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Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
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- Expressversand

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HBA1 Pre-design Chimera RNAi

Catalog # : H00003039-R01

規格 : [10 nmol] [20 nmol]

List All

Specification

Product Homo sapiens hemoglobin, alpha 1 (HBA1), mRNA.

Description:

Reactivity: Human

Supplied DEPC water

Product:

Target Refseq: NM_000558

Storage Store at -20°C, do not exceed 4 - 5 freeze-thaw cycles to ensure
Instruction: product integrity.

Note: Position of the Chimera RNAi.



Application Image

RNAi Knockdown

Publication Reference

- [dsCheck: highly sensitive off-target search software for double-stranded RNA-mediated RNA interference.](#)
Naito Y, Yamada T, Matsumiya T, Ui-Tei K, Saigo K, Morishita S. *Nucleic Acids Res.* 2005 Jul 1;33(Web Server issue):W589-91.
- [Functional dissection of siRNA sequence by systematic DNA substitution: modified siRNA with a DNA seed arm is a powerful tool for mammalian gene silencing with significantly reduced off-target effect.](#)
Ui-Tei K, Naito Y, Zenno S, Nishi K, Yamato K, Takahashi F, Juni A, Saigo K. *Nucleic Acids Res.* 2008 Apr;36(7):2136-51. Epub 2008 Feb 11.
- [Guidelines for the selection of highly effective siRNA sequences for mammalian and chick RNA interference.](#)
Ui-Tei K, Naito Y, Takahashi F, Haraguchi T, Ohki-Hamazaki H, Juni A, Ueda R, Saigo K. *Nucleic Acids Res.* 2004 Feb 9;32(3):936-48. Print 2004.
- [siDirect: highly effective, target-specific siRNA design software for mammalian RNA interference.](#)
Naito Y, Yamada T, Ui-Tei K, Morishita S, Saigo K. *Nucleic Acids Res.* 2004 Jul 1;32(Web Server issue):W124-9.

Applications

RNAi Knockdown

Gene Information

Entrez GeneID: 3039

Gene Name: HBA1

Gene Alias: HBH,HBA-T3

Gene hemoglobin, alpha 1

Description:

Omim ID: [141800](#)

Gene Ontology: [Hyperlink](#)

Gene 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]

Other Designations: alpha 1 globin,alpha one globin,alpha-1 globin,alpha-1-globin,hemoglobin alpha 1 globin chain,hemoglobin alpha-1 chain

Related Disease

[Abortion](#), [Spontaneous Albuminuria](#) [alpha-Thalassemia](#) [Anemia](#) [Anemia](#), [Aplastic Anemia](#), [Hypochromic Anemia](#), [Sickle Cell Anemia](#), [sickle cell](#) [beta-Thalassemia](#) [Dyslipidemias](#) [Fetal Diseases](#) [Genetic Predisposition to Disease](#) [Glucosephosphate Dehydrogenase Deficiency](#) [Hemolysis](#) [Hypertension](#), [Pulmonary Iron Overload](#) [Malaria](#) [Malaria](#), [Cerebral Malaria](#), [Falciparum](#)

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