Mild phenylketonuria

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Mild phenylketonuria. ORPHA:79253

Mild phenylketonuria is a rare form of phenylketouria (PKU, see this term), an inborn error of amino acid metabolism, characterized by symptoms of PKU of mild to moderate severity.