Open Peer Review on Qeios

Hartnup disease

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hartnup disease. ORPHA:2116

Hartnup disease is a rare metabolic disorder belonging to the neutral aminoacidurias and characterized by abnormal renal and gastrointestinal transport of neutral amino acids (tryptophan, alanine, asparagine, glutamine, histidine, isoleucine, leucine, phenylalanine, serine, threonine, tyrosine and valine).