Motor neuropathy in porphobilinogen deaminase–deficient mice imitates the peripheral neuropathy of human acute porphyria

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Acute porphyrias are inherited disorders caused by partial deficiency of specific heme biosynthesis enzymes. Clinically, porphyrias are manifested by a neuropsychiatric syndrome that includes peripheral neuropathy. Although much is known about the porphyrias’ enzyme defects and their biochemical consequences, the cause of the neurological manifestations remains unresolved. We have studied porphyric neuropathy in mice with a partial deficiency of porphobilinogen deaminase (PBGD). PBGD-deficient mice (PBGD–/–) imitate acute porphyria through massive induction of hepatic δ-aminolevulinic acid synthase by drugs such as phenobarbital. Here we show that PBGD–/– mice develop impairment of motor coordination and muscle weakness. Histologically, femoral nerves of PBGD–/– mice exhibit a marked decrease in large-caliber (>8 μm) axons and ultrastructural changes consistent with primary motor axon degeneration, secondary Schwann cell reactions, and axonal regeneration. These findings resemble those found in studies of affected nerves of patients with acute porphyria and thus provide strong evidence that PBGD deficiency causes degeneration of motor axons without signs of primary demyelination, thereby resolving a long-standing controversy. Interestingly, the neuropathy in PBGD–/– mice developed chronically and progressively and in the presence of normal or only slightly (twofold) increased plasma and urinary levels of the putative neurotoxic heme precursor δ-aminolevulinic acid. These data suggest that heme deficiency and consequent dysfunction of hemeproteins can cause porphyric neuropathy.

logical systems in vitro, including interactions with γ-aminobutyric acid (GABA) receptors (8) and free radical formation (9). However, many of these effects occur at relatively high concentrations that are unlikely to exist in tissues during acute attacks. Arguments against the ALA hypothesis are the high excretion of ALA in many asymptomatic patients and the lack of clinical signs of neurological dysfunction after ALA administration in healthy volunteers or patients with acute porphyria (reviewed in ref. 4).

Extensive studies in rodents involving oral and parenteral administration of ALA for up to several weeks also failed to produce significant toxic, behavioral, or pharmacological effects (10). However, ALA given in the setting of normally regulated heme synthesis (i.e., without partial deficiency of a specific enzyme) may not reflect the pathophysiological situation of porphyria. Therefore, the question whether ALA can produce some, or even all, of the manifestations of acute porphyria has remained a controversial issue. Connected with this uncertainty is the unresolved question regarding actual ALA concentrations in the central and peripheral nervous systems or to what extent ALA overproduced in the liver may cross the blood-brain barrier (11).

In the present study, we have used behavioral tests, light and electron microscopy, and electrophysiological techniques to examine the development of peripheral neuropathy of PBGD−/− mice of different ages. Our data document a chronic and progressive axonal motor neuropathy that morphologically resembles the neuropathy of porphyria patients. Further experiments were carried out to determine the concentration of ALA in brain and peripheral tissue in PBGD−/− mice. Our data reveal that the motor neuropathy in PBGD−/− mice develops at normal or only slightly increased ALA concentration, making neurotoxicity of ALA an unlikely mechanism of porphyric neuropathy.

Methods

Animals and treatments. PBGD-deficient mice used in this study have previously been described (5). PBGD was partially inactivated by gene targeting so that one allele of PBGD had a loss of function disruption and the other allele a partial disruption. Compound heterozygous mice (designated PBGD-deficient, or PBGD−/−) have 25–30% of normal PBGD activity. Age- and gender-matched normal C57BL/6 wild-type (+/+ ) mice were used as controls. Mice were maintained in a 12-hour light/12-hour dark cycle and had free access to standard rodent laboratory chow and drinking water. The animals were studied at various ages: old (15.5–20.5 months), middle (six months), and young (two to three months). To biochemically imitate a human porphyric attack, mice were treated with increasing doses (100, 110, 120, and 125 mg/kg) of phenobarbital intraperitoneally for four consecutive days (5).

