Porphyric Neuropathy: An Ultrastructural and Quantitative Case Study

P.S. THORNER, J.M. BILBAO, A.A.F. SIMA and S. BRIGGS

SUMMARY: We report a case of acute neuropathy in a 46 year old female with porphyria variegata. Histologic, electron microscopic, and quantitative examinations of peripheral nerves were performed at onset of the neuropathy and at autopsy. The results revealed severe qualitative and quantitative changes in myelinated and unmyelinated fibers showing features indicative of an axonopathy with a distribution in keeping with a dying-back phenomenon.

RÉSUMÉ: Nous rapportons un cas fatal de neuropathie aigue chez une femme de 46 ans avec porphyrie variegata. Des examens histologiques, de microscopie électronique et quantitatifs des nerfs périphériques furent obtenus au début de la neuropathie et à l'autopsie. On montra ainsi des changements de structure importants et quantitatifs au niveau des fibres myélinisées et non myélinisées. Ces changements indiquent une axonopathie compatible avec le phénomène du "dying-back".

INTRODUCTION

The hepatic porphyrias are characterized by acute attacks of neurological dysfunction which often result in the patient's death (Goldberg, 1959; Eales & Linder, 1962; Eales, 1963; Bloomer, 1976). The clinical presentation includes colicky abdominal pain, neuropsychiatric disorders, and peripheral neuropathy. After a single attack the neuropathy may be reversible whereas repeated or prolonged attacks cause irreversible peripheral nerve damage (Berg, 1945; Flügel and Kruschky, 1977; Wochnik-Dyjas, et al, 1978).

The time course of the neuropathy may be rapidly progressive, reaching flaccid paralysis in days (Goldberg, 1959), or proceed at a slower pace evolving over months. There are conflicting reports regarding the basic pathology of porphyric neuropathy. Early authors (Denny-Brown & Sciarra, 1945; Gibson & Goldberg, 1956; Campbell, 1963), supported segmental demyelination as the primary process whereas recent investigators like Cavanagh and co-workers (Cavanagh & Mellick, 1965; Cavanagh & Ridley, 1967; Schoental & Cavanagh, 1977), and others (Sweeney, et al, 1970; Tschudy, et al, 1975), suggest that axonal degeneration is the basic structural change in porphyric neuropathy.

This report describes the qualitative and quantitative changes in a sural nerve biopsy and in post mortem material obtained from the tibial nerve, ventral and dorsal sacral roots.

CASE HISTORY

The patient was a 46 year old Caucasian woman with a past history of depression (1973, 1977) and abdominal pain with diarrhea in 1978. She was on no medica-

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TABLE 1

Results of Laboratory Investigations

<table>
<thead>
<tr>
<th></th>
<th>BLOOD</th>
<th>URINE</th>
<th>STOOL</th>
</tr>
</thead>
<tbody>
<tr>
<td>δ-ALA</td>
<td>0.69 mg% (N &lt; 0.30)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Porphobilinogen</td>
<td>POSITIVE (N = negative)</td>
<td>POSTIVE (N = negative)</td>
<td>-</td>
</tr>
<tr>
<td>Uroporphyrins</td>
<td>250 mg/d (N &lt; 30)</td>
<td>24mg/ day (N = 0)</td>
<td>143 mg% (N &lt;100)</td>
</tr>
<tr>
<td>Coproporphyrins</td>
<td>803 µg/d (N=75-275)</td>
<td>265mg/ day (N=50-250µg/day)</td>
<td>1384 mg% (N &lt; 700)</td>
</tr>
<tr>
<td>Protoporphyrins</td>
<td>-</td>
<td>-</td>
<td>2050 mg% (N &lt; 1500)</td>
</tr>
</tbody>
</table>

CSF examination, CT scan of the head and a myelogram were all normal. A diagnosis of porphyria variegata was considered and confirmed by laboratory studies (Table 1).

Three distant female relatives had experienced symptoms suggestive of porphyria but there was no biochemical confirmation of the diagnosis.

During her hospital course the patient suffered three bouts of aspiration pneumonia which were treated with broad spectrum antibiotics. In late November she developed increasing agitation and systemic candidiasis. She died on November 25, 1979.

