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Electrophysiologic evidence of polyneuropathy in a cat with signs of bilateral brachial plexus neuropathy

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**Case Description**—A 2-year-old spayed female domestic shorthair cat was examined because of bilateral thoracic limb weakness of acute onset.

**Clinical Findings**—Clinical signs included muscle atrophy, paresis, depressed spinal reflexes, hyperesthesia of the thoracic limbs, and reduced jaw muscle tone. Pelvic limb reflexes were normal. Results of a neurologic examination were suggestive of multifocal lesions involving both brachial plexuses and the trigeminal nerves. Abnormal nerve conduction across the brachial plexus and delayed late potentials were found on electrodiagnostic testing, and diffuse subclinical involvement of other regions of the peripheral nervous system was confirmed on the basis of abnormal electromyographic findings for the masticatory muscles and conduction block of the peroneal nerve.

**Treatment and Outcome**—No specific treatments were given, and neurologic signs resolved within a month. A relapse occurred 2 months after the first episode, with clinical signs affecting both the pelvic and the thoracic limbs on this occasion. Again, the condition resolved without specific treatment, and 13 months after the initial episode, the cat reportedly was normal.

**Conclusions and Clinical Relevance**—Findings suggested that brachial plexus neuropathy can be a multifocal disease in cats, even if clinically apparent neurologic deficits are initially subtle or absent, and that electrodiagnostic techniques can be used to identify subclinical involvement of the peripheral nerves. (J Am Vet Med Assoc 2009;234:240–244)

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**ABBREVIATION**

| CMAP | Compound muscle action potential |

A 2-year-old 4.2-kg (9.2-lb) neutered female domestic shorthair cat was referred to the Queen’s Veterinary School Hospital of the University of Cambridge 8 days after an acute onset of thoracic limb weakness. According to the owner, the cat initially had right thoracic limb lameness, but signs progressed within 2 days to bilateral thoracic limb weakness. The cat lived indoors and had been vaccinated 7 months previously. Results of a CBC, serum biochemical panel, and urinalysis performed prior to referral were unremarkable, as were results of serologic testing for evidence of exposure to feline coronavirus, *Toxoplasma* spp, FIV, and FeLV. No obvious lesions were seen on lateral radiographic views of the vertebral column.

On initial examination at the Queen’s Veterinary School Hospital, the cat had signs of bilateral thoracic limb paresis (ie, palmigrade stance and lack of extension of the elbow and shoulder joints), but pelvic limb motion appeared normal. Neurologic examination revealed marked abnormalities of postural reactions (ie, wheelbarrow, hopping, and hemiwalking) in both thoracic limbs, but pelvic limb postural reactions were normal. There was bilateral atrophy of the thoracic limb muscles, and the cat had severe signs of resentment in response to thoracic limb manipulation. Thoracic limb myotatic and flexor reflexes were markedly depressed, whereas pelvic limb reflexes were normal. Jaw tone, assessed by use of a standard method, was found to be reduced. The cutaneous trunci reflex was absent bilaterally. No signs of pain were evident during manipulation of the neck. Signs were considered to be most likely a result of lesions involving both brachial plexuses and the motor branches of the trigeminal nerves. A focal lesion of the gray matter between spinal cord segments C6 and T2 was considered a possibility but was thought to be unlikely because of the extensive loss of muscular function in the thoracic limbs, compared with the pelvic limbs, and a lesion in this location would not explain the decreased jaw tone.

Results of a follow-up CBC and serum biochemical profile were unremarkable, other than high serum creatine kinase activity (587 U/L; reference range, 49 to 151 U/L). Ten days after the first clinical signs were noticed by the owner, the cat was anesthetized with propofol and sevoflurane, and concentric-needle electromyography of the muscles of the head, proximal and distal muscle groups of the right and left thoracic and pelvic limbs, and muscles of the cervical, thoracic, and lumbar portions of the vertebral column was performed. Abnormal spontaneous electrical activity, including fibrillation potentials and positive sharp waves ranging from 1+ to 2+ (reference range, 0 to +), was recorded in all proximal and distal muscle groups in both thoracic limbs. Fibrillation potentials were also recorded in the masseter muscles (2+) and...
Table 1—Results of electrophysiologic testing in a cat with polyneuropathy and clinical signs of bilateral brachial plexus neuropathy.

