Homozygotyczna hipercholesterolemia rodzinna
Kod Orpha: 391665 Kod OMIM: 144010

Opis choroby *

Definicja
A rare disorder of lipid metabolism characterized by severely elevated low-density lipoprotein cholesterol levels and subsequent premature formation of atherosclerotic plaques in the coronary arteries, proximal aorta, and other arteries, significantly increasing the risk of cardiovascular disease at an early age. Xanthomas of the skin and in tendons are also a hallmark of the disease. Lethality is high due to early complications, in particular myocardial infarction.

Dane

Klasyfikacja
Choroba

Synonimy
HoFH
HoFH

Kod ORPHA
391665

Kod OMIM
144010

Kod ICD10
E78.0

Kod ICD11
-

*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.