

## Opis choroby \*

### Definicja

A rare, genetic, metabolic disease with connective tissue and eye involvement, characterized by progressive ectopic mineralization and fragmented elastic fibers in the skin, retina and vascular walls.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Gronblad-Strandberg-Touraine syndrome

PXE

Zespół Gronblada, Strandberga i Touraine'a

PXE

#### Kod ORPHA

758

#### Kod OMIM

264800

#### Kod ICD10

Q82.8

#### Kod ICD11

EC40

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#### \*Źródło

orphanet