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Barth syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Barth syndrome. ORPHA:111

Barth syndrome (BT HS) is an inborn error of phospholipid metabolism characterized by dilated cardiomyopathy (DCM), skeletal myopathy, neutropenia, growth delay and organic aciduria.