DISEASE:
Familial isolated restrictive cardiomyopathy

NAME:	Familial isolated restrictive cardiomyopathy
DESCRIPTION:	A rare genetic cardiac disease characterized by restrictive ventricular filling due to high ventricular stiffness that results in severe diastolic dysfunction in the absence of dilated or hypertrophied ventricles.
ORPHACODE:	75249
SYNONYMS:	Familial or idiopathic restrictive cardiomyopathy
XREF(S):	Orphanet OMIM OMIM OMIM OMIM ICD-10 OMIM OMIM
ANALYTE(S):	FLNC KIF20A TNNI3 TNNT2 MYPN

CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Cardiomyopathy, hereditary (gene panel)
- Cardiomyopathy: hypertrophic cardiomyopathy, dilated cardiomyopathy, restrictive cardiomyopathy, left ventricular non-compaction cardiomyopathy, arrhythmogenic right ventricular cardiomyopathy (gene panel)
- Cardiopathies, hereditary (gene panel)
- Hypertrophic cardiomyopathy (gene panel)

Related Laboratories

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

Related Analytes

- filamin C
- kinesin family member 20A
- myopalladin
- troponin I3, cardiac type
- troponin T2, cardiac type

Related Gene Panels

- Cardiomyopathy (genepanel) - UZA
- Cardiomyopathy, hereditary (208 genes) - VUB
- Cardiopathies, hereditary (102 genes) - KUL
- Hypertrophic cardiomyopathy (75 genes) - IPG
