DISEASE:
Peroxisomal acyl-CoA oxidase deficiency

NAME:	Peroxisomal acyl-CoA oxidase deficiency
DESCRIPTION:	Peroxisomal acyl-CoA oxidase deficiency is a rare neurodegenerative disorder that belongs to the group of inherited peroxisomal disorders and is characterized by hypotonia and seizures in the neonatal period and neurological regression in early infancy.
ORPHACODE:	2971
SYNONYMS:	Pseudo-NALD Pseudo-neonatal adrenoleukodystrophy Pseudoadrenoleukodystrophy
XREF(S):	Orphanet OMIM MeSH ICD-10
ANALYTE(S):	ACOX1
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