FLOTCH syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. FLOTCH syndrome. ORPHA:2045

FLOTCH syndrome is a rare, genetic, cutaneous disorder characterized by leuchonychia and multiple, recurrent pilar cysts, associated or not with ciliar dystrophy and/or koilonychia. Renal calculi have also been reported.