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Metachromatic leukodystrophy

INSERM

Source

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

Metachromatic leukodystrophy. ORPHA:512

Metachromatic leukodystrophy (MLD) is a rare lysosomal storage disorder characterized by intralysosomal accumulation of sulfatides in various tissues, leading to progressive deterioration of motor and neurocognitive function.