DISEASE:
PEHO syndrome

NAME:	PEHO syndrome
DESCRIPTION:	PEHO (Progressive encephalopathy with Edema, Hypsarrhythmia and Optic atrophy) syndrome is a rare neurodegenerative disorder belonging to the group of infantile progressive encephalopathies.
ORPHACODE:	2836
SYNONYMS:	Progressive encephalopathy with edema, hypsarrhythmia and optic atrophy Progressive encephalopathy-optic atrophy syndrome
XREF(S):	Orphanet ICD-10 MeSH OMIM
ANALYTE(S):	KIF1A ZNHIT3
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Epilepsy (gene panel)
- Neuropathy (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Antwerpen

Related Analytes

- kinesin family member 1A
- zinc finger HIT-type containing 3

Related Gene Panels

- Neuropathy (148 genes) - IPG
- Rare epilepsy with developmental delay (> 240 genes) - UZA