LETTER



Guillain–Barre/Miller–Fisher overlap syndrome or acute, motor and sensory, axonal neuropathy with cranial nerve involvement?

Josef Finsterer 厄

Department Neurology, Neurology & Neurophysiology Center, Vienna, Austria

KEYWORDS Guillain–Barre syndrome; cranial nerves; side effect; SARS-CoV-2 vaccination; Miller–Fisher syndrome

Letter to the Editor

With interest, we read the article by Liang et al. about a 64 y-old male with Guillain-Barre syndrome (GBS) Miller-Fisher syndrome (MFS) overlap syndrome starting 11 d after the second jab of the Sinovac-CoronaVac vaccine.¹ Intravenous immunoglobulins (IVIG) and acupuncture resulted in complete recovery at the last followup, 54 d after dismissal. The study is attractive but raises concerns that require further discussion.

We disagree with the diagnosis GBS-MFS overlap syndrome.¹ According to the nerve conduction studies (NCSs) the patient had axonal, sensorimotor neuropathy.¹ Because there was obvious dissociation cyto-albuminique,¹ the patient should be classified as acute, motor and sensory, axonal neuropathy (AMSAN) with cranial nerve involvement rather than GBS-MFS overlap. Cranial nerves affected in the index patient were the 6th and 7th cranial nerve.¹ AMSAN is more prevalent in the Eastern compared to the Western hemisphere, where acute, inflammatory demyelinating polyneuropathy (AIDP) prevails. Arguments against MFS are that the patient had facial palsy and that ophthalmoparesis was incomplete. Only bulb abduction was impaired, whereas in MFS there is usually ophthalmoparesis for all degrees of eye movements.

Missing are the cytokine and chemokine levels and the titers of the glial acidic fibrillary protein (GAFP) in the CSF. These parameters have been found elevated in GBS or central nervous system disease following a SARS-CoV-2 vaccination.²

We disagree with the notion that the index is the first presenting with MFS-GBS overlap syndrome.¹ Though rare, GBS-MFS has been previously reported in single patients.³

Missing is the information whether the autonomic innervation of the heart was impaired or not. AMSAN commonly is associated with autonomic dysfunction, which is why we should know if blood pressure regulation was impaired or not and whether supra-ventricular or ventricular arrhythmias were detected on long-term electrocardiogram (ECG) recordings or telemetry. Particularly, we should know if the Valsalva maneuver was positive or negative, if there was postural tachycardia syndrome (POTS), increased or decreased heart rate variability on frequency analysis of long-term (ECG) recordings, and an abnormal tilt test. We should also know if there were clinical or echocardiographic signs of heart failure and if troponin or pro-brain natriuretic peptide (pro-BNP) were elevated or not.

The term "right facial diplegia" is misleading.¹ Do the authors mean facial diplegia with right-sided predominance or do they truly mean right-sided facial palsy without affection of the contralateral side?

Overall, the study carries obvious limitations that require reevaluation and discussion. Clarifying these weaknesses would strengthen the conclusions and could improve the study. MFS is questionable if the facial nerves are affected in addition to cranial nerves innervating extra-ocular eye muscles.

Author contribution

SM: design, literature search, discussion, first draft, critical comments, final approval, JF: literature search, discussion, critical comments, final approval.

Disclosure statement

No potential conflict of interest was reported by the author(s).

Funding

The author(s) reported that there is no funding associated with the work featured in this article.

Data availability

All data are available from the corresponding author.

ORCID

Josef Finsterer (D) http://orcid.org/0000-0003-2839-7305

CONTACT Josef Finsterer 🖾 fifigs1@yahoo.de 🖬 Neurology & Neurophysiology Center, Postfach 20, Vienna 1180, Austria.

© 2022 The Author(s). Published with license by Taylor & Francis Group, LLC.

This is an Open Access article distributed under the terms of the Creative Commons Attribution-NonCommercial-NoDerivatives License (http://creativecommons.org/licenses/by-nc-nd/4.0/), which permits non-commercial re-use, distribution, and reproduction in any medium, provided the original work is properly cited, and is not altered, transformed, or built upon in any way.

References

- Liang H, Cao Y, Zhong W, Ma Z, Liu J, Chen H. Miller-Fisher syndrome and Guillain-Barre syndrome overlap syndrome following inactivated COVID-19 vaccine: case report and scope review. Hum Vaccin Immunother. 2022 Oct 31;2125753. doi:10.1080/ 21645515.2022.2125753.
- Gigli GL, Vogrig A, Nilo A, Fabris M, Biasotto A, Curcio F, Miotti V, Tascini C, Valente M. HLA and immunological features of SARS-CoV -2-induced Guillain-Barré syndrome. Neurol Sci. 2020 Dec;41 (12):3391–94. doi:10.1007/s10072-020-04787-7.
- 3. Dang YL, Bryson A. Miller-Fisher syndrome and Guillain-Barre syndrome overlap syndrome in a patient post Oxford-AstraZeneca SARS-CoV-2 vaccination. BMJ Case Rep. 2021 Nov 30;14(11): e246701. doi:10.1136/bcr-2021-246701.