Mild hyperphenylalaninemia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Mild hyperphenylalaninemia. ORPHA:79651

Mild hyperphenylalaninemia (HPA) is a rare form of phenylketonuria (see this term), an inborn error of amino acid metabolism, characterized by mild symptoms of HPA.