

**HMBS, 1-361aa, Human, His tag, E.coli**

**Cat.NO.: TP02465**

3th Edition

**Synonyms:** Porphobilinogen deaminase, PBG-D, PBGD, PORC, UPS.

**Description:** Porphobilinogen deaminase, also known as HMBS, is a member of the hydroxymethylbilane synthase superfamily. It is a cytoplasmic enzyme found in the heme synthesis pathway. Deficiency of HMBS causes errors in pyrrole metabolism which in turn leads to an inherited autosomal disorder called acute intermittent porphyria (AIP) which is characterized by acute attacks of neurological dysfunctions with hypertension, tachycardia, peripheral neurologic disturbances, abdominal pain and excessive amounts of aminolevulinic acid and porphobilinogen in the urine.

**Form:** Liquid. In 20mM Tris-HCl buffer (pH 8.0) containing 1mM DTT, 10% glycerol, 0.1M NaCl

**Molecular Weight:** 41.9kDa (385aa) confirmed by MALDI-TOF

**Sequences:**

MGSSHHHHHSSGLVPRGSHMGSHMSGNGNAAATAEENSPKMRVIRVGTRKSQLARIQTDSVVATLKASYPGLQ  
FEIIMSTTGDKILDALSIGEKSLFTKELEHALEKNEVDLVVHSLKDLPTVLPPGFTIGAICKRENPHDAVVFHPKFV  
GKTLETLPKSVVGTSSLRRAAQLQRKFPHFLEFRSIRGNLNTRLRKLDEQQEFSAILATAGLQRMGWHNRVQGILH  
PEECMYAVGQAGLVEVRAKDQDILDVGLHDPETLLRCIAERAFLRHLEGGCSVPVAVHTAMKDGQLYLTGGV  
WSLDGSDSIQETMQATIHVPAQHEDGPEDDPQLVGITARNIPRGPQLAAQNGLISLANLLLSKGAKNILDVARQLND  
AH

**Purity:** > 95% by HPLC

**Concentration:** 1 mg/ml (determined by Bradford assay)

**Endotoxin Level:** <1.0 EU per 1 ug of protein (determined by LAL method)

**Storage:** Can be stored at +4°C short term (1-2 weeks). For long term storage, aliquot and store at -20°C or -70°C. Avoid repeated freezing and thawing cycles.