Familial isolated dilated cardiomyopathy

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Familial isolated dilated cardiomyopathy. ORPHA:154

Familial isolated dilated cardiomyopathy is a rare, genetically heterogeneous cardiac disease characterized by dilatation leading to systolic and diastolic dysfunction of the left and/or right ventricles, causing heart failure or arrhythmia.