Opis choroby *

Definicja

A rare non-hereditary condition characterized by gastrointestinal stromal tumors (GIST, intramural mesenchymal tumors of the gastrointestinal tract with neuronal or neural crest cell origin), pulmonary chondromas and extraadrenal paragangliomas.

Dane

Klasyfikacja
Choroba

Kod ORPHA
139411

Kod OMIM
604287

Kod ICD10
D44.8

Kod ICD11
2F7A.0

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*Źródło

orphanet