DISEASE:
Friedreich ataxia

NAME:	Friedreich ataxia
DESCRIPTION:	Friedreich ataxia (FRDA) is an inherited neurodegenerative disorder classically characterized by progressive gait and limb ataxia, dysarthria, dysphagia, oculomotor dysfunction, loss of deep tendon reflexes, pyramidal tract signs, scoliosis, and in some, cardiomyopathy, diabetes mellitus, visual loss and defective hearing.
ORPHACODE:	95
SYNONYMS:	FA FRDA
XREF(S):	Orphanet MeSH MedDRA OMIM OMIM ICD-10
ANALYTE(S):	FXN
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- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Brussel VUB
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