Wolcott-Rallison syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Wolcott-Rallison syndrome. ORPHA:1667

Wolcott-Rallison syndrome (WRS) is a very rare genetic disease, characterized by permanent neonatal diabetes mellitus (PNDM) with multiple epiphyseal dysplasia and other clinical manifestations, including recurrent episodes of acute liver failure.