Mucolipidosis type II

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Mucolipidosis type II. ORPHA:576

Mucolipidosis II (MLII) is a slowly progressive lysosomal disorder characterized by growth retardation, skeletal abnormalities, facial dysmorphism, stiff skin, developmental delay and cardiomegaly.