

Zespół Sticklera

Kod Orpha: 828 Kod OMIM: 614284

Opis choroby *

Definicja

A rare group of genetic connective tissue disorders characterized by ophthalmic, auditory, orofacial and articular manifestations. The two main clinical forms are clinically distinguished by the vitreous phenotype; stickler type 1 by a vestigial vitreous gel in the immediate retrolental space, bordered by a distinct folded membrane, and Stickler type 2 by sparse and irregularly thickened bundles of fibers throughout the vitreous cavity.

Dane

Klasyfikacja

Choroba

Synonimy

Hereditary progressive arthroophthalmopathy

Kod ORPHA

828

Kod OMIM

614284

Kod ICD10

Q87.0

Kod ICD11

LD2F.1Y

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.