HGD Recombinant Rabbit mAb

Catalog: BS50067 Host: Rabbit Reactivity: Human

BackGround:

This gene encodes the enzyme homogentisate 1,2 dioxy-genase. This enzyme is involved in the catabolism of the amino acids tyrosine and phenylalanine. Mutations in this gene are the cause of the autosomal recessive metabolism disorder alkaptonuria.[provided by RefSeq, May 2010]

Product:

1mg/ml in 50mM Tris-Glycine(pH 7.4), 0.15M NaCl, 40% Glycerol, 0.01% sodium azide and 0.05% BSA.

Molecular Weight:

50 kDa

Swiss-Prot:

Q93099

Purification&Purity:

Affinity Purification

Applications:

WB: 1:1000-1:5000
IHC-P: 1:200-1:500
ICC/IF: 1:200-1:500

Storage&Stability:

Store at 4℃ short term. Aliquot and store at -20℃ long term. Avoid freeze-thaw cycles.

Specificity:

The antibody detects endogenous levels of it.

DATA:

Note:

For research use only, not for use in diagnostic procedure.

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