

Opis choroby *

Definicja

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.

Dane

Klasyfikacja	Synonimy
Zespół wad wrodzonych SGS	SGS

Kod ORPHA	Kod OMIM	Kod ICD10
798	269150	Q87.0

Kod ICD11
LD27.0Y

[*Źródło](#)

orphanet