Zaburzenie neurorozwojowe związane z PLAA
Kod Orpha: 521426 Kod OMIM: 617527

Opis choroby *

Definicja
A rare genetic neurological disorder characterized by infantile onset of progressive leukoencephalopathy, microcephaly, severe global developmental delay, and spasticity resulting in quadriparesis and posture deformation. Additional features include an abnormally exaggerated startle reflex, seizures, dystonia, and hypomimia or amimia, as well as progressive chest deformities and contractures of large and hyperextensibility of small joints, among others. Thin corpus callosum is a prominent feature in brain imaging, in addition to white matter abnormalities consistent with leukoencephalopathy.
Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
PLAAND
PLAAND

Kod ORPHA
521426

Kod OMIM
617527

Kod ICD10
Q87.8

Kod ICD11
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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.