At autopsy systemic candidiasis involving the kidneys, myocardium, brain and the spinal cord was noted. Bilateral aspiration pneumonia was seen. The relevant neuropathology is described in detail.

MATERIAL AND METHODS

The sural nerve obtained by biopsy, and ventral and dorsal S1 roots and tibial nerve dissected at post-mortem examination, 7 hours after death, were fixed in cacodylate buffered 2% glutaraldehyde, post-fixed in Millonig-buffered 1% osmium tetroxide, dehydrated and embedded in Epon-Araldite. Cross and longitudinal 1 micron thick sections were stained with toluidine blue for light microscopy. Ultra-thin sections were stained with uranyl acetate and lead citrate and examined under the electron microscope. In addition, spinal root ganglia, spinal cord and skeletal muscle were fixed in neutral formaldehyde and examined under the light microscope.

Caliber-spectra and fiber densities of myelinated fibers were calculated from photomicrographs at a final magnification of 1000. In each nerve all myelinated fibers of one randomly chosen fascicle were counted. Caliber-spectra and fiber densities of unmyelinated fibers were calculated from unselected electron micrographs with a final magnification of 15,000. The axon-myelin ratios were examined in randomly chosen fibers, which showed no axon or myelin destruction, at a final magnification of 15,000, using the methods of Robertson and Sima (1980). Linear regressions for the relationship between the natural log. of axis cylinder area and myelin thickness as measured by the number of lamellae were calculated (cf. Dyck et al, 1981).

Control nerves were obtained 8 hours post-mortem from a 52 year old woman who died without neurological disease. The morphometric measurements were performed on a Hewlett-Packard 9874A Digitizer interfaced with a Hewlett-Packard 9825A Computer.

RESULTS

Structural Changes:

Sural Nerve Biopsy: Light microscopic examination of thick sections revealed a marked reduction in the number of large myelinated fibers (Fig. 1). Electron microscopy showed extensive axonal degenerative changes consisting of dissolution of the axoplasmic organelles, granular degeneration of the axoplasm and ingrowth of the inner Schwann cell lip (Fig. 2), sometimes forming honeycomb structures at the axon-Schwann cell interface. Many of the affected fibers showed well preserved myelin sheaths. Other fibers revealed a later stage of axonal degeneration, namely complete disintegration of the axon (Fig. 3), collapse and fragmentation of the myelin and the development of denervated Schwann cells (Fig. 3). Some of the smaller myelinated fibers were uninvolved. Unmyelinated fibers showed axonal fragmentation and retraction of Schwann cell cytoplasm leaving axons directly apposed to basement lamina (Fig. 4). Large unmyelinated fibers frequently demonstrated a separation of neurofilaments and tubules with the neurofilaments in a central position (Fig. 5).

Tibial Nerve: Light microscopic examination of thick sections showed depletion of large myelinated fibers and a large number of free foamy macrophages in the endoneurium (Fig. 6). In the electron microscope small-sized myelinated fibers showed marked axonal atrophy. Others contained accumulation of membranous lipid structures in the axons. Clusters of regenerating units consisting of axons with thin myelin sheaths and duplicated basement membranes were occasionally found. The changes in the unmyelinated fibers were similar to those described in the sural nerve. In particular, there was a large number of single axon-Schwann cell units with pronounced Schwann cell banding. Occasionally unmyelinated fibers contained dense accumulations of misdirected 10nm neurofilaments which caused axonal distention (Fig. 7).

Ventral and Dorsal roots: Changes in the spinal roots were less severe than in the tibial and sural nerves. Many myelinated fibers appeared normal, whereas others showed complex mem-
branous Schwann cell processes invading the axons, and accumulation of organelle debris in the axons. Axonal atrophy was marked, particularly in large sized fibers. In addition, the ventral root contained numerous regenerating units. (Fig. 8).

