

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

798

Kod OMIM

269150

Kod ICD10

Q87.0

Kod ICD11

LD27.0Y

*Źródło

orphanet