Behavioral studies. Standard behavioral tests were selected to evaluate motor function in PBGD+/− mice (12). The rotorod test measures motor coordination and proprioceptive sensitivity. It determines the time (in seconds) that mice can stay on a rotating dowel turning at 12 rpm. In addition, the motor coordination was studied on an inclined plane, where the angle at which mice can no longer remain on the plane is determined. The walk-on-a-rod test evaluates the time required (in seconds) to cross a 40-cm-long rod 10 mm in diameter. The hot plate test examines the escape reaction: the time (in seconds) that elapses until the mouse jumps out of a glass cylinder (17 cm high and 9 cm in diameter) with the base plate at 48°C. Muscular endurance was quantified by determination of the time (in seconds) an animal keeps hold of a weight (32 g) with two of its legs when lifted by the tail. Maximal strength (in newtons) was measured with the mouse pulling a piece of wire connected to an isometric transducer. Pain threshold was determined by the tail-flick test and the occurrence of the paw-lick response during hot plate tests. In the tail-flick test, an intense light beam is focused on the tail, and the time (in seconds) that passes until the animal flicks its tail is determined.

Light and electron microscopy. The femoral nerve (comprising branches of the quadriceps muscle and the cutaneous saphenous nerve) was processed for light and electron microscopy as recently described (13, 14). Snap-frozen and unfixed cryosections (6 μm) were subjected to indirect immunoperoxidase staining with antibodies against the myelin-specific protein zero (P0) (P0, mAb anti-P0, clone D41E4 provided by S.L. Miller, Children’s Hospital, Philadelphia, Pennsylvania, USA; ref. 15) and against the glial fibrillar acidic protein (GFAP) (GFAP, polyclonal rabbit anti-GFAP purchased from DAKO A/S, Golstrup, Denmark) and included negative controls for the secondary peroxidase–labeled antibodies (goat anti-mouse IgG [Fc] P0-Ab and goat anti-rabbit IgG [whole molecule] P0-Ab; Sigma Chemical Co., St. Louis, Missouri, USA). Counts of myelinated axons were performed on micrographs of toluidine blue-stained semi-thin plastic sections (1 μm) of the quadriceps muscle branch of the femoral nerve at a final magnification of ×800.

Electrophysiology. Investigation of nerve conduction characteristics of facial and sciatic nerves in 21 (15.5–20.5 months old) PBGD−/− mice and 16 gender- and age-matched control mice were performed with methods described previously (16). Electromyography in five PBGD−/− and four control mice was done with a Tönnies digital electromyograph (Tönnies, Würzburg, Germany) with a concentric needle electrode. Motor unit action potentials (MUAPs) were elicited by gently pressing on the opposite limb to induce a withdrawal response. Microdialysis and ALA measurements. Microdialysis (17) allows continuous monitoring of substances from the extracellular space in tissues and organs in vivo with minimal perturbation of the physiological system, i.e., without contamination by blood. The microdialysis probe (CMA12; shaft length 6 mm, diameter 0.5 mm, membrane length 2 mm, pore size 20 kDa; Carnegie Medicine AB, Stockholm, Sweden) was placed by a stereotaxic device into the striatum to estimate central nervous system concent-

Table 1

<table>
<thead>
<tr>
<th>Test</th>
<th>Mouse</th>
<th>Age (months)</th>
</tr>
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<tbody>
<tr>
<td>Rotorod</td>
<td>PBGD−/−</td>
<td>151.2 ± 29.3</td>
</tr>
<tr>
<td></td>
<td>control</td>
<td>180^a</td>
</tr>
<tr>
<td>Walk on rod</td>
<td>PBGD−/−</td>
<td>3.0 ± 0.5</td>
</tr>
<tr>
<td></td>
<td>control</td>
<td>2.2 ± 0.2</td>
</tr>
<tr>
<td>Inclined plane (°)</td>
<td>PBGD−/−</td>
<td>80.2 ± 2.2</td>
</tr>
<tr>
<td></td>
<td>control</td>
<td>88.4 ± 5.0</td>
</tr>
<tr>
<td>Escape reaction (s)</td>
<td>PBGD−/−</td>
<td>9.0 ± 2.8</td>
</tr>
<tr>
<td></td>
<td>control</td>
<td>10.0 ± 1.1</td>
</tr>
<tr>
<td>Muscular endurance (N)</td>
<td>PBGD−/−</td>
<td>51.8 ± 4.0</td>
</tr>
<tr>
<td></td>
<td>control</td>
<td>56.8 ± 2.0</td>
</tr>
<tr>
<td>Maximal strength (N)</td>
<td>PBGD−/−</td>
<td>0.025 ± 0.002</td>
</tr>
<tr>
<td></td>
<td>control</td>
<td>0.028 ± 0.002</td>
</tr>
</tbody>
</table>

Data represent mean ± SEM (n = 5). PBGD−/− mice are compared with age-matched wild-type control mice. 

^aP < 0.001. *P < 0.01. +P < 0.05. ^45 s, time limit of the test.
The overall diameter of all quadriceps nerves investigated was reduced by 90% (from 139 to 14; Figure 1, e–g). ANOVA combined with the Scheffé post hoc test.

Electrophysiological and electromyographical data. Behavioral tests were analyzed by ANOVA combined with the Scheffé post hoc test.

**Results**

**Behavioral studies of motor function.** Motor dysfunction was evident in all behavioral tests (Table 1) in old PBGD–/– mice (age 16 months) compared with age-matched controls. In particular, in the rotarod test, old PBGD–/– mice were unable to stay on the turning rod and unable to escape from a cylinder placed on a hot plate (escape reaction). Motor disturbance was already demonstrable at six months for the rotarod, inclined plane, and maximal strength tests, as well as for the escape reaction, but for three-month-old mice the differences were not statistically significant.

Sensory deficits were not obvious in PBGD–/– mice of any age. Thus, pain threshold, measured with tail-flick or pawlick response during a hot plate test, was not different in PBGD–/– and control mice (data not shown).

**Light microscopy and immunohistochemistry of peripheral nerves.** The most striking finding in quadriceps nerves from 17-month-old PBGD–/– mice was a reduction in the number of large axons (diameter >8 μm). These large axons decreased by 90% (from 139 to 14; Figure 1, e–g). The overall diameter of all quadriceps nerves investigated also was strongly reduced in old PBGD–/– mice. At six months of age, the reduction of large-caliber axons was ~60% (from 141 to 53; Figure 1, c, d, and g). At this age, extended regions of the endoneurium were occupied by cellular profiles that were devoid of myelin (Figure 1d). Contorted myelin and myelin ovoids were also observed, indicating degenerative events. In quadriceps nerves from two-month-old PBGD–/– mice, neither loss of large-caliber axons nor myelin profiles suggesting degeneration were observed (Figure 1, a, b, and g). Immunohistochemistry with antibodies against the myelin component P0 confirmed the light microscopy findings in aged mice (Figure 2, a and c). A prominent feature was the increased expression of GFAP, an established marker of nonmyelinating and denervated Schwann cells (Figure 2).

**Electron microscopy.** Striking ultrastructural abnormalities were found in quadriceps nerves in PBGD–/– mice six and 17 months old. In addition to the reduced numbers of large-caliber fibers, the presence of basal lamina–covered Schwann cell profiles devoid of axons and myelin was the most frequent pathological alteration (Figure 3, a and b). These structures represent most probably denervated Schwann cells (bands of Büngner) and were often situated close to axon-Schwann cell units with normally shaped or thin myelin (Figure 3a). The bands of Büngner were occasionally associated with axonal growth cones (Figure 3c) or with nonmyelinated small-caliber axons (Figure 3d).

Another striking characteristic was the presence of myelinated axons with features suggesting axonal degeneration, such as overall high electron density, disorganized neurofilaments and neurotubules, and clusters of electron-dense material apposed to the axolemma (Figure 4, a and b). Axonal membranes were often discontinuous, and the periaxonal compartment often was poorly defined (Figure 4b). Absence of axonal mitochondria (Figure 4, a and b) was also frequently observed. Another typical feature was the occurrence of myelin-containing Schwann cells that were devoid of an axon and characterized by a contorted myelin structure reminiscent of Wallerian degeneration (Figure 4, c and d). Approximately 5–10% of myelinated axon-Schwann cell units exhibited the described degenerative phenotype. Nonmyelinating Schwann cells ensheathing bundles of small-caliber axons were also abnormally organized in the quadriceps nerve of six- and 17-month-old mice, as

![Figure 1](https://example.com/figure1.png)
reflected by slender cell processes protruding into the endoneurial space (Figure 4e). By contrast, both myelinated and nonmyelinated fibers appeared normal in the saphenous nerve, the major sensory cutaneous branch of the femoral nerve (Figure 4f).

In summary, the morphological findings are consistent with degeneration of large-caliber axons, secondary Schwann cell reactions, and axonal regenerations.

Electrophysiological investigations. We found a statistically significant decrease of the sciatic nerve motor response (M-response) amplitude in PBGD−/− mice (Figure 5a). All other sciatic nerve conduction studies were normal in the PBGD−/− mice compared with control mice (Table 2). The shape of the M-response was usually biphasic or triphasic. Slight temporal dispersion occurred in five PBGD−/− mice but in only one control mouse (Figure 5a). Four out of the five PBGD−/− mice with temporal dispersion had low M-response amplitudes (<10 mV). However, very low M-response amplitudes also occurred in six PBGD−/− mice and two control mice with no temporal dispersion. Mixed-afferent nerve conduction was normal in the sciatic nerve of PBGD−/− mice (Table 2). Facial nerve motor conduction studies, including the absolute refractory time, were normal in PBGD−/− mice (data not shown).

Electromyography studies were performed with the greater gluteal muscle, the quadriceps, and the gastrocnemius muscle in control and PBGD−/− mice. Our main finding was a significant increase of MUAP amplitudes in all investigated muscles of PBGD−/− mice (Table 2 and Figure 5c). The short duration of MUAPs and the rapid recruitment of muscle units precluded a detailed analysis of single MUAP duration. We detected polyphasic MUAPs in three out of five PBGD−/− mice (all in the greater gluteal muscle, three out of 93 MUAPs analyzed; Figure 5b), fibrillation in one PBGD−/− mouse (quadriceps muscle), and fasciculations in the greater gluteal and quadriceps muscles of one PBGD−/− mouse. These observations are likely to underestimate the true degree of abnormality because of the high background activity. No pathological spontaneous activity occurred in control mice. The recruitment pattern consisted of normal-appearing recruitment of MUAPs with an amplitude of 250–1,000 μV, with an interference pattern on maximal withdrawal in control mice. In PBGD−/− mice, however, early recruitment of high-amplitude MUAPs occurred on muscle contraction, and no dense interference pattern was obtained (Figure 5c). Also, the discharge frequency of motor units was slightly higher in PBGD−/− mice (range 30–140 Hz) compared with control mice (range 14–70 Hz).

These findings are characteristic of a chronic axonopathy predominantly affecting motor fibers. There is no electrophysiological evidence for primary demyelination or sensory nerve involvement.

