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Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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EFEMP1 Pre-design Chimera RNAi

Catalog # : H00002202-R02

規格 : [10 nmol] [20 nmol]

List All

Specification

Product Description: Homo sapