A 21-year-old Pakistani woman with a history of 2 seizures at 20 months of age and consanguineous parents was diagnosed with premature ovarian failure and hyperprolactinemia. Her neurologic examination was normal. Brain imaging showed confluent white matter T2/fluid-attenuated inversion recovery hyperintensities with periventricular cavitations (figure). Genetic testing showed homozygosity for a p.Lys273Arg missense mutation in the \textit{EIF2B2} gene (NM_014239.3: c.818A>G), which had previously been reported as disease-causing,\textsuperscript{1} and confirmed vanishing white matter (VWM) ovarioleukodystrophy. At 2-year follow-up, she remains neurologically asymptomatic with stable imaging. There is phenotypic variation in VWM disease and neurologic deterioration is associated with stressors (head trauma, fever).\textsuperscript{1,2}

**AUTHOR CONTRIBUTIONS**

Dr. Mukerji conceptualized and designed the study, interpreted the data, and drafted the manuscript. Dr. Eichler interpreted the data and revised the manuscript for intellectual content.

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**DISCLOSURE**

The authors report no disclosures relevant to the manuscript. Go to Neurology.org for full disclosures.

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