Dacryocystitis-osteopoikilosis syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Dacryocystitis-osteopoikilosis syndrome. ORPHA:1562

Dacryocystitis - osteopoikilosis is an exceedingly rare autosomal dominant disorder reported in only a few patients to date and is characterized by dacryocystitis due to lacrimal canal stenosis,and osteopoikilosis (demonastratedradiologically as discrete spherical osteosclerotic lesions of 2-10mm in diameter).