DISEASE:
Schinzel-Giedion syndrome

NAME:	Schinzel-Giedion syndrome
DESCRIPTION:	Schinzel-Giedion syndrome (SGS) is an ectodermal dysplasia syndrome chiefly characterized by a distinctive facial dysmorphism, hydronephrosis, severe developmental delay, typical skeletal malformations, and genital and cardiac anomalies.
ORPHACODE:	798
SYNONYMS:	SGS
XREF(S):	Orphanet OMIM MedDRA ICD-10
ANALYTE(S):	SETBP1
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

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