DISEASE:
8p11.2 deletion syndrome

NAME:	8p11.2 deletion syndrome
DESCRIPTION:	8p11.2 deletion syndrome is a contiguous gene syndrome characterized by the association of congenital spherocytosis, dysmorphic features, growth delay and hypogonadotropic hypogonadism.
ORPHACODE:	251066
SYNONYMS:	Del(8)(p11.2) Monosomy 8p11.2
XREF(S):	Orphanet ICD-10
ANALYTE(S):	ANK1
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

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