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Fibronectin glomerulopathy

INSERM

Source

INSERM. (1999). *Orphanet: an online rare disease and orphan drug data base*. [*Fibronectin glomerulopathy*](#). ORPHA:84090

Fibronectin glomerulopathy is a hereditary kidney disease characterized by proteinuria, type IV renal tubular acidosis, microscopic hematuria and hypertension that may lead to end-stage renal failure in the second to sixth decade of life.