<table>
<thead>
<tr>
<th>Variable</th>
<th>Tibial nerve</th>
<th>Peroneal nerve</th>
<th>Radial nerve</th>
<th>Ulnar nerve</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Affected cat</td>
<td>Reference values</td>
<td>Affected cat</td>
<td>Reference values</td>
</tr>
<tr>
<td>CMAP amplitude (mV)</td>
<td>24.4</td>
<td>20.9 ± 3.4</td>
<td>17.0</td>
<td>30.9 ± 6.6</td>
</tr>
<tr>
<td>Distal M-wave latency (ms)</td>
<td>4.6</td>
<td>29.0 ± 6.2</td>
<td>3.5</td>
<td>34.7*</td>
</tr>
<tr>
<td>Motor conduction velocity (m/s)</td>
<td>90</td>
<td>101.4 ± 12.9</td>
<td>109</td>
<td>88.3 ± 17.8</td>
</tr>
<tr>
<td>Sensory conduction velocity (m/s)</td>
<td>ND</td>
<td>80.2 ± 7.9</td>
<td>90</td>
<td>85.3 ± 6.8</td>
</tr>
<tr>
<td>M-wave residual latency (ms)</td>
<td>1.46</td>
<td>1.8 ± 0.4</td>
<td>0.42</td>
<td>1.9 ± 0.6</td>
</tr>
<tr>
<td>F-wave latency (ms)</td>
<td>12.9</td>
<td>9.5 ± 1.0</td>
<td>ND</td>
<td>NA</td>
</tr>
<tr>
<td>F ratio</td>
<td>5.45</td>
<td>1.75 ± 0.2</td>
<td>ND</td>
<td>NA</td>
</tr>
</tbody>
</table>

Reference values are given as mean ± SD and have been published previously.  
*Reference values were not available; reported values represent results of electrophysiologic testing in a clinically normal cat.  
NR = Not recorded. ND = Not determined. NA = Not available.
conversation 13 months after the first episode, the owner indicated that the cat appeared normal.

**Discussion**

Bilateral dysfunction of the brachial plexus is apparently uncommon in cats and dogs and to our knowledge, there is only one previously published report of brachial plexus neuropathy in a cat. Importantly, although the cat described in the present report had bilateral brachial plexus lesions, we were also able to identify subclinical involvement of other peripheral motor and sensory nerves through electrodiagnostic testing.

In the cat described in the present report, signs of pain were clearly elicited by manipulation of the thoracic limbs, and results of electrodiagnostic testing were suggestive of radial nerve sensory dysfunction. In contrast, signs of pain were not described in a previous report of a cat with brachial plexus neuropathy, although a report of a dog with brachial plexus neuropathy described intermittent periods of pain. Jaw tone in the cat described in the present report was reduced clinically, and this finding was confirmed by the abnormal electromyographic findings for the masseter muscles. Mild abnormalities were also seen during electromyography of the left cranial tibial muscle, and peroneal nerve conduction was abnormal, even though results of neurologic examination of the pelvic limbs were normal. In the previous report of a cat with brachial plexus neuropathy, results of electromyography were unremarkable and nerve conduction velocity of the tibial nerve was normal; however, the peroneal nerve was not tested. Interestingly, severe dyspnea developed in this cat and has also been identified in cats with acute idiopathic polyneuropathy. Because of the additional findings in the cat described in the present report, it is necessary to consider other causes of polyneuropathy, as well as the possibility of systemic disease.
not accurate to characterize the condition as a brachial plexus neuropathy. Rather, it appeared that the cat had an acute polyneuropathy with clinical signs of bilateral brachial plexus neuropathy.

Electromyographic findings and low CMAP amplitudes in the cat described in the present report suggested that the nerve lesions in this case were predominately a result of multifocal axonal loss. In the previous report, conduction velocity of the median nerve was markedly reduced (5.5 m/s), suggesting that lesions were the result of a demyelinating process. In the cat described in the present report, the temporal dispersion and loss of amplitude of CMAPs recorded following stimulation of the proximal portion of the radial nerve, compared with results obtained following stimulation of the distal portion of the radial nerve, suggested that there was involvement of the brachial plexus. However, it was difficult to determine whether this finding represented early demyelination, as this would imply severe involvement of large-diameter myelinated fibers and slow clinical recovery. Temporal dispersion is typically subclinical, and thoracic limb weakness in this cat was most likely attributable to disruption of axonal conduction. The prolonged duration and polyphasic morphology of the CMAPs obtained following stimulation of the proximal portion of the radial nerve can probably be explained by duration-dependent phase cancellation or focal axonal injury. Stimulation of the nerve roots proximal to the brachial plexus can be technically challenging, and amplitude of the recorded response depends to a great extent on the site of stimulation. The lack of proximity of the proximal stimulating electrodes to the nerve roots could have, in itself, caused substantial temporal dispersion. However, when the same recording technique was used in a clinically normal cat, good recordings of CMAPs were obtained after supramaximal stimulation of the C8 nerve root. On the other hand, electrodiagnostic testing of the peroneal nerve is much easier and accurate in small animals, and a true conduction block without temporal dispersion was recorded when results for the proximal stimulation site (trochanteric fossa) were compared with results for the distal stimulation site (stifle joint), which was considered suggestive of demyelination. In the pelvic limbs, only the left cranial tibial muscle had electromyographic abnormalities suggestive of axonal loss. In the early stages of peripheral nerve demyelinating processes associated with mild axonal loss, clinical signs (weakness) are lacking, consistent with findings for the cat described in the present report. Interestingly, the pelvic limbs were affected when the cat had a relapse of clinical signs, suggesting that more severe (axonal) lesions of peripheral nerves in the pelvic limb occurred during the second episode. Finally, values for F-wave latency and the F ratio give definitive evidence of proximal peripheral nerve involvement.

