Zespół FRAXF
Kod Orpha: 100974 Kod OMIM:

Opis choroby *

Definicja
FRAXF syndrome was originally identified in a family with developmental delay and an expanded CCG repeat at the folate-sensitive FRAXF fragile site. Since this initial description, FRAXF has been associated with a range of manifestations but no clear phenotype has been established.
Dane

Klasyfikacja
Choroba

Kod ORPHA
100974

Kod OMIM
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Kod ICD10
Q99.2

Kod ICD11
-

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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.