

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):	<u>Orphanet</u> <u>OMIM</u> <u>MedDRA</u> <u>ICD-10</u>
ANALYTE(S):	<u>SETBP1</u>
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