Opis choroby *

Definicja

Shprintzen□Goldberg omphalocele syndrome is a very rare inherited malformation syndrome characterized by omphalocele, scoliosis, mild dysmorphic features (downslanted palpebral fissures, s-shaped eyelids and thin upper lip), laryngeal and pharyngeal hypoplasia and learning disabilities.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
3164

Kod OMIM
182210

Kod ICD10
Q79.2

Kod ICD11
LD2F.1Y

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*Źródło

orphanet