Open Peer Review on Qeios

Tyrosinemia type 3

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Tyrosinemia type 3. ORPHA:69723

Tyrosinemia type 3 is an inborn error of tyrosine metabolism characterised by mild hypertyrosinemia and increased urinary excretion of 4-hydroxyphenylpyruvate, 4-hydroxyphenyllactate and 4-hydroxyphenylacetate.