## Opis choroby *

Definicja

A rare, systemic disease characterized by persistent vomiting with confusion, lethargy, disorientation, hyperreflexia, hyperventilation, and tachycardia, with rapid progression to seizures, non-inflammatory encephalopathy, coma and death. It typically develops between 12 hours and 3 weeks after recovery from a viral illness, such as upper respiratory tract infection or gastroenteritis. Hepatomegaly, acute hepatic steatosis, fatty liver degeneration and multiple laboratory abnormalities are associated.

Dane

**Klasyfikacja**
Choroba

**Kod ORPHA**
3096

**Kod OMIM**
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**Kod ICD10**
G93.7

**Kod ICD11**
8E46

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*Źródło

orphanet