motor conduction in neuralgic amyotrophy. We find that in most cases the lesion is axonal degeneration in nature, but in some there is evidence of proximal conduction block.

Methods
PATIENTS AND HEALTHY CONTROLS
Patients presenting with sudden onset of shoulder pain followed up to 4 weeks later by weakness of upper limb muscles were included in the study. Patients with possible cervical radiculopathy, sensorimotor neuropathy secondary to systemic conditions, local trauma, and anterior horn cell disease were excluded. Of an original group of 14 patients presenting with suggestive features, six were excluded for these reasons. Over a period of 11 months, eight patients (including one woman) were recruited. In addition, 10 healthy controls were studied using identical techniques. All gave their informed consent and were free from any neurological conditions. Their ages ranged from 27 to 51 years. The interval between the onset of symptoms to the time of study ranged from 2 weeks to 8 months.

NERVE CONDUCTION STUDIES AND EMG
All patients had peripheral nerve conduction studies, including median and ulnar sensory studies, median and ulnar motor conduction, and minimal F wave latency using conventional techniques. Needle EMG was performed as clinically indicated.

MOTOR ROOT AND ERB’S POINT STIMULATION
This was provided by a high voltage electrical stimulator (Digitimer Ltd, Hertfordshire, England) producing a peak output voltage of 750
V. To achieve maximal stimulation, 14 mm diameter surface electrodes with an interelectrode distance of 14 cm were used. Stimuli with a 5 μs rise time and 100 μs decay were applied over the cervical vertebral column via silver electrodes covered in saline soaked lint and mounted on a straight plastic holder. The cathode was placed in the midline between the appropriate vertebral spines. With this method, the motor roots are excited 2–4 cm distal to the anterior horn cell body, corresponding to the exit foramen of the motor root from the spinal column. 

Stimulation was performed over the C7/T1 interspace and moved upwards sequentially for C7, C6, and C5 root stimulation. Cathodal stimulation at Erb’s point was also performed using the same stimulator. It was clearly important to ensure that supramaximal stimulation was delivered at each site. To this end, responses were collected at increasing stimulation intensity, until it was certain that a further increase of at least 10% beyond maximal stimulation intensity did not produce a larger response.

### Table 1: Clinical features of patients

<table>
<thead>
<tr>
<th>Patient</th>
<th>Age</th>
<th>Sex</th>
<th>Interval (months)</th>
<th>Shoulder pain</th>
<th>Duration (weeks)</th>
<th>Family history</th>
<th>Patient history</th>
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<tbody>
<tr>
<td>1</td>
<td>29</td>
<td>M</td>
<td>2</td>
<td>R</td>
<td>2</td>
<td>–</td>
<td>–</td>
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<tr>
<td>2</td>
<td>67</td>
<td>M</td>
<td>8</td>
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<td>3</td>
<td>–</td>
<td>–</td>
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<tr>
<td>3</td>
<td>51</td>
<td>M</td>
<td>1</td>
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<td>–</td>
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<tr>
<td>4</td>
<td>68</td>
<td>M</td>
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<td>L</td>
<td>3</td>
<td>4</td>
<td>–</td>
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<tr>
<td>5</td>
<td>48</td>
<td>M</td>
<td>2.0</td>
<td>R,L</td>
<td>3</td>
<td>–</td>
<td>R side 2 years ago</td>
</tr>
<tr>
<td>6</td>
<td>62</td>
<td>M</td>
<td>2.0</td>
<td>R</td>
<td>4</td>
<td>2</td>
<td>–</td>
</tr>
<tr>
<td>7</td>
<td>37</td>
<td>F</td>
<td>2.5</td>
<td>L</td>
<td>2</td>
<td>–</td>
<td>R side 9 years ago</td>
</tr>
<tr>
<td>8</td>
<td>58</td>
<td>M</td>
<td>0.5</td>
<td>R</td>
<td>2</td>
<td>–</td>
<td>–</td>
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</table>

R=Right; L=left. Interval refers to the time in months from the onset of symptoms to the initial examination and neurophysiological study. Duration refers to the duration of shoulder pain. Patient 4 had two affected siblings. Patient 5 had salmonella enteritis 1 week before onset. Patient 8 had a 20 year history of limb girdle dystrophy.

### Results

The clinical features of all eight patients are summarised in Table 1. Of the eight patients, five had sensory symptoms, two had recurrent neuralgic amyotrophy, with one on the ipsilateral (patient 6) and one on the contralateral side (patient 7), two gave a history of trauma or infection (patients 5 and 7) and one patient had a positive family history of neuralgic amyotrophy (patient 4). Four patients (1, 2, 5, and 7) showed predominantly upper plexus involvement, three (3, 4, and 8) lower plexopathy, and patient 6 had diffuse abnormalities referable to the upper and lower plexus.

