Familial long QT syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Familial long QT syndrome. ORPHA:768

Congenital long QT syndrome (LQTS) is a hereditary cardiac disease characterized by a prolongation of the QT interval at basal ECG and by a high risk of life-threatening arrhythmias.