



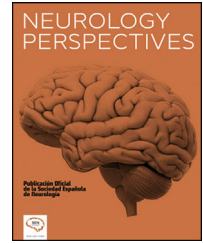
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SCIENTIFIC LETTER

Guillain-Barre syndrome after vaccination for Covid-19. The first report in Latin America

Síndrome de Guillain-Barre posterior a vacunación para Covid-19. Primer reporte en Latinoamérica

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With the onset of the COVID-19 (SARS-CoV2) pandemic in Wuhan-China at the end of 2019,¹ multiple clinical trials have been conducted seeking medical solutions to mitigate the impact of the virus. Vaccination has been the most important tool to combat infectious agents and their complications.² In the case of COVID-19, vaccines from different manufacturers have been developed with an action-focused on preventing serious cases and mortality. The Sinovac-life-science inactivated virus vaccine is one of them, with effectiveness studies in different age groups and use in different countries of the world within their vaccination plans.³

Vaccines can have rare, varied adverse effects including neurological entities. Guillain-Barre syndrome (GBS) is associated with multiple infectious agents including COVID-19,⁴ it has also been reported after vaccination of polio, hepatitis B, rabies, and influenza in periods from 2 days to 3 weeks⁵; recently one case of GBS has been reported with the Pfizer vaccine for COVID-19⁶ and two cases in the Johnson & Johnson vaccine studies.⁷ Although this does not generate a causal relationship, it creates the need to follow

up all patients vaccinated for COVID-19 who develop neurological pathologies, to increase the evidence and favor a precise medical approach with timely treatments in complex neurological conditions.

We report the case of a 73-year-old man, from Cali-Colombia, who consults a national reference university hospital in March 2021, with four-day symptoms consisting of upper limb (UL) paresis, posterior lower limb involvement (LL) associated with an inability to walk, and dyspnea. Four days before the onset of symptoms, the patient had received the first dose of vaccination for COVID-19-Sinovac. Upon admission to the emergency department, he presented flaccid symmetric quadriparesis with muscle strength in UL of 1/5 and bilateral LL 2/5, generalized hyporeflexia, without sensory alteration or sphincter involvement, without upper motor neuron signs; they suspected motor poly-neuro-radicalopathy with a descending pattern and a high risk of ventilatory failure, for which he was admitted to the intensive care unit with oxygen support, 24 h later with deterioration of the respiratory pattern and a requirement for orotracheal intubation. The patient had a history of GBS 20 years ago, without information on medical history or management, does not report other relevant personal or family history, nor febrile, gastrointestinal, or respiratory episodes in the last month.

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Table 1 Patient studies.