The etiology of brachial plexus neuropathy in animals is not known. In a cat described previously, vaccination 1 month prior to the onset of clinical signs was proposed as an inciting cause, and in a dog, an allergic mechanism was suggested. However, the reason why the brachial plexus should have been specifically involved early in the course of the disease in these animals was not explained. In some humans with neuralgic amyotrophy, which is also known as Parsonage-Turner syndrome, acute idiopathic brachial neuritis, and serum neuritis, lymphocytes have increased blastogenic activity when cultured with extracts of brachial plexus nerves but not when cultured with extracts of sacral plexus nerves, suggesting an immune-mediated response directed at epitopes specific to nerves derived from the brachial plexus may play a role. An immune-mediated response is also suggested by the presence of complement-dependent antibodies against peripheral nerve myelin, histologic evidence of mononuclear inflammatory infiltrates in the brachial plexus, and electron microscopic evidence of demyelinated axons and onion-bulb formation.

It was not possible to determine the underlying cause of the polyneuropathy in the cat described in the present report, but there probably are a limited number of peripheral nerve diseases in cats associated with spontaneous recovery, such as immune-mediated peripheral nerve diseases. Brachial plexus biopsy would have been necessary to fully characterize the condition, but was declined by the owner. Although biopsy of a peripheral nerve, such as the peroneal nerve, would have been less technically demanding than brachial plexus biopsy, results of histologic examination of nerve biopsy specimens from dogs with brachial plexus neuropathy have been nonspecific. On the other hand, examination of peripheral nerve biopsy specimens may have provided information regarding demyelination, remyelination, axonal loss, and whether there was an inflammatory component to the neuropathy.

Findings in the present case revealed that brachial plexus neuropathy can be a multifocal disease in cats, even if clinically apparent neurologic deficits are initially subtle or absent. This is of importance, as it may affect how veterinarians approach such cases. Electrodiagnostic testing was crucial in confirming the multifocal involvement in this case, although nerve and muscle biopsy would have been necessary to confirm the underlying etiology and pathologic abnormalities.

References

Selected abstract for JAVMA readers from the American Journal of Veterinary Research

Use of tick surveys and serosurveys to evaluate pet dogs as a sentinel species for emerging Lyme disease
Sarah A. Hamer et al

Objective—To evaluate dogs as a sentinel species for emergence of Lyme disease in a region undergoing invasion by *Ixodes scapularis*.

Sample Population—353 serum samples and 78 ticks obtained from dogs brought to 18 veterinary clinics located in the lower peninsula of Michigan from July 15, 2005, through August 15, 2005.

Procedures—Serum samples were evaluated for specific antibodies against *Borrelia burgdorferi* by use of 3 serologic assays. Ticks from dogs were subjected to PCR assays for detection of pathogens.

Results—Of 353 serum samples from dogs in 18 counties in 2005, only 2 (0.6%) contained western blot analysis–confirmed antibodies against *B burgdorferi*. Ten of 13 dogs with *I scapularis* were from clinics within or immediately adjacent to the known tick invasion zone. Six of 18 *I scapularis* and 12 of 60 noncompetent vector ticks were infected with *B burgdorferi*. No ticks were infected with *Anaplasma phagocytophilum*, and 3 were infected with *Babesia* spp.

Conclusions and Clinical Relevance—Serosurvey in dogs was found to be ineffective in tracking early invasion dynamics of *I scapularis* in this area. Tick chemoprophylaxis likely reduces serosurvey sensitivity in dogs. Ticks infected with *B burgdorferi* were more common and widely dispersed than seropositive dogs. In areas of low tick density, use of dogs as a source of ticks is preferable to serosurvey for surveillance of emerging Lyme disease.

Impact for Human Medicine—By retaining ticks from dogs for identification and pathogen testing, veterinarians can play an important role in early detection in areas with increasing risk of Lyme disease. (Am J Vet Res 2009;70:49–56)