QUANTITATIVE CHANGES

Caliber Spectra:
Myelinated fibers: The tibial and the sural nerves and the unifascicular spinal roots showed marked shifts of their caliber-spectra toward thinner diameters (Fig. 9). These shifts were more marked in the tibial and sural nerves and less marked in the ventral and dorsal roots. The fiber densities were decreased in the sural and tibial nerves, whereas they were increased in the spinal roots, particularly in the

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ventral root. The total fiber number showed an absolute decrease in the dorsal root and a slight increase in the ventral root (Table 2).

Unmyelinated fibers: Unmyelinated fiber sizes were shifted toward thinner diameters in both the tibial and the sural nerves. The tibial nerve showed a small population (approximately 2.5%) of unmyelinated fibers with larger diameters than were found in the control nerve (Fig. 9). These fibers represent in part those exhibiting axonal swelling due to dense accumulations of neurofilaments. The densities of unmyelinated fibers per area unit was markedly increased in the tibial nerve and marginally increased in the sural nerve (Table 2).

Axon-myelin ratio: Both the ventral root and the tibial nerve showed significantly decreased ratios between the axonal area and the number of myelin lamellae (Fig. 10), indicating severe axonal atrophy.

Light microscopic examination: Dorsal root ganglia showed mild loss of neurons. The ventral horns of the
spinal cord showed moderate neuronal loss and chromatolysis of preserved neurons. In skeletal muscle, grouped fiber atrophy was seen.

DISCUSSION
The present study has demonstrated severe qualitative and quantitative changes in myelinated and unmyelinated nerve fibers in a patient with porphyric neuropathy. The sural nerve biopsy obtained during the early course of the disease revealed acute degenerative changes such as axonal sequestration by the inner Schwann cell lip and granular degeneration of the axoplasm in myelinated axons. These changes were accompanied by only minor structural changes of the Schwann cell and myelin sheath and are therefore indicative of a primary axonal involvement. More severe changes consisted of complete axonal disintegration and Wallerian degeneration. The caliber-spectrum analysis demonstrated that large myelinated fibers were especially involved in the disease process. In contrast, Anzil and Dozic (1978), found small-sized fibers to be mainly involved in porphyric neuropathy.

The examination of the tibial nerve 9 weeks later revealed less acute degenerative changes. There was marked axonal atrophy, (objectively demonstrated by the axon-myelin ratio), numerous free foamy macrophages and a large number of regenerating units, indicating a more longstanding process. In the spinal roots, the axonal degenerative and atrophic changes were less obvious, although the ventral root revealed marked regenerative activity. This was also reflected by 1) a slight absolute increase in the number of myelinated fibers, presumably representing remyelination of sprouting fibers, and 2) a significantly increased fiber density of the ventral root, since more small remyelinating fibers can be harbored per area unit. The presence of occasional degenerating neurons in the spinal ganglia and the ventral horns suggests that the disease process had progressed to the cell somata.

The qualitative changes and the topographic difference in severity of quantitative changes in the peripheral nerves is suggestive of a primary axonopathy of dying-back type. (Schoental & Cavanagh, 1977; Spencer & Schaumburg, 1977; Cavanagh, 1979; Sima, et al, 1981). Early investigators (Denny-Brown & Scirra, 1945; Gibson & Goldberg, 1956; Campbell, 1963) have suggested that demyelination is the primary disease process in porphyric neuropathy. In the present study demyelination was certainly a prominent pathologic feature, although this must be interpreted as being secondary to axonal degeneration, based on the results of more recent investigative techniques used here such as quantitative morphology and ultrastructure.

One might however argue that the axonal atrophy could possibly be due to disuse during the patient’s hospitalization (Eisen et al, 1973). However this seems unlikely because of the sudden onset of the patient's neuromuscular complaints and the rapid progression to a flaccid areflexic quadriplegia.

The structural changes reported in the present case show similarities with those recently reported in mice after the injection of a synthetic porphyrin, tetrathenylporphinesulfonate (Sima, et al, 1981). In this experimental model, peripheral motor nerves exhibited progressive axonal degenerative changes with secondary demyelination, evolving from the motor nerve terminals (Felix & Sima, 1981), to the ventral roots (Sima, et al, 1981).

