

## **Product datasheet for TP303742**

## OriGene Technologies, Inc.

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## HBA2 (NM\_000517) Human Recombinant Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Recombinant protein of human hemoglobin, alpha 2 (HBA2), 20 µg

Species: Human
Expression Host: HEK293T

**Expression cDNA Clone** >RC203742 protein sequence or AA Sequence: Red=Cloning site Green=Tags(s)

MVLSPADKTNVKAAWGKVGAHAGEYGAEALERMFLSFPTTKTYFPHFDLSHGSAQVKGHGKKVADALTN

Α

VAHVDDMPNALSALSDLHAHKLRVDPVNFKLLSHCLLVTLAAHLPAEFTPAVHASLDKFLASVSTVLTSK

YR

**TRTRPLEQKLISEEDLAANDILDYKDDDDKV** 

Tag: C-Myc/DDK

Predicted MW: 15.1 kDa

**Concentration:** >0.05 μg/μL as determined by microplate BCA method

**Purity:** > 80% as determined by SDS-PAGE and Coomassie blue staining

**Buffer:** 25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol

**Preparation:** Recombinant protein was captured through anti-DDK affinity column followed by

conventional chromatography steps.

**Note:** For testing in cell culture applications, please filter before use. Note that you may experience

some loss of protein during the filtration process.

Storage: Store at -80°C.

Stability: Stable for 12 months from the date of receipt of the product under proper storage and

handling conditions. Avoid repeated freeze-thaw cycles.

**RefSeq:** NP 000508

**Locus ID:** 3040

UniProt ID: P69905





RefSeq ORF:

RefSeq Size: 622

Cytogenetics: 16p13.3

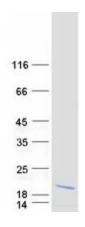
Synonyms: ECYT7; HBA-T2; HBH

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Summary: The human alpha globin gene cluster located on chromosome 16 spans about 30 kb and

includes seven loci: 5'- zeta - pseudozeta - mu - pseudoalpha-1 - alpha-2 - alpha-1 - theta - 3'. The alpha-2 (HBA2) and alpha-1 (HBA1) coding sequences are identical. These genes differ slightly over the 5' untranslated regions and the introns, but they differ significantly over the 3' untranslated regions. Two alpha chains plus two beta chains constitute HbA, which in normal adult life comprises about 97% of the total hemoglobin; alpha chains combine with delta chains to constitute HbA-2, which with HbF (fetal hemoglobin) makes up the remaining 3% of adult hemoglobin. Alpha thalassemias result from deletions of each of the alpha genes as well as deletions of both HBA2 and HBA1; some nondeletion alpha thalassemias have also been reported. [provided by RefSeq, Jul 2008]

## **Product images:**



Coomassie blue staining of purified HBA2 protein (Cat# TP303742). The protein was produced from HEK293T cells transfected with HBA2 cDNA clone (Cat# [RC203742]) using MegaTran 2.0 (Cat# [TT210002]).