DISEASE:
Fryns syndrome

NAME:	Fryns syndrome
DESCRIPTION:	A rare multiple congenital anomaly syndrome characterized by congenital diaphragmatic hernia (CDH) and pulmonary hypoplasia, distal limb hypoplasia and facial anomalies in addition to variable expression of additional birth defects.
ORPHACODE:	2059
SYNONYMS:	Diaphragmatic hernia-abnormal face-distal limb anomalies syndrome
XREF(S):	Orphanet MeSH ICD-10 OMIM
ANALYTE(S):	PIGN
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