ALA concentrations. Only a slight increase (approximately twofold) in the plasma concentration and urinary excretion of ALA was observed in PBGD−/− mice (Table 3). In ECF near the tibial nerve of hindlimbs, ALA concentrations were determined to be ~15–25% of the plasma levels in both PBGD−/− and control mice. In striatum of PBGD−/− and control mice, ALA was undetectable without drug administration. Drugs such as phenobarbital induce ALA synthase in the liver and precipitate acute attacks of porphyria. We therefore treated mice with phenobarbital to increase the concentration of plasma ALA to levels observed in patients during acute attacks of porphyria (2–13 × 10⁻⁶ M; Table 3) (11). As expected, phenobarbital treatment caused a massive (~50-fold) increase of ALA in plasma in PBGD−/− mice, whereas the increase in controls was only twofold (Table 3). Under these conditions, the brain (striatum) ECF concentration of ALA rose from undetectable to ~6 × 10⁻⁸ M in PBGD−/− mice and from undetectable to ~2 × 10⁻⁹ M in controls. The plasma-to-brain ratio of 120:1 and 250:1 suggests a high blood-brain barrier for ALA. Of special interest in this investigation is the fact that PBGD−/− mice developed motor neuropathy without any drug treatment.

Discussion

PBGD−/− mice develop progressive muscle weakness and loss of motor coordination with increasing age, as evidenced in behavioral tests. The disturbance of motor

![Figure 2](Image)

**Figure 2** Immunostaining of myelin with antibodies against P0 (red color) and GFAP (red color). Cryosections of the femoral nerve of a control mouse (a and b) and a mouse with PBGD deficiency (c and d), both at 16 months of age. Scale bar: 20 μm.
function is most likely caused by a continuous decrease in the number of larger-caliber axons in motor nerves, with a reduction of 60% and 90% at six and 17 months of age, respectively (Figures 1 and 2). The predominant affliction of the larger-caliber axons has also been noted in the few nerve biopsies performed in patients with porphyric peripheral neuropathy (6, 19). Apparently, degenerated axons have the ability to regenerate. The observation that the loss of large-caliber axons is higher than the decrease in the total number of axons can be explained by the regrowth and remyelination of axons that acquire and maintain a small-caliber phenotype (Figure 1g). The increased GFAP staining in the peripheral nerves of PBGD−/− mice (Figure 2, b and d) also is consistent with increased Schwann cell activity associated with regeneration of axons. A further argument in favor of axonal regrowth is the decreased recruitment and polyphasia of MUAPs in PBGD−/− mice. The increase of MUAP amplitudes also is consistent with enlargement of motor unit size due to reinnervation, as is the observation of axonal growth cones and axonal sprouts (Figure 3, c and d). Electromyographic findings in human porphyric neuropathy usually also reveal decreased recruitment of MUAPs shortly after onset of weakness (20, 21).

The conspicuous absence of pathological changes in the sensory saphenous nerve again matches the clinical experience in porphyria patients, as the peripheral neuropathy is chiefly a motor paralysis; sensory disturbances are minimal and late, if they appear at all (2, 4, 6, 22).

Our interpretation of the structural data is supported by the electrophysiological studies, including nerve conduction and electromyographic experiments in PBGD−/− mice. In particular, there were no signs of primary demyelination (temporal dispersion, slowing of nerve conduction, increase of latencies; Table 2 and Figure 5). Mild admixed demyelination, indicated by mild temporal dispersion, occurred only in the most severely affected PBGD−/− mice with a very low M-response. The most prominent electromyography (EMG) finding in PBGD−/− mice was a severe chronic neurogenic pattern with decreased MUAP recruitment, increased MUAP amplitudes, and polyphasia. Although we may not have achieved maximal innervation, we show that normal mice achieve a dense interference pattern under our experimental conditions, whereas PBGD−/− mice recruit high-amplitude MUAPs relatively early without achieving a dense interference pattern. This is due to a loss of motor

