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Case Report
Leptomeningeal and Intraparenchymal Blood Barrier Disruption in a MOG-IgG-Positive Patient

Seyed Hamidreza Mohseni, Hanne Pernille Bro Skejoe, Jens Wuerfel, Friedemann Paul, Markus Reindl, Sven Jarius, and Nasrin Asgari

1Department of Radiology, Slagelse Hospital, Slagelse, Denmark
2Department of Radiology, Aleris-Hamlet Hospital, Copenhagen, Denmark
3Medical Image Analysis Center Basel and Department of Biomedical Engineering, University Basel, Switzerland
4Clinical and Experimental Multiple Sclerosis Research Center and NeuroCure Clinical Research Center, Department of Neurology, Charité-Universitätsmedizin Berlin, Germany
5Experimental and Clinical Research Center, Max Delbrueck Center for Molecular Medicine and Charité-Universitätsmedizin Berlin, Berlin, Germany
6Clinical Department of Neurology, Medical University Innsbruck, Innsbruck, Austria
7Molecular Neuroimmunology Group, Department of Neurology, University Hospital Heidelberg, Heidelberg, Germany
8Department of Neurology, Slagelse Hospital, Institute of Regional Health Research, Denmark
9Department of Neurobiology, Institute of Molecular Medicine, University of Southern Denmark, Denmark

Correspondence should be addressed to Nasrin Asgari; nasgari@health.sdu.dk

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1. Introduction
Pathogenic immunoglobulin G (IgG) autoantibodies directed to myelin oligodendrocyte glycoprotein (MOG), an oligodendrocytic protein localized to the outer surface of the myelin sheaths, have recently been identified in patients with inflammatory CNS demyelination [1–6]. This discovery has led to the recognition of a new disease entity, now often referred to as “MOG encephalomyelitis” (MOG-EM) [2]. Phenotypically, MOG-EM may present as optic neuritis (ON), (often longitudinally extensive) transverse myelitis, brainstem encephalitis, encephalitis, or any combination of these syndromes and, especially in children, as acute demyelinating encephalomyelitis (ADEM) [3–6]. Myelitis is often associated with leptomeningeal (pia-arachnoid) contrast enhancement (LME). LME is a sign of leptomeningeal barrier impairment and leakage of contrast agent into the subarachnoid space. Understanding LME patterns may help in the differential diagnosis of the various forms of myelitis [7–10]. So far, little is known about LME in MOG-IgG-positive myelitis.
Leptomeningeal enhancement Intraparenchymal enhancement Longitudinally extensive transverse myelitis

Figure 1: Spinal cord MRI. Sagittal gadolinium-enhanced T1-weighted images with fat saturation showing linear leptomeningeal thickening and enhancement (a) and intramedullary parenchymal patchy enhancement (b), correlating with diffuse T2-weighted hyperintense signal increase extending from C2 to Th3 (c).

2. Case Report

A 30-year-old woman with no previous history of systemic inflammatory disease or neoplastic diseases developed loss of vision in the left eye and two days later in the right eye due to acute ON, followed by tetraparesis two weeks later. Spinal cord MRI obtained prior to treatment revealed LME, intraparenchymal contrast enhancement corresponding to the site of LME, and longitudinal extensive transverse myelitis (LETM) extending from C2 to Th3 (Figure 1).

Cerebral MRI was normal. Cerebrospinal fluid contained 122 leukocytes/mm³ with polymorphonuclear predominance; oligoclonal bands were not determined. Aquaporin-4 (AQP4)-IgG was negative. Accordingly, seronegative neuromyelitis optica was suspected by that time. Follow-up MRI demonstrated resolution of LME four months later.

Retrospective testing by means of two cell-based assays employing fixed and live HEK293 cells, respectively, transfected with full-length human MOG revealed the presence of MOG-IgG antibodies in a serum sample taken at onset [1, 2]. MOG-IgG seropositivity was confirmed in a second sample taken four years later.

3. Discussion

Inflammation in demyelinating diseases of the CNS is commonly associated with blood-brain barrier (BBB) disruption. Leptomeningeal involvement has recently been recognized as an important feature in multiple sclerosis pathogenesis [8]. LME has also been observed in AQP4-IgG-positive neuromyelitis optica spectrum disorder [9, 10]. Here, we present a case of MOG encephalomyelitis (MOG-EM) [2] with a longitudinally extensive demyelinating spinal cord lesion in which the blood-CNS barriers were disrupted, as demonstrated by gadolinium-enhanced MRI. Our findings demonstrate that spinal cord LME may occur also in MOG-EM, one of the most important differential diagnoses of MS. Notably, LME, which indicates an abnormally permeable leptomeningeal-blood barrier, was accompanying intraparenchymal BBB breakdown during an attack of acute myelitis, as visualised by contrast enhancement on T1-weighted imaging. This finding suggests that meningeal inflammation may have occurred as a bystander reaction following MOG-IgG-related parenchymal inflammation associated with subpial demyelination. Lesions involving the peripheral portions of the spinal cord indeed occur in a substantial number of cases of MOG-IgG-positive myelitis, as has recently been shown [3]. Similarly, cortical brain lesions have been described in MOG-EM, some of which were associated with LME [3, 11], and patients with MOG-IgG-positive ON may commonly present with perioptic contrast enhancement [3]. Future studies should systematically assess the presence of LME in MOG-IgG-related myelitis as well as its potential value in discriminating MOG-EM and other demyelinating diseases affecting the spinal cord.
Glossary

AQP4: Aquaporin-4  
BBB: Blood-brain barrier  
CNS: Central nervous system  
CSF: Cerebrospinal fluid  
IgG: Immunoglobulin G  
LETM: Longitudinal extensive transverse myelitis  
LME: Leptomeningeal enhancement  
MOG: Myelin oligodendrocyte glycoprotein  
MRI: Magnetic resonance imaging  
ON: Optic neuritis.

Conflicts of Interest

Seyed Hamidreza Mohseni, Nasrin Asgari, Hanne Pernille Bro Skejoe, Friedemann Paul, and Sven Jarius report no conflicts of interest. The University Hospital and Medical University of Innsbruck (Austria, employer of Markus Reindl) receive payments for antibody assays (MOG, AQP4, and other autoantibodies) and for MOG and AQP4 antibody validation experiments organized by Euroimmun (Lübeck, Germany).

Authors’ Contributions

Seyed Hamidreza Mohseni contributed to acquisition of data, interpretation of results, and drafting and revising the manuscript. Hanne Pernille Bro Skejoe contributed to MRI reevaluation, interpretation of results, and revising the manuscript. Jens Wuerfel contributed to MRI reevaluation, interpretation of results, and revising the manuscript. Friedemann Paul contributed to interpretation of results and revising the manuscript. Markus Reindl contributed to determination of MOG-IgG and revising the manuscript. Sven Jarius contributed to determination of MOG-IgG and revising the manuscript. Nasrin Asgari contributed to study concept and design, acquisition of data, interpretation of results, and revising the manuscript. All authors read and approved the final version of the manuscript.

References
