## CORRESPONDENCE

## Guillain-Barré Syndrome Associated with SARS-CoV-2

TO THE EDITOR: From February 28 through March 21, 2020, in three hospitals in northern Italy, we examined five patients who had Guillain-Barré syndrome after the onset of coronavirus disease 2019 (Covid-19), the disease caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2). During that period, an estimated 1000 to 1200 patients with Covid-19 were admitted to these hospitals. Four of the patients in this series had a positive nasopharyngeal swab for SARS-CoV-2 at the onset of the neurologic syndrome, and one had a negative nasopharyngeal swab and negative bronchoalveolar lavage but subsequently had a positive serologic test for the virus. Detailed case reports are provided in the Supplementary Appendix, available with the full text of this letter at NEJM.org.

The first symptoms of Guillain–Barré syndrome were lower-limb weakness and paresthesia in four patients and facial diplegia followed by ataxia and paresthesia in one patient (Table 1). Generalized, flaccid tetraparesis or tetraplegia evolved over a period of 36 hours to 4 days in four patients; three received mechanical ventilation. The interval between the onset of symptoms of Covid-19 and the first symptoms of Guillain–Barré syndrome ranged from 5 to 10 days (Table 1 and Fig. S1 in the Supplementary Appendix). None of the patients had dysautonomic features.

On analysis of the cerebrospinal fluid (CSF), two patients had a normal protein level and all the patients had a white-cell count of less than 5 per cubic millimeter. Antiganglioside antibodies were absent in the three patients who were tested. In all the patients, a real-time polymerase-chain-reaction assay of the CSF was negative for SARS-CoV-2. Results of electrophysiological studies are shown in Table S1. Compound muscle action potential amplitudes were low but could be obtained; two patients had prolonged motor distal latencies. On electromyography, fibrillation potentials were pres-

ent in three patients initially; in another patient, they were absent initially but were present at 12 days. The findings were generally consistent with an axonal variant of Guillain–Barré syndrome in three patients and with a demyelinating process in two patients. Magnetic resonance imaging, performed with the administration of gadolinium, showed enhancement of the caudal nerve roots in two patients, enhancement of the facial nerve in one patient, and no signal changes in nerves in two patients. Additional laboratory findings are shown in Table S2.

All the patients were treated with intravenous immune globulin (IVIG); two received a second course of IVIG and one started plasma exchange. At 4 weeks after treatment, two patients remained in the intensive care unit and were receiving mechanical ventilation, two were undergoing physical therapy because of flaccid paraplegia and had minimal upper-limb movement, and one had been discharged and was able to walk independently.

The interval of 5 to 10 days between the onset of viral illness and the first symptoms of Guillain–Barré syndrome is similar to the interval seen with Guillain–Barré syndrome that occurs during or after other infections.<sup>2</sup> Although many infectious agents have been associated with Guillain–Barré syndrome, there may be a propensity for preceding infection with *Campylobacter jejuni*, Epstein–Barr virus, cytomegalovirus, and Zika virus. There have been reports of an association between Guillain–Barré syndrome and coronavirus infections.<sup>3,4</sup>

On the basis of this observational series involving five patients, it is not possible to determine whether severe deficits and axonal involvement are typical features of Covid-19-associated Guillain-Barré syndrome. We could not determine the effect of reduced vital capacity due to neuromuscular failure from Guillain-Barré syndrome in these patients, but such an effect might be considered if findings on chest imag-

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Table 1. Characteristics of Five Patients with Guillain–Barré Syndrome after the Onset of Covid-19.*	Treatment and Outcomes at Week 4	Received 2 cycles of IVIG; had poor outcomes, including persistence of severe upper-limb weak- ness, dysphagia, and lower-limb paraplegia	Received IVIG; had improvements, including decrease in ataxia and mild decrease in facial weakness	Received 2 cycles of IVIG; had poor outcomes, including ICU admission owing to neuromuscular respiratory failure and flaccid tetraplegia	Received IVIG; had mild improvement but unable to stand 1 mo after onset	Received IVIG and plasma exchange; had bacterial pneumonia during IVIG treatment, which delayed plasma exchange
	MRI Results	Head: normal Spine: enhancement of caudal nerve roots	Head: enhancement of facial nerve bilaterally Spine: normal	Head: normal Spine: enhancement of caudal nerve roots	Head: normal Spine: normal	Head: not performed Spine: normal
	Antiganglioside Antibodies:;	Negative	Not tested	Negative	Not tested	Negative
	CSF Findings†	Day 2 (first lumbar puncture): normal protein level; no cells; negative PCR assay for SARS- CoV-2 Day 10 (second lumbar puncture): protein level, 101 mg/dl; white-cell count, 4 per mm³; negative PCR assay for SARS- CoV-2	Day 3: protein level, 123 mg/dl; no cells; negative PCR assay for SARS-CoV-2	Day 3: protein level, 193 mg/dl; no cells; negative PCR assay for SARS-CoV-2	Day 5: normal protein level; no cells; negative PCR assay for SARS-CoV-2	Day 3: protein level, 40 mg/dl; white-cell count, 3 per mm³; CSF:serum albumin ratio, 1.2%; negative PCR assay for SARS-CoV-2
	Neurologic Signs and Symptoms	Flaccid areflexic tetraplegia evolving to facial weakness, upper-limb paresthesia (36 hr), and respiratory failure (day 6)	Facial diplegia and generalized areflexia evolving to lower- limb paresthesia with ataxia (day 2)	Flaccid tetrapares is and facial weakness evolving to areflexia (day 2) and respiratory failure (day 5)	Flaccid areflexic tetraparesis and ataxia (day 4)	Facial weakness, flaccid areflexic paraplegia (days 2–3), and respiratory failure (day 4)
	Onset of Neurologic Syndrome	7 Days after fever, cough, and ageusia	10 Days after fever and pharyngitis	10 Days after fever and cough	5 Days after cough and hyposmia	7 Days after cough, ageusia, and anos- mia
Table 1.	Patient No.	1	2	м	4	25

\* Covid-19 denotes coronavirus disease 2019, CSF cerebrospinal fluid, ICU intensive care unit, IVIG intravenous immune globulin, MRI magnetic resonance imaging, PCR polymerase chain reaction, and SARS-CoV-2 severe acute respiratory syndrome coronavirus 2.

† On CSF analysis, all the patients had a normal glucose level and IgG index and a polyclonal pattern on electrophoresis. The normal range for the protein level is 15 to 45 mg per

deciliter. An enzyme-linked immunosorbent assay was used to test for antibodies to GM1, GQ1b, and GD1b.

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ing are not commensurate with the severity of respiratory insufficiency. Guillain–Barré syndrome with Covid-19 should be distinguished from critical illness neuropathy and myopathy, which tend to appear later in the course of critical illness than Guillain–Barré syndrome.

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Disclosure forms provided by the authors are available with the full text of this letter at NEJM.org.

This letter is dedicated to the loving memory of Dr. Arrigo Moglia.

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