DISEASE:
Anhidrotic ectodermal dysplasia-immunodeficiency-osteopetrosis-lymphedema syndrome

NAME:	Anhidrotic ectodermal dysplasia-immunodeficiency-osteopetrosis-lymphedema syndrome
DESCRIPTION:	This syndrome is characterized by severe immunodeficiency, osteopetrosis, lymphedema and anhidrotic ectodermal dysplasia.
ORPHACODE:	69088
SYNONYMS:	OL-EDA-ID
XREF(S):	Orphanet ICD-10 OMIM OMIM
ANALYTE(S):	IKBKG
CREATED:	13 May 2019 - 01:02
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RELATED CONTENT

Related Genetic Tests

- Primary lymphedema / fetal hydrops (gene panel)

Related Laboratories

- Centre de Génétique Médicale UCL

Related Analytes

- inhibitor of nuclear factor kappa B kinase regulatory subunit gamma