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# Fructose-1,6-bisphosphatase deficiency

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Fructose-1,6-bisphosphatase deficiency. ORPHA:348*

Fructose-1,6-biphosphatase (FBP) deficiency is a disorder of fructose metabolism (see this term) characterized by recurrent episodes of fasting hypoglycemia with lactic acidosis, that may be life-threatening in neonates and infants.