

## 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

<b>Klasyfikacja</b> Choroba	<b>Synonimy</b> Hereditary progressive arthroophthalmopathy
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<b>Kod ORPHA</b> 828	<b>Kod OMIM</b> 614284	<b>Kod ICD10</b> Q87.0
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<b>Kod ICD11</b> LD2F.1Y
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### \*Źródło

orphanet