Table 2

<table>
<thead>
<tr>
<th>Nerve conduction and electromyography</th>
<th>Control (+/+)</th>
<th>PBGD−/−</th>
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<tbody>
<tr>
<td></td>
<td>(n = 18)</td>
<td>(n = 23)</td>
</tr>
<tr>
<td><strong>Nerve conduction of sciatic nerve</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Proximal stimulation</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Latency (ms)</td>
<td>1.5 ± 0.2</td>
<td>1.6 ± 0.3</td>
</tr>
<tr>
<td>Amplitude (mV)</td>
<td>14 ± 4</td>
<td>10 ± 5</td>
</tr>
<tr>
<td>F-latency (ms)</td>
<td>4.4 ± 0.6</td>
<td>4.7 ± 1.1</td>
</tr>
<tr>
<td>Distal stimulation</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Latency (ms)</td>
<td>0.9 ± 0.2</td>
<td>1.0 ± 0.3</td>
</tr>
<tr>
<td>Amplitude (mV)</td>
<td>17 ± 5</td>
<td>12 ± 5</td>
</tr>
<tr>
<td>F-latency (ms)</td>
<td>4.8 ± 0.6</td>
<td>4.9 ± 1.6</td>
</tr>
<tr>
<td>Motor nerve conduction</td>
<td>41 ± 12</td>
<td>41 ± 9</td>
</tr>
<tr>
<td>Velocity (ms)</td>
<td>56 ± 17</td>
<td>51 ± 12</td>
</tr>
</tbody>
</table>

The data are expressed as mean ± SD. *Difference of means between control and PBGD−/− mice; P < 0.05, double-sided U test.

Electromyography

<table>
<thead>
<tr>
<th>Muscle</th>
<th>Control (+/+)</th>
<th>PBGD−/−</th>
</tr>
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<tbody>
<tr>
<td></td>
<td>(n = 4)</td>
<td>(n = 5)</td>
</tr>
<tr>
<td>Gluteus maximus</td>
<td>425 µV (100–3,000; n = 50)</td>
<td>960 µV (200–6,000; n = 93)</td>
</tr>
<tr>
<td>Quadriceps</td>
<td>400 µV (180–1,400; n = 53)</td>
<td>970 µV (190–3,000; n = 38)</td>
</tr>
<tr>
<td>Gastrocnemius</td>
<td>390 µV (120–840; n = 41)</td>
<td>1,600 µV (500–3,390; n = 15)</td>
</tr>
</tbody>
</table>

The data are given as medians (range, number of motor units analyzed). *The differences of the medians between the control and PBGD−/− mice were significantly different in all three muscles; P < 0.05, double-sided U test.
units and corresponds to the lower compound muscle action potential (CMAP) amplitudes in the sciatic nerve conduction studies.

The results of these nerve conduction studies in PBGD−/− mice are consistent with electrophysiological findings in patients with porphyric neuropathy who suffer from an axonal neuropathy that predominantly affects motor fibers with little electrophysiological evidence of demyelination (6, 20–22). EMG findings in human porphyric neuropathy usually reveal decreased recruitment of MUAPs shortly after onset of weakness, with pathological spontaneous activity first appearing in proximal muscles and later spreading to distal muscles (20, 21).

The behavioral, structural, and electrophysiological data in PBGD−/− mice provide convincing evidence that porphyric neuropathy is primarily an axonal motor neuropathy and that the additional changes described most likely are the result of repeated events leading to degeneration and regeneration.

