A two-month-old boy presented with breastfeeding difficulty, hypoactivity, hyporeactivity, and seizures. Neuroimaging showed multiple areas of cytotoxic and vasogenic edema in the midbrain, cerebellum, basal ganglia, and brain hemispheres, with lactate peak (►Figure 1). Exome sequencing revealed a heterozygous mutation c.597dup (p.His200Serfs*25) in the SCL19A3 gene. The SCL19A3 gene encodes thiamine transporter-2. Mutations can result in thiamine metabolism dysfunction syndrome-2, and they can present as early as infantile Leigh-like syndrome,¹ a potentially treatable disease. This patient was submitted to thiamine and biotin replacement, but his evolution confirmed a poor prognosis (►Figure 2).² The poor outcome warrants genetic counseling for the families.

Conflict of Interest
The authors have no conflict of interest to declare.

References

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Figure 1  Brain magnetic resonance imaging (MRI) scan at two months of age. Axial diffusion (A,B); fluid-attenuated inversion recovery (FLAIR) (C); apparent diffusion coefficient (ADC) map (D,E); and T2-weighted imaging (F). Symmetric areas of cytotoxic (red arrows) and vasogenic (asterisk) edema in the midbrain, superior vermis, subcortical white matter of the cerebral hemispheres, genu of the corpus callosum, internal capsules, and basal ganglia, mainly in the putamina and thalami. There are also some cystic changes in the occipital lobes (blue arrows). Magnetic resonance spectroscopy (MRS) (G) with intermediate TE (144 ms) showing the lactate inverted peaks at 1.3 ppm chemical shift (green arrow).

Figure 2  Two-month follow-up after treatment with biotin and thiamine. Axial diffusion (A); FLAIR (B); T2- (C) and T1-weighted (D) multiplanar gradient-recalled (MPGR) imaging. Resolution of the areas with restriction diffusion and interval encephalomalacia and necrosis in the basal ganglia bilaterally, mainly in the thalami (arrows). There is diffuse cerebral atrophy and holohemispheric subdural hematomas (asterisk), due to stretching of the bridging veins.