Open Peer Review on Qeios

Lipoid proteinosis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Lipoid proteinosis. ORPHA:530

Lipoid proteinosis (LP) is a rare genodermatosis characterized clinically by mucocutaneous lesions, hoarseness developing in early childhood and, at times, neurological complications.