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
-

Kod ICD10
G93.7

Kod ICD11
8E46

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*Źródło

orphanet