A number of hypotheses have been proposed to explain the pathogenesis of porphyric neuropathy, but none has been directly tested. Increased ALA synthase activity, accumulation of ALA, and heme deficiency in the liver are biochemical features common to all acute porphyrias. Explanations that link these biochemical changes to neuronal dysfunction implicate ALA as a neurotoxin and/or propose decreased functions of specific hemeproteins. The almost invariable association of clinical symptoms with high levels of ALA in serum and urine and the demonstration of in vitro effects of ALA in neurophysiological test systems, albeit usually observed at concentrations unlikely to occur in acute porphyria, led to the prevailing consideration of ALA as a neurotoxin. However, the results reported here reveal that highly increased ALA levels are not a prerequisite for peripheral neuropathy, as mice developed progressive neuropathology without high levels of ALA. Rather, the data suggest that heme deficiency in the liver or nervous system could lead to decreased levels of key hemeproteins such as mitochondrial cytochromes, tryptophan dioxygenase, nitric oxide synthase, or soluble guanylate cyclase in various tissues, resulting in indirect or direct effects in motor neurons.

Preliminary investigations of a number of hemeproteins support this possibility. Thus, mitochondria isolated from liver of PBGD−/− mice displayed decreased oxygen consumption presumably due to decreased function of hemeproteins of the respiratory chain, and tryptophan levels were elevated in the brain of PBGD−/− mice, although only after phenobarbital treatment (Lindberg, R.L.P., et al., unpublished observations), similar to studies reported by Litman and Correia (23) in a model of chemically induced porphyria in rats. On the other hand, extensive studies in our laboratory of heme-dependent dimerization, enzymatic activity, and induction of neuronal nitric oxide synthase did not reveal differences between PBGD−/− mice and controls (Lindberg, R.L.P., et al., unpublished observations). Thus, the role of heme deficiency and its differential effects on various hemeproteins in the generation of porphyric neuropathy require further study.

Many patients carrying the specific enzyme defects of acute porphyrias have no clinical symptoms. Clinical manifestations reflecting neuropathy frequently are precipitated by drugs, hormones, and fasting. These factors are known to induce or increase the inducibility of hepatic ALA synthase (1, 2, 6). After an acute attack, most symptoms are reversible.

However, in the animal model of acute porphyria described here, the neuropathy developed progressively with age and in the absence of high levels of ALA. In
PBGD–/– mice, therefore, axonal neuropathy most likely develops as a consequence of subacute or chronic heme deficiency. Mild chronic peripheral neuropathy was also documented by Mustajoki and Seppäläinen (24) in patients with latent porphyria, even in patients who never had acute symptoms, by decreased conduction velocity of slow motor fibers.

The animal model of acute porphyria described here thus shows similarities and differences to the human disease. For instance, although phenobarbital resulted in a massive induction of hepatic ALA synthase and consequent accumulation of plasma ALA to levels observed in patients during acute attacks (Table 3), mice did not manifest acute and reversible symptoms such as paralysis or seizures, and the motor disturbance was not visibly aggravated by the drug treatment. Moreover, there were no obvious signs of autonomic neuropathy such as tachycardia or a change in eating behavior in PBGD–/– mice, except for a slight but significant increase in blood pressure (Lindberg, R.L.P., unpublished observations).

The reasons for these discrepancies are unclear at this time. We consider it unlikely that the differences in phenotype are due to the lower activity of PBGD in mice (~30% of normal activity) compared with the 50% decrease in AIP patients. The V_{max} of PBGD in humans is normally about twice as high as the V_{max} (uninduced) of ALA synthase (25), making it the second rate-limiting step. In C57BL/6 mice, however, we found the PBGD activity to be four to five times higher than that of ALA synthase. In our model, therefore, the enzyme deficiency was adjusted to the species difference between mouse and human liver to produce the same drug-induced biochemical changes in ALA synthase induction and ALA accumulation. Since acute porphyria shows extremely variable penetrance and expressivity, PBGD–/– C57BL/6 mice may reflect only one of the possible phenotypes of the human disease.

In conclusion, we show that mice deficient in PBGD develop motor dysfunction and peripheral neuropathological features closely resembling those of patients with inherited acute porphyrias. The axonal neuropathy occurs in the absence of high levels of ALA. These data suggest that heme deficiency and consequent dysfunction of hemeproteins can cause porphyric neuropathy.
Acknowledgments

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