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## Dilated cardiomyopathyhypergonadotropic hypogonadism syndrome

**INSERM** 

## Source

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

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

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