Peripheral Neuropathy Evaluations of Patients With Prolonged Long COVID

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Abstract

Background and Objectives
Recovery from severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) infection appears exponential, leaving a tail of patients reporting various long COVID symptoms including unexplained fatigue/exertional intolerance and dysautonomic and sensory concerns. Indirect evidence links long COVID to incident polyneuropathy affecting the small-fiber (sensory/autonomic) axons.

Methods
We analyzed cross-sectional and longitudinal data from patients with World Health Organization (WHO)-defined long COVID without prior neuropathy history or risks who were referred for peripheral neuropathy evaluations. We captured standardized symptoms, examinations, objective neurodiagnostic test results, and outcomes, tracking participants for 1.4 years on average.

Results
Among 17 patients (mean age 43.3 years, 69% female, 94% Caucasian, and 19% Latino), 59% had ≥1 test interpretation confirming neuropathy. These included 63% (10/16) of skin biopsies, 17% (2/12) of electrodiagnostic tests and 50% (4/8) of autonomic function tests. One patient was diagnosed with critical illness axonal neuropathy and another with multifocal demyelinating neuropathy 3 weeks after mild COVID, and ≥10 received small-fiber neuropathy diagnoses. Longitudinal improvement averaged 52%, although none reported complete resolution. For treatment, 65% (11/17) received immunotherapies (corticosteroids and/or IV immunoglobulins).

Discussion
Among evaluated patients with long COVID, prolonged, often disabling, small-fiber neuropathy after mild SARS-CoV-2 was most common, beginning within 1 month of COVID-19 onset. Various evidence suggested infection-triggered immune dysregulation as a common mechanism.

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Go to Neurology.org/NN for full disclosures. Funding information is provided at the end of the article.

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Severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) can cause long-term disability (long COVID) with new neurologic manifestations after even mild infections. Reports of peripheral neuropathy include Guillain-Barré syndrome, mononeuritis multiplex, brachial plexitis, cranial neuropathies, and orthostatic intolerance, although some studies included patients with potentially contributory conditions. Various long COVID symptoms overlap with those of small-fiber polyneuropathy (SFN). Hence, we prospectively analyzed a cross-section of patients with long COVID evaluated for incident neuropathy.

### Methods

**Standard Protocol Approvals, Registrations, and Patient Consents**

This retrospective analysis was approved by the hospitals’ ethical review committee (1999P009042). Although participant consent was not required, all 17 provided verbal consent and 16 signed agreements for participation and publication of anonymized results.

**Study Design**

Inclusion required no known prior neuropathy or risks plus confirmation of SARS-CoV-2 infection according to