### Table 2: Proximal electrical stimulation CMAP amplitudes (mV)

<table>
<thead>
<tr>
<th>Patient</th>
<th>Right ADM</th>
<th>EDC</th>
<th>Biceps</th>
<th>Deltoïd</th>
<th>Left ADM</th>
<th>EDC</th>
<th>Biceps</th>
<th>Deltoïd</th>
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<td>1</td>
<td>10.8</td>
<td>11.0</td>
<td>15.8</td>
<td>17.2</td>
<td>28.9</td>
<td>24.1</td>
<td>15.8</td>
<td>20.4</td>
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<td>2</td>
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<td>4.0</td>
<td>5.8</td>
<td>14.4</td>
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<td>18.6</td>
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<td>9.9</td>
<td>4.7</td>
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<td>17.3</td>
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<td>14.3</td>
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<td>8.0</td>
<td>6.6</td>
<td>6.7</td>
<td>20.0</td>
<td>20.7</td>
<td>4.5</td>
<td>5.0</td>
</tr>
<tr>
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<td>6.5</td>
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<td>3.1</td>
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<tr>
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<td>10.6</td>
<td>9.3</td>
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<td>3.5</td>
<td>2.6</td>
<td>2.6</td>
</tr>
<tr>
<td>8r</td>
<td>1.4</td>
<td>2.3</td>
<td>2.0</td>
<td>2.2</td>
<td>3.5</td>
<td>3.5</td>
<td>2.8</td>
<td>2.8</td>
</tr>
</tbody>
</table>

N=Neck; E=Erb’s point; ADM=abductor digitii minimi; EDC=extensor digitorium communis. r=repeat study. —=not done.
The results are summarised in table 2 and figs 1 and 2. All patients studied tolerated the procedure well. Conduction block was defined as a root to Erb’s CMAP amplitude ratio less than 2 SD of the healthy controls’ values for each muscle. Control mean (SD) ratio values for the right ADM, EDC, biceps brachii, and deltoid were 0.93 (0.05), 0.93 (0.08), 1.04 (0.24), and 0.95 (0.09) respectively. The corresponding values for the left side were 0.94 (0.19), 0.91 (0.17), 0.94 (0.16), and 0.96 (0.13). Conduction block between root exit foramina and Erb’s point was demonstrable in three patients (2, 6, and 8), who exhibited ratios of 0.37, 0.39, and 0.64 in the right deltoid, biceps, and abductor digiti minimi respectively. Ratios of the negative peak areas of CMAPs were also considered and identified the same three patients as having significant conduction block. With the exception of patient 1, who had no motor conduction abnormality, all other patients had small CMAPs from both root and Erb’s point stimulation, indicating an underlying loss of axons.

Repeat studies on the three patients with conduction block at various intervals after the initial study yielded additional information on disease progression. In patient 2, the conduction block seen in the right deltoid gave way to an axon loss lesion with reduction of CMAP amplitudes from both root and Erb’s point stimulation. The initial conduction block in the right biceps in patient 6 disappeared with increase of CMAP amplitudes evoked from root and Erb’s stimulation and a ratio increasing to normal. However, this was not the case in patient 8, whose repeat study showed increased conduction block from 0.64 to 0.61 in the right ADM.

**Discussion**

The presence of conduction block is well documented in acute or chronic demyelinating polyneuropathy and multifocal motor neuropathy with conduction block. However, focal conduction block, particularly in the most proximal motor root segments, has not been shown previously in neuralgic amyotrophy. Most existing studies employed conventional techniques, which are not helpful in localising proximal lesions precisely. In our experience and others, F waves were not useful in this respect; presumably surviving or unblocked axons are able to generate an F response at the normal latency. Needle EMG is useful in identifying active denervation changes, and paraspinal muscle sampling with its inherent disadvantages of technical difficulty and myotomal overlap only localises the lesion proximal or distal to the posterior primary rami. By contrast, we find percutaneous electrical root stimulation a useful, well tolerated test for localising motor root diseases.

