

Opis choroby *

Definicja

A rare autosomal recessive connective tissue disorder characterized by tortuosity and elongation of the large and medium-sized arteries and a propensity towards aneurysm formation, vascular dissection, and stenosis of the pulmonary arteries.

Dane

Klasyfikacja	Synonimy
Zespół wad wrodzonych ATS	ATS

Kod ORPHA	Kod OMIM	Kod ICD10
3342	208050	I77.1

Kod ICD11
LD28.Y

*Źródło

orphanet