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Hemoglobin alpha Rabbit pAb

Catalog Number: bs-9889R

Target Protein: Hemoglobin alpha

Concentration: 1mg/ml

Form: Liquid Host: Rabbit

Clonality: Polyclonal

Isotype: IgG

Applications: IHC-P (1:100-500), IHC-F (1:100-500), IF (1:50-200)

Reactivity: Mouse (predicted:Human, Sheep, Cow)

Predicted MW: 15 kDa Entrez Gene: 3039 Swiss Prot: P69905

Source: KLH conjugated synthetic peptide derived from human Hemoglobin alpha: 1-100/142.

Purification: affinity purified by Protein A

Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.

Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.

Background: he 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

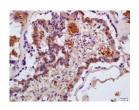
delta chains to constitute HbA-2, which with HbF (fetal hemoglobin) makes up the

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]

VALIDATION IMAGES



Tissue/cell: mouse placenta tissue; 4% Paraformaldehyde-fixed and paraffin-embedded; Antigen retrieval: citrate buffer (0.01M, pH 6.0), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum,C-0005) at 37°C for 20 min; Incubation: Anti-Hemoglobin alpha Polyclonal Antibody, Unconjugated(bs-9889R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining