Johnson neuroectodermal syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Johnson neuroectodermal syndrome. ORPHA:2316

Johnson neuroectodermal syndrome is characterised by alopecia, anosmia or hyposmia, conductive deafness with malformed ears and microtia and/or atresia of the external auditory canal, and hypogonadotropic hypogonadism.