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Fabry disease

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Fabry disease. ORPHA:324

Fabry disease (FD) is a progressive, inherited, multisystemic lysosomal storage disease characterized by specific neurological, cutaneous, renal, cardiovascular, cochleo-vestibular and cerebrovascular manifestations.