Pediatric-onset systemic lupus erythematosus (SLE) encompasses diverse symptoms, such as headache, seizures, stroke, depression, psychosis, cognitive impairment, chorea, and neuropathy. Neuropsychiatric lupus occurs in one-fourth of cases. Although 95%–97% of the children survive, the disease flares in childhood in 20%, and 25% sustain permanent neuropsychiatric injury. The paucity of biomarkers, mostly studied in blood; lack of modern neuroimmunologic studies; and poorly defined treatment modalities have hindered management.

We describe a child with extensive peripheral and CNS manifestations and multiorgan involvement. Multiple cellular and cytokine/chemokine markers indicated profound neuroinflammation with some components responsive, others resistant, to 3-agent immunotherapy.

Methods. Description of the laboratory methods is provided at Neurology.org/nn.

Case report. A previously healthy 11-year-old African American girl was hospitalized for intractable headaches, myalgias, malar rash, intermittent paresthesias, foot drop, and 25-pound weight loss over 5 months. SLE was diagnosed based on speckled antinuclear antibodies >1:640, DNA-DS 257 IU/mL, anti-histone 321 U/mL, and anti-Smith 386 U/mL. CSF studies revealed high total immunoglobulin G (IgG) and IgG synthesis rate (table e-1). Head MRI disclosed transient fluid-attenuated inversion recovery T2 hyperintensities and mild cerebral atrophy (figure e-1). EMG/nerve conduction studies (NCS) exhibited sensorimotor neuropathy, conduction block at the fibular head, and absent F-waves and H-reflex, suggesting acute demyelinating peripheral neuropathy. Classical features of acute inflammatory demyelinating polyneuropathy, such as acute ascending radiculopathy, cranial nerve involvement, respiratory compromise, and albuminocytologic dissociation, were absent. Other abnormalities included membranous lupus nephritis (International Society of Nephrology/Renal Pathology Society class V), a small pericardial effusion, leukopenia, anemia, and thrombocytopenia.

Results. Neuroimmunologic studies. In CSF (figure 1), pretreatment inflammatory cytokine/chemokine concentrations were strikingly elevated: CXCL13 (331-fold), CCL19 (15-fold), BAFF (5.7-fold), CXCL12 (5.2-fold), CXCL10 (>143-fold), interleukin-6 (IL-6) (7.4-fold). The CSF/serum ratio was raised at 0.17 for CXCL13 (controls 0.01) and >1.2 for CXCL10 (controls 0.65), indicating intrathecal chemokine secretion. In serum, CXCL13 (23-fold) and CXCL10 (88-fold) concentrations were marked, less so for CCL21 (1.5-fold), whereas CCL17 was low (≈88%).

After 6 weeks of immunotherapy, the 13-fold elevated CSF B-cell frequency and high concentration of CXCL13, CXCL12, and CCL19 plunged to the control median, but BAFF and CXCL10 remained elevated. Increased T-cytotoxic/suppressor cell frequency and reduced T-helper/inducer frequency lowered the T-helper (Th)/T-suppressor ratio (table e-1). The initially elevated natural killer cell percentage decreased. CSF IL-6 plummeted 93% from 13.3 to 0.95 pg/mL (controls 0–7.5). Total IgG declined, the IgG synthesis rate remained elevated, and CSF leukocytosis abated.

Discussion. In this case of debilitating pediatric neurolupus, the main findings were (1) major upregulation of CSF B cells and inflammatory cytokines/chemokines despite minimal pleocytosis and no oligoclonal bands; (2) evidence of increased Th1 and decreased Th2 immune responses, (3) incomplete prediction of CSF perturbations by blood testing; (4) downregulation of many, not all, clinical and...
immunologic abnormalities on multimodal disease-modifying therapy; and (5) prolonged clinical and central/peripheral electrophysiologic recovery.

In CSF, the overexpression of lymphocyte chemoattractants (CXCL13, CCL19, CXCL12, CXCL10) and homeostatic cytokines (BAFF) may contribute to central B and T cell recruitment and proliferation/survival in neurolupus. Lymphoid chemokines (CCL19, CXCL13) are relevant because they may form ectopic lymphoid structures within the CNS in chronic neuroinflammation.

Posttreatment reduction in CSF B-cell frequency, CXCL13, CCL19, CXCL12, and IL-6 concentrations led to clinical improvement, whereas high BAFF and CXCL10 persisted despite it. CXCL10 is a potent inflammatory mediator in Th1 immune responses, induces migration of certain T-cell subsets, and is involved in lupus nephritis. The extreme CSF CXCL10 elevation is similar to that in adult-onset SLE.

In serum, SLE-induced reduction in CCL17, which attracts CXCR4-bearing Th2 cells, is consistent with suppressed expression of CCL17 in lupus-prone MRL/lpr mice. Increased CCL21 would recruit distinct T- and B-cell subpopulations by binding to CXCR7, as does CCL19. Serum CXCL13 and CXCL10 were predictably elevated.

Multiagent immunotherapy prevents isolating the effect of any one agent. However, from our research in a paraneoplastic disorder, high-dose glucocorticoids probably account for the corrective effects on chemokines/cytokines. Immune cellular phenotypic changes are most likely attributable to cyclophosphamide. IVIG has neither effect, but modulates antibodies and unmeasured parameters.

These findings, requiring confirmation in other children with neurolupus, have clinical implications. CSF lymphocyte subset analysis, available at hospital flow cytometry laboratories, characterized neuroinflammation and raised the level of alert more clearly than routine CSF studies. No one finding was specific for neurolupus, but the combined immunology panel showed commonalities and differences compared to multiple sclerosis and paraneoplastic disorders. It can be incorporated into translational research on pediatric neurolupus to identify biomarkers of disease activity, treatment response, or prognosis. CSF posttreatment testing for no evidence of disease activity addresses questions about the type and duration of immunotherapy. Because swift and sufficient handling of neuroinflammation is crucial, more rapid, brain-directed targeting of the B-cell/humoral component should be considered in the initial therapeutic approach.

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Figure 1 CSF inflammatory markers before and after treatment

(A) CD19 + CD3^- B cell frequency. (B) CXCL13 concentration. (C) CCL19 concentration. (D) BAFF concentration. (E) CXCL12 concentration. (F) CXCL10 concentration. CSF is normally analyzed undiluted, but even at a 1:20 dilution, the patient's massive CXCL10 concentration exceeded the highest concentration of standard. The control median and 95% confidence intervals from 18 to 25 children with noninflammatory neurologic disorders are shown in colored lines.
B.K.A. wrote the case report, and M.R.P. wrote the manuscript. Z.Y.W. performed and analyzed the electrophysiologic studies. All authors reviewed/revised and finalized the manuscript prior to submission.

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