DISEASE:
Familial dilated cardiomyopathy with conduction defect due to LMNA mutation

NAME:	Familial dilated cardiomyopathy with conduction defect due to LMNA mutation
DESCRIPTION:	A rare familial cardiomyopathy characterized by left ventricular enlargement and/or reduced systolic function preceded or accompanied by significant conduction system disease and/or arrhythmias including bradyarrhythmias, supraventricular or ventricular arrhythmias. Disease onset is usually in early to mid-adulthood. Sudden cardiac death may occur and may be the presenting symptom. In some cases, it is associated with skeletal myopathy.
ORPHACODE:	300751
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	LMNA
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RELATED CONTENT

Related Genetic Tests

- Cardiomyopathy, hereditary (gene panel)
- Hypertrophic cardiomyopathy (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- lamin A/C

Related Gene Panels

- Cardiomyopathy, hereditary (208 genes) - VUB
- Hypertrophic cardiomyopathy (75 genes) - IPG