### Table 1 Participants, Objective Tests, and Treatments

<table>
<thead>
<tr>
<th>ID</th>
<th>Age (y)</th>
<th>Sex</th>
<th>Latino</th>
<th>Race</th>
<th>Day of initial EDX</th>
<th>EDX report</th>
<th>Day of initial skin biopsy</th>
<th>END lower leg</th>
<th>END upper thigh</th>
<th>Day of initial AFT</th>
<th>AFT interpretation</th>
<th>Therapy</th>
<th>Patient-reported improvement since nadir</th>
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<td>202</td>
<td>Diagnostic</td>
<td>Diagnostic</td>
<td>203</td>
<td>Borderline</td>
<td></td>
<td>50%</td>
</tr>
<tr>
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<td>F</td>
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<td>White</td>
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<td>236</td>
<td>Diagnostic</td>
<td>Diagnostic</td>
<td>189</td>
<td>Normal</td>
<td>Pred</td>
<td>0%</td>
</tr>
<tr>
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<td>F</td>
<td>No</td>
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<td>140</td>
<td>Normal</td>
<td>527</td>
<td>Diagnostic</td>
<td>(&lt;1.0%)</td>
<td>189</td>
<td>Normal</td>
<td>Pred</td>
<td>90%</td>
</tr>
<tr>
<td>4</td>
<td>37.0</td>
<td>F</td>
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<td>Mixed</td>
<td>272</td>
<td>Normal</td>
<td>299</td>
<td>Normal (60.2%)</td>
<td>Normal</td>
<td>189</td>
<td>Diagnostic (&lt;1.0%)</td>
<td>IVIg, pred, HC</td>
<td>70%</td>
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<tr>
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<td>293</td>
<td>Normal (95.3%)</td>
<td>230</td>
<td>Normal</td>
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<td></td>
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<td>35%</td>
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<td>409</td>
<td>Normal</td>
<td>354</td>
<td>Diagnostic</td>
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<td>40%</td>
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<tr>
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<td>41.1</td>
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<td>White</td>
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<td>Normal</td>
<td>409</td>
<td>Diagnostic</td>
<td>(&lt;1.0%)</td>
<td>597</td>
<td>Borderline</td>
<td>IVIg</td>
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<td>376</td>
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<td></td>
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</tr>
<tr>
<td>10</td>
<td>31.0</td>
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<td>No</td>
<td>White</td>
<td>63</td>
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<td>Normal</td>
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<td>Abnormal</td>
<td>IVIg, pred, MP</td>
<td>88%</td>
</tr>
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<td>Normal (54.5%)</td>
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<td></td>
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<tr>
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<td>White</td>
<td>215</td>
<td>Normal</td>
<td>213</td>
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<td>(&lt;1.0%)</td>
<td></td>
<td></td>
<td>IVIg</td>
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<td>44</td>
<td>Normal</td>
<td>Normal</td>
<td>49</td>
<td>Abnormal</td>
<td>IVIg, pred, MP</td>
<td>88%</td>
</tr>
<tr>
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<td>F</td>
<td>No</td>
<td>White</td>
<td>250</td>
<td>Diagnostic</td>
<td>236</td>
<td>Incomplete</td>
<td>Pred</td>
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<td></td>
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<tr>
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<td>White</td>
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<td>Abnormal</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>75%</td>
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</tbody>
</table>

Abbreviations: AFT = composite autonomic function testing; EDX = electrodiagnostic testing; END = skin biopsy epidermal neurite density; HC = hydrocortisone; IVIg = IV immunoglobulin therapy; MAGNET = Mass. General Neuropathy Exam Tool; MP = methylprednisolone; Pred = prednisone. EDX occurred on average on D258 ± 29 of illness; AFT occurred on average on D192 ± 17. Skin biopsy testing occurred on average on D264 ± 32; results are reported as % of predicted normative END if calculated by the laboratory (with ≤5% of predicted END diagnostic), or normal/diagnostic if laboratory had not performed statistical modeling. Patient 6’s nondiagnostic but abnormally high END suggested intraneuronal inflammation with exuberant axonal regeneration and their initially normal AFT was abnormal when repeated.
guidelines of the World Health Organization (WHO). COVID severity classification followed WHO guidelines. Inclusion required meeting the WHO definition of long COVID (onset of symptoms within 90 days of the first day of COVID symptoms that last for >2 months). Participants were enrolled upon COVID confirmation and neuromuscular referral before record review or most testing and treatment. Participants documented neuropathy symptoms via online REDCap surveys, and their neurologists documented standardized in-person and occasional telehealth neuropathy examinations. Because most participants had received symptom-relieving medications at varying doses, we analyzed only potentially preventive treatments, all of which were immunotherapies. Parametric analyses were used with variability represented by standard errors.

**Data Availability**
Any anonymized data not published within the article will be shared by request from any qualified investigator.

### Results

Among 17 patients with SARS-CoV-2 onset between February 21, 2020, and January 19, 2021, treated in 10 states/territories (Table 1), 16 had mild COVID. The one (#9) with severe COVID (1 month stay in intensive care with ventilatory support) had electrodiagnostically confirmed sensorimotor polyneuropathy ascribed to critical care illness in addition to SFN. Medical histories and comprehensive blood screening (not shown) identified none with conventional neuropathy risks nor evidence of systemic dysimmunity. Imaging of the brain or spine, if performed, was unrevealing.

