Saldino-Mainzer syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Saldino-Mainzer syndrome. ORPHA:140969

Saldino-Mainzer syndrome is characterised by the association of renal disease, retinal pigmentary dystrophy, cerebellar ataxia and skeletal dysplasia.