Lesch-Nyhan syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Lesch-Nyhan syndrome. ORPHA:510

Lesch-Nyhan syndrome (LNS) is the most severe form of hypoxanthine-guanine phosphoribosyltransferase (HPRT) deficiency (see this term), a hereditary disorder of purine metabolism, and is associated with uric acid overproduction (UAO), neurological troubles, and behavioral problems.