Participants’ ages averaged 43.3 ± 3.3 years on COVID D1, and 68.8% were female; 18.8% were Latino, and 94.1% were Caucasian. Diagnostic tests for neuropathy (Table 1) revealed that 16.7% electrodiagnostic studies were abnormal, whereas 62.5% (10/16) of lower leg skin biopsies pathologically confirmed SFN, as corroborated by 50% of upper thigh biopsies and autonomic Table 2 Initial Symptom Scores

<table>
<thead>
<tr>
<th>ID</th>
<th>Time since COVID onset</th>
<th>Total symptom severity</th>
<th>Sensory severity</th>
<th>Cardiovascular severity</th>
<th>Gastrointestinal severity</th>
<th>Urinary/sexual severity</th>
<th>Miscellaneous severity</th>
<th>Pain score (0–10)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>14.9</td>
<td>58.8%</td>
<td>62.5%</td>
<td>50.0%</td>
<td>41.7%</td>
<td>31.3%</td>
<td>67.5%</td>
<td>6</td>
</tr>
<tr>
<td>2</td>
<td>3.0</td>
<td>67.6%</td>
<td>87.5%</td>
<td>54.2%</td>
<td>75.0%</td>
<td>50.0%</td>
<td>62.5%</td>
<td>8</td>
</tr>
<tr>
<td>3</td>
<td>14.5</td>
<td>19.1%</td>
<td>25.0%</td>
<td>12.5%</td>
<td>12.5%</td>
<td>0.0%</td>
<td>30.0%</td>
<td>2</td>
</tr>
<tr>
<td>4</td>
<td>14.0</td>
<td>25.0%</td>
<td>9.4%</td>
<td>8.3%</td>
<td>16.7%</td>
<td>0.0%</td>
<td>27.5%</td>
<td>1</td>
</tr>
<tr>
<td>5</td>
<td>3.9</td>
<td>22.8%</td>
<td>21.9%</td>
<td>29.2%</td>
<td>8.3%</td>
<td>0.0%</td>
<td>37.5%</td>
<td>1</td>
</tr>
<tr>
<td>6</td>
<td>8.9</td>
<td>39.7%</td>
<td>37.5%</td>
<td>16.7%</td>
<td>16.7%</td>
<td>50.0%</td>
<td>65.0%</td>
<td>6</td>
</tr>
<tr>
<td>7</td>
<td>6.7</td>
<td>61.8%</td>
<td>53.1%</td>
<td>50.0%</td>
<td>70.8%</td>
<td>37.5%</td>
<td>80.0%</td>
<td>1</td>
</tr>
<tr>
<td>8</td>
<td>0.1</td>
<td>22.1%</td>
<td>25.0%</td>
<td>8.3%</td>
<td>16.7%</td>
<td>25.0%</td>
<td>30.0%</td>
<td>1</td>
</tr>
<tr>
<td>9</td>
<td>3.0</td>
<td>37.5%</td>
<td>78.1%</td>
<td>20.8%</td>
<td>37.5%</td>
<td>6.3%</td>
<td>27.5%</td>
<td>8</td>
</tr>
<tr>
<td>10</td>
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<td>80.9%</td>
<td>100.0%</td>
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<td>91.7%</td>
<td>25.0%</td>
<td>90.0%</td>
<td>10</td>
</tr>
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<td>11</td>
<td>9.7</td>
<td>9.6%</td>
<td>12.5%</td>
<td>8.3%</td>
<td>0.0%</td>
<td>0.0%</td>
<td>17.5%</td>
<td>0</td>
</tr>
<tr>
<td>12</td>
<td>8.1</td>
<td>19.1%</td>
<td>46.9%</td>
<td>0.0%</td>
<td>0.0%</td>
<td>0.0%</td>
<td>27.5%</td>
<td>7</td>
</tr>
<tr>
<td>13</td>
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<td>62.5%</td>
<td>57.5%</td>
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<tr>
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<td>59.4%</td>
<td>45.8%</td>
<td>58.3%</td>
<td>87.5%</td>
<td>72.5%</td>
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<td>15</td>
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<td>15.6%</td>
<td>12.5%</td>
<td>8.3%</td>
<td>12.5%</td>
<td>30.0%</td>
<td>0</td>
</tr>
<tr>
<td>16</td>
<td>0.1</td>
<td>30.1%</td>
<td>21.9%</td>
<td>25.0%</td>
<td>33.3%</td>
<td>0.0%</td>
<td>50.0%</td>
<td>7</td>
</tr>
<tr>
<td>17</td>
<td>0.7</td>
<td>58.8%</td>
<td>50.0%</td>
<td>41.7%</td>
<td>66.7%</td>
<td>62.5%</td>
<td>70.0%</td>
<td>7</td>
</tr>
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<td>Mean</td>
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<td>26.5%</td>
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<td>SEM</td>
<td>1.3</td>
<td>5.3%</td>
<td>6.8%</td>
<td>4.8%</td>
<td>7.3%</td>
<td>6.7%</td>
<td>5.4%</td>
<td>0.9</td>
</tr>
</tbody>
</table>