Clinical and neurophysiological evolution of the three patients in which conduction block was demonstrated is of some interest. Patient 2 presented with a typical history of neuralgic amyotrophy with no obvious precipitating cause. His condition did not improve and when
seen 8 months later a conduction block was detected recording from the right deltoid. He had repeat studies 7 months later (15 months after initial presentation) which disclosed findings consistent with an axon loss lesion. Clinically, he did not show improvement in muscle power in his right deltoid, although the distal muscles were considerably stronger. Patient 6 had a previous episode of neuralgic amyotrophy which recovered completely 2 years before this recurrent episode. His initial studies performed 2 months after onset showed a conduction block in the right biceps brachii. Repeat studies 9 months later (11 months after the recurrent presentation) showed resolution of the block and improved CMAP amplitudes. Clinically, he had significant improvement in power of both proximal and distal muscles. Patient 8 presented with typical symptoms of neuralgic amyotrophy, again with no obvious precipitating cause. He had a 20 year history of limb girdle muscular dystrophy confirmed on biopsy, with minimal functional deficit. Initial studies within 2 weeks of onset of his symptoms showed a conduction block recording from the right ADM, the weakest muscle. Repeat studies 1 month later (6 weeks after initial presentation) showed further reduction of CMAP amplitudes and persisting conduction block. He did not experience any clinical improvement in muscle power.

It should be pointed out, however, that many peripheral nerve disorders are characterised by a mixture of demyelinating and axonal degeneration features. Few cases of neuropathy in general fall precisely into pure axonal or demyelinating categories. A neuropathy from extensive demyelination is often associated with axonal degeneration. Conversely, axonal neuropathy may lead to secondary paranodal demyelination. Whereas it is well known that the distal portion of a sectioned nerve undergoes secondary demyelination as a result of axonal degeneration, the mechanism of paranodal segmental demyelination from localised changes or diffuse metabolic insults is less well understood. Various postulates have been put forward, ranging from hypotheses akin to pre-Wallerian degeneration mechanisms to local processes causing a reduction in volume of the axon cylinder with disturbed ultrastructure of transport organelles. It is therefore appropriate to consider whether demyelinating or axonal loss processes coexisted or if one preceded the other.

Pathological studies favour an immune mediated demyelinating pathogenesis over an axon loss mechanism in neuralgic amyotrophy. Sierra et al reported decreased CD3 and CD8 lymphocyte subsets in five of six patients with acute neuralgic amyotrophy. Nerve biopsies of a patient with relapsing brachial plexus neuropathy and bilateral upper trunk conduction block showed florid multifocal mononuclear inflammatory plexus infiltrates. A postmortem based study showed widespread demyelination. A focal immune mediated mechanism of demyelination has been postulated in interleukin-2 associated brachial plexopathy. Immune complex formation has also been implicated as a cause of acute brachial plexus neuropathy and the Guillain-Barré syndrome associated with serum sickness. It is known that antiperipheral nerve myelin antibodies are associated with demyelination in Guillain-Barré syndrome. Vriesendorp et al showed that these antibodies and soluble terminal activation products were increased in the acute phase of neuralgic amyotrophy and decreased subsequently during clinical recovery. They concluded that complement dependent antibody mediated demyelination may participate in the initial phase of nerve damage or augment an ongoing process. By contrast, there remains minimal evidence to date favouring acute axonal destruction in response to a direct or indirect immunological insult, as seen in a few cases of axonal Guillain-Barré syndrome established by postmortem examination.

It can be deduced from our clinical and electrophysiological findings that although neuralgic amyotrophy is thought to be a condition predominantly characterised by axon loss, focal demyelination may play a significant part in its pathogenesis. Focal demyelination manifesting as conduction block may be detected as early as 2 weeks after the onset of the condition (patient 2) in the clinically most affected muscle. From here, the condition may takes two courses: resolution of conduction block with clinical improvement (patient 6) or progression to axonal degeneration with protracted clinical deficit (patient 2). It is possible that conduction blocks could be detected in more muscles if proximal stimulation studies were performed closer to the time of initial presentation. Hence, it is logical that therapeutic interventions should be directed to the group of patients showing electrophysiologically confirmed conduction block, with the aim of eradicating the block and possibly minimising subsequent axonal degeneration.

Most studies on neuralgic amyotrophy have described it as a self limiting condition with a good prognosis. In general, it is noted that 80% to 90% of patients would be expected to recover by 3 years, leaving 10% to 20% with considerable residual muscle weakness. There are several reports, however, of less favourable outcomes. A retrospective evaluation of 44 patients 1 to 7 years after the illness onset showed that 32 still had objective deficits and 17 experiencing intermittent complaints of radicular origins, particularly in patients with jobs of a manual nature. Recovery can begin very late in the illness, sometimes up to 6 months. With early and severe muscle wasting, the prospect of useful power returning can be poor. These cases are especially prone to late complications of scapulocostal pain syndromes, which can leave the patient with considerable morbidity. It might therefore be justifiable to identify these cases early with a view to therapeutic intervention such as intravenous immunoglobulin, which has been shown to be useful in other focal demyelinating diseases of the peripheral nerve.
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