Dilated cardiomyopathy-hypergonadotropic hypogonadism syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Dilated cardiomyopathy-hypergonadotropic hypogonadism syndrome. ORPHA:2229

This syndrome is characterized by the association of dilated cardiomyopathy and hypergonadotropic hypogonadism (DCM-HH).