Data represent summaries of participant’s initial responses to the REDCap online Small-fiber Symptom Survey, a comprehensive survey of symptoms associated with small-fiber polyneuropathy. Scores are reported as % of full scale (100% maximally present) where never = 0%, rarely = 25%, sometimes = 50%, often = 75%, and always = 100%. The sensory domain reflects positive and negative sensory symptoms and the cardiovascular domain including tachycardia and orthostatic symptoms plus extremity edema and color changes. Gastrointestinal symptoms include upper and lower dysfunction. The miscellaneous domain includes physical and mental fatigue, poor sleep, headaches, vision, hearing, sweating, movement/muscle concerns, and distal hair loss. See eTable 1, links.lww.com/NXI/A697, for scores for individual symptoms.
function tests. Initial SFN symptom scores (Table 2) were abnormal—reduced to 40.7% of ideal on average—with pain scores averaging 4.8/10. Initial neuromuscular examinations (Table 3) averaged 77.0% of ideal, with reduced/abnormal distal pin and vibration sensations and absent Achilles reflexes most prevalent. Participants 9 and 15 had distal muscle weakness and atrophy. Some patients were initially evaluated early in the course and others later, and investigations continued for months. Sixteen participants with 2020 onset had >1 year follow up, with the latest onset on 1/19/21. See Figure 1 (case 15) and eFigure 1, links.lww.com/NXI/A697, (case 13) for longitudinal details.

Treatments comprised corticosteroids in 35.3% (6/17) and IV immunoglobulins (IVIg) in 35.3% (6/17). Five were initially dosed at 2.0 g/kg/4 weeks and 1 at 1.6 g/kg/4 weeks. The 5 patients who received repeated IVIg, and their neurologists, reported benefit (e.g., Figure 1, Table 1). eFigure 1 reports patient 13’s graphed symptom and examination scores before and during IVIg. Patients’ impressions of recovery varied (averaging 51.8 ± 6.7%), reflecting varying illness severity, treatment status, and assessment timing.

### Discussion

Neuromuscular evaluations proved useful in most of these patients with long COVID. However some symptoms, exam changes and test results may have been false-negative, given that assessments were not often optimally timed (e.g., #6) and many patients reported care delays. This reported case of multifocal motor neuropathy (Figure 1) increases the spectrum of COVID-associated dysimmune neuropathies. Critical illness neuropathy—reported in approximately 10% of intubated patients with COVID—is attributed to various prolonged insults including intense inflammation and nerve compressions. Inherent study limitations include bias toward referrals for sensory neuropathy and underpowering. The initial evaluations reported occurred at varying times during the illness and treatment, whereas longitudinal assessments at standardized intervals are ideal for diagnostic and treatment decisions. Timing also complicates analysis of blood testing for immune markers (not shown). We screened patients with newly diagnosed neuropathy for all common established causes of distal sensory neuropathy, including routinely measuring ANA, ESR, IgG anti–SS-A/SS-B antibodies, and complement

### Table 3 Neuropathy Examination Scores

<table>
<thead>
<tr>
<th>ID</th>
<th>Timing of initial examination (mo)</th>
<th>Total normality</th>
<th>Vital sign normality</th>
<th>Leg/toe appearance</th>
<th>Toe strength</th>
<th>Toe position sensation</th>
<th>Toe vibration sensation</th>
<th>Leg/toe light touch sensation</th>
<th>Leg/toe pin sensation</th>
<th>Achilles reflexes</th>
</tr>
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<tr>
<td>1</td>
<td>14.9</td>
<td>53%</td>
<td>50%</td>
<td>100%</td>
<td>100%</td>
<td>100%</td>
<td>0%</td>
<td>33%</td>
<td>20%</td>
<td>0%</td>
</tr>
<tr>
<td>2</td>
<td>6.0</td>
<td>94%</td>
<td>100%</td>
<td>100%</td>
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<td>100%</td>
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</tr>
<tr>
<td>8</td>
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Data represent initial neurologists’ responses to the MAGNET (Mass. General Neuropathy Exam Tool).5 Scores are reported as % of full scale (normal examination findings).
components C3 and C4, the most productive markers of dysimmunity in initially idiopathic SFN. We did not detect evidence of Sjögren syndrome, and other inflammatory markers were only occasionally elevated. Interpretation is complex as early elevations could be nonspecifically associated with acute COVID, and many months later, inflammation and markers might have subsided leaving residual axonopathy as the proximate cause of current symptoms. Regeneration can take up to 2 years or be incomplete.