Serum test							
Test	Results	Test	Results	Test	Results	Test	Results
RT-PCR SARS-CoV-2	Negative	C3	88 mg/dL (Normal: 86–250 mg/dL)	C4	23.5 mg/dL	ENAS	(Normal: 12–72 mg/dL) 0.2 (Negative < 1)
Hemogram	Leukocytes 11,700/mm ³ , Hemoglobin: 13.5 g/dL Platelets: 204000/mm ³	Urine	Negative	porphobilinogen		Vitamin B12	354 pg/ml
Procalcitonin	0.07 ng/dL	Thyroid-Stimulating	3.2 mUI/L	Hormone		HCV antibody test	Negative
Urine and blood culture	Negative	and HBSAg		Carcinoembryonic	1.28	Antigen	(Normal: 0–3 ng/ml)
C-Reactive Protein	0.8	Prostate-specific	0.99	antigen	(Normal: 0–4 ng/ml)	Ca 19–9	5.1 (Normal: 0–37 U/ml)
Ferritin	136						
Nontreponemal tests and HIV	Negative						
Arterial blood gases	Metabolic acidosis						
Creatinine /// BUN	0.73 mg/dL /// 20 mg/dL						
CEREBROSPINAL FLUID TEST							
Color: Clear. Opening pressure: 21 cm h ₂ o							
Glucose: 106 mg/DL - LDH: 65 U/L - RBCs count: 70/mm ³ - Lymphocytes: 0/mm ³ - Proteins: 43 mg/DL - WBC count: 0/mm ³							
Bacilloscopy: Negative -Treponemal Test: Negative - cryptococcal antigen: Negative - Indian ink: Negative - Gram: Negative - CMV: Negative							
IMAGING TEST							
<ul style="list-style-type: none"> • Simple and contrast cervical and dorsal spine MRI • Simple and Contrast Brain MRI 				<ul style="list-style-type: none"> • No evidence of myelopathy or root compressions • No evidence of intracranial injuries 			
NEUROCONDUCTION STUDIES							
Sensory nerve conduction	Site	Peak (ms)	Amp (μV)	Site 1	Site 2	Dist (cm)	Vel (m/s)
Median-left and right (2 nt digit)	Wrist	NR (N < 3,6)	NR (N > 10)	Wrist	2 nt digit	14	NR (N > 40)
Ulnar-left and right (Lat Mall)	Calf	NR (N < 4,2)	NR (N > 5)	Calf	Lateral Mall	14	NR (N > 38)
Motor nerve conduction	Site	Peak (ms)	Amp (mV)	Site 1	Site 2	Dist (cm)	Vel (m/s)
Median-left and right (Abd poll brev)	Wrist	NR (N < 4)	NR (N > 5)	Elbow	Wrist		NR (N > 50)
Peroneo-left and right (Ext dig brev)	Ankle	NR (N < 6.1)	NR (N > 2.5)				
Tibial-left and right (Abd hall brev)	Ankle	NR (N < 6.1)	NR (N > 3)				
Ulnar-left and right (abd 5th digit)	Wrist	NR (N < 3,8)	NR (N > 3)	Elbow	Wrist		NR (N > 50)
MRI: magnetic resonance imaging, HIV: human immunodeficiency virus, BUN: blood urea nitrogen, ENAS: extractable nuclear antigen antibodies, HCV: hepatitis c virus, HBSAg: hepatitis b Surface antigen, RBC: red blood cell; WBC: White blood cell, LDH: lactate dehydrogenase, CMV: cytomegalovirus, 2 nt:second, Lat: lateral, Mall: Malleolus, 5th: fifth, abd poll brev: abductor pollicis brevis, ext dig brev: extensor digitalis brevis, abd hall brev: abductor hallux brevis, abd: abductor, amp: amplitude, dist: distance, vel: velocity, N:normal, NR: no response.							

Lumbar puncture was performed with opening pressure of 21 cm/H₂O, cytochemical without alterations, chest X-ray without signs of pneumonia or pleural effusions. The studies of the patient show a normal biochemical, metabolic, and infectious profile (Table 1), simple and contrasted magnetic resonance imaging (MRI) of the brain and total spine without findings of clinical relevance, and an electrophysiological study of four extremities that reported acute demyelinating motor and sensory polyneuropathy. (Table 1).

Neurology evaluates reporting flaccid quadriplegia, hyporeflexia, and acute ventilatory failure due to GBS, for

which plasmapheresis with 3% albumin was started (five sessions), the first being performed 72 h after admission. After the second plasmapheresis, the patient partially regains movement in the LL and UL with limited finger mobility, without cranial nerve involvement and communication through gestures. At the end of the plasmapheresis sessions, it is assessed by psychiatry, finding bilateral muscle strength in UL: elbow flexion and extension: 2/5, wrist flexion-extension: 2/5, without complete mobility of the fingers of the hands; LL: adductors 4/5, knee flexion-extension: 3/5, foot dorsiflexion: 1/5, foot plantarflexion:

Table 2 Rehabilitation in Guillain-Barre syndrome (GBS).

GBS-rehabilitation goals.