Qualitative and quantitative changes of unmyelinated fibers in porphyric neuropathy have not previously been reported, although the clinical picture of acute porphyria suggests that this fiber population is most likely involved in the disease process (Tschudy, et al, 1975; Becker & Kramer, 1977). In the present case, unmyelinated fibers showed axonal degeneration with sequestration of axonal material and marked atrophy. Similar structural changes of unmyelinated fibers have

### TABLE 2

<table>
<thead>
<tr>
<th>Fiber Type</th>
<th>Fiber Number/Fascicle</th>
<th>Fiber Density (Fibers/mm²)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Ventral Root</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>P</td>
<td>4436</td>
<td>14,786</td>
</tr>
<tr>
<td>C</td>
<td>3655</td>
<td>4,641</td>
</tr>
<tr>
<td><strong>Dorsal Root</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>P</td>
<td>4502</td>
<td>9,829</td>
</tr>
<tr>
<td>C</td>
<td>9741</td>
<td>8,776</td>
</tr>
<tr>
<td><strong>Tibial Nerve</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>P</td>
<td>885</td>
<td>5,205</td>
</tr>
<tr>
<td>C</td>
<td>2384</td>
<td>6,425</td>
</tr>
<tr>
<td><strong>Sural Nerve</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>P</td>
<td>409</td>
<td>2,133</td>
</tr>
<tr>
<td>C</td>
<td>1807</td>
<td>6,146</td>
</tr>
<tr>
<td><strong>Unmyelinated Fibers</strong></td>
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<td></td>
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<tr>
<td>Tibial Nerve</td>
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<td></td>
</tr>
<tr>
<td>P</td>
<td>549</td>
<td>115,336</td>
</tr>
<tr>
<td>C</td>
<td>358</td>
<td>40,498</td>
</tr>
<tr>
<td>Sural Nerve</td>
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<td></td>
</tr>
<tr>
<td>P</td>
<td>333</td>
<td>69,957</td>
</tr>
<tr>
<td>C</td>
<td>367</td>
<td>61,319</td>
</tr>
</tbody>
</table>

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previously been reported in experimental diabetes (Sima & Robertson, 1979; Sima, 1980). Many unmyelinated fibers were found to be denuded from the ensheathing Schwann cells, a finding that has been reported in painful diabetic neuropathy (Brown, et al, 1976), and in different models of experimental diabetic neuropathy (Sima & Robertson, 1979; Sima, 1980). An interesting observation was a small population of unmyelinated fibers that exhibited axonal swelling due to dense accumulations of neurofilaments. This finding is similar to what is encountered in myelinated fibers in the experimental model referred to previously (Sima, et al, 1981). This observation may indicate that disturbed axonal transport could be of pathogenetic significance in the development of porphyric neuropathy (Schoental & Cavanagh, 1977; Cavanagh, 1979; Felix & Sima, 1981; Sima, et al, 1981).

Although the mechanism(s) responsible for porphyric neuropathy is (are) poorly understood, it is generally felt that the precursors to porphyrins, delta-ALA and/or PBG are responsible for the neuropathy (Kramer, et al, 1973; Shanley, et al, 1975; Brodie, et al, 1976; Sima, et al, 1981), since these substances are elevated only in those porphyrias which show neurological manifestations. It has been suggested that these substances may inhibit enzymes and cofactors (Cavanagh &
Ridley, 1967; Schoental & Cavanagh, 1977; Wochnik-Dyjas et al, 1978; Cavanagh, 1979; Sima et al, 1981), essential for the functional and structural integrity of the axon. In porphyria variegate a defect in ferrochelatase gives it the characteristic delta-ALA and PBG (Brodie, et al, biochemical profile of elevated porphyrins and secondary elevation of delta-ALA and PBG (Brodie, et al, 1977; Becker and Kramer, 1977; Becker, et al, 1977). In summary, the present study has demonstrated an axonal neuropathy affecting distal nerves more severely than proximal ones in a patient with porphyria variegate. Furthermore, severe structural and quantitative changes of unmyelinated fibers have been demonstrated for the first time in porphyric neuropathy.

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REFERENCES


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