These results identify small-fiber neuropathy as most prevalent in this small group of patients with long COVID, also known as post-acute sequelae of SARS CoV-2 infection. In SFN, the small-diameter unmyelinated and/or thinly myelinated sensory and autonomic fibers are predominantly affected, although most patients with severe or advanced polyneuropathy, e.g., case 9, develop large- and small-fiber damage. The small fibers are disproportionately vulnerable, with their lack of myelin exposing them to environmental stressors including immunity, while inability to use saltatory conduction increases metabolic demand, and cytoplasmic paucity limits axonal regeneration. However, small-fiber axons grow throughout life to reinnervate continuously dividing tissues such as the skin and to help repair injuries. If toxic conditions improve, axon elongation and sprouting accelerate to increase the probability of reinnervating enough target cells to resolve symptoms.

Here, most patients treated with sustained IVIg, the primary treatment for inflammatory neuropathy, with preliminary evidence of effectiveness for dysimmune SFN, perceived improvement (e.g., Figure 1, eFigure 1, links.lww.com/NXI/A697). Some treated only with corticosteroids did as well; participant 3 reported that prednisone helped her toward 90% improvement and was discontinued only because of adverse effects. Others improved substantially without immunotherapy (e.g., case 17), documenting spontaneous recovery and need to individualize treatment decisions.

The hypothesis that some long COVID symptoms reflect underlying small-fiber pathology is supported by research observation of small-fiber loss applying in vivo corneal confocal microscopy to patients with long COVID. As with other post-COVID neurologic illnesses, susceptibility to inflammatory mediators appears essential. Autopsy study of post-COVID patients identified neuritis with perivascular macrophage infiltrates but no viral antigens, implicating inflammatory immune responses rather than direct infection. In addition, 1/4th of human DRG neurons express mRNA for SARS-CoV-2–associated receptors and deploy ACE2 protein. Thus, virus or spike protein fragments may attach to them, promoting formation of antibodies that can also target adjacent neural epitopes. Here, the slightly delayed onsets, prolonged postinfectious courses, and apparent responses to continued immunotherapy suggested dysimmune mechanisms.

This report strengthens evidence linking several idiopathic multisymptom conditions—including SFN and fibromyalgia—with dysimmunity, sometimes incident to infections or vaccinations. As with COVID-incident Guillain-Barré syndrome and all referral-based case series, the current cases neither confirm causality nor the clinical significance or magnitude of any association. However, identifying small-fiber neuropathy and multifocal motor neuropathy in 1 small sample of patients with WHO-defined long COVID provides rationale and preliminary data for larger investigations and may influence interim medical evaluations of similar patients.

Acknowledgment
The authors gratefully acknowledge patient contributions including details from medical personnel. They also thank many colleagues who referred patients, facilitated studies, or contributed

Figure 1 Case 15: Prolonged COVID-incident Multifocal Motor Neuropathy
data including Heather M. Downs, BS, Lisa Paul, NP, Shibani Mukerji, MD, PhD, Khosro Farhad, MD, Pedro Steven Buarque de Macedo, MD, Yancy Seamans, FNP, Joseph Tornabene, MD, Matthew P. Wicklund, MD, Jennifer Curtin, MD, Yair Mina, MD, Sara Dehbashi, MD, and Madeleine C. Klein, BS.

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**Disclosure**

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### Appendix  Authors

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<tr>
<th>Name</th>
<th>Location</th>
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<tr>
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<td>Drafting/revision of the manuscript for content, including medical writing for content, and major role in the acquisition of data</td>
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### References

Peripheral Neuropathy Evaluations of Patients With Prolonged Long COVID
Anne Louise Oaklander, Alexander J. Mills, Mary Kelley, et al.

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