- Physical therapy: Maintenance of joint mobility, stretching, muscle strengthening, motor control, positioning, edema control.
- Occupational therapy: retraining in self-care, grippers and grips, transfers, energy saving.
- Phonoaudiology: vocal and swallowing muscles, ventilatory muscles, augmentative communication measures.
- Orthotic handling.
- Pain control.
- Mobility and walking assistance devices.
- Adaptations: home and family environment.
- Personal goals and social integration.

2/5, toe flexion-extension: 1/5; right patellar reflex + / +++ +, which was previously absent. A rehabilitation plan was started with objectives (Table 2), in addition to orthotic management for proper positioning, prevention of contractures, and pain management.

The patient presented in this case meets clinical and electrophysiological criteria for GBS⁸ with a favorable response to medical treatment, in addition to initiating symptoms after COVID-19 vaccination in a period similar to that reported with other infectious agents.⁵ Vaccination in the COVID-19 pandemic is an appropriate and widely applied intervention, which generates the need to study possible complications to offer the best medical treatment in the shortest time. Although, observational studies are required to define a solid causal relationship, a basis is created for the reporting of future cases and their treatment. To our knowledge, this case presented in March 2021 is the first report in Latin America.

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Ethical consideration

The patient gave his consent for the use of clinical and paraclinical information, it was authorized by the ethics committee of the Hospital Universitario del Valle, Cali, Colombia.

Declaration of Competing Interest

We declare that we have no conflict of interest.

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References

1. Wang D, Hu B, Hu C, Zhu F, Liu X, Zhang J, et al. Clinical characteristics of 138 hospitalized patients with 2019 novel Coronavirus-infected pneumonia in Wuhan, China. *JAMA*. 2020;323:1061–9. <https://doi.org/10.1001/jama.2020.1585>.
2. Canoui E, Launay O. History and principles of vaccination. *Rev Mal Respir*. 2019;36:74–81. <https://doi.org/10.1016/j.rmr.2018.02.015>.
3. Palacios R, Patiño EG, de Oliveira Piorelli R, Conde MTRP, Batista AP, Zeng G, et al. Double-blind, randomized, placebo-controlled phase III clinical trial to evaluate the efficacy and safety of treating healthcare professionals with the adsorbed COVID-19 (inactivated) vaccine manufactured by sinovac - PROFISCOV: A structured summary of a study protocol for a randomised controlled trial. *Trials*. 2020;21:853. <https://doi.org/10.1186/s13063-020-04775-4>.
4. Vallejo Serna R, Cantor González JF, Arce Gálvez L. Síndrome de Guillain-Barré asociado a COVID-19: diagnóstico, tratamiento y rehabilitación. *Neurol Perspect*. 2021;1:104–6. <https://doi.org/10.1016/j.neurop.2021.03.003>.
5. Chen Y, Zhang J, Chu X, Xu Y, Ma F. Vaccines and the risk of Guillain-Barré syndrome. *Eur J Epidemiol*. 2020;35:363–70. <https://doi.org/10.1007/s10654-019-00596-1>.
6. Waheed S, Bayas A, Hindi F, Rizvi Z, Espinosa PS. Neurological complications of COVID-19: Guillain-Barre syndrome following Pfizer COVID-19 vaccine. *Cureus*. 2021;13:2–5. <https://doi.org/10.7759/cureus.13426>.
7. Márquez Loza AM, Holroyd KB, Johnson SA, Pilgrim DM, Amato AA. Guillain-Barré Syndrome in the placebo and active arms of a COVID-19 vaccine clinical trial: temporal associations do not imply causality. *Neurology*. 2021 <https://doi.org/10.1212/WNL.0000000000011881>.
8. Leonhard SE, Mandarakas MR, Gondim FAA, Bateman K, Ferreira MLB, Cornblath DR, et al. Diagnosis and management of Guillain-Barré syndrome in ten steps. *Nat Rev Neurol*. 2019;15:671–83. <https://doi.org/10.1038/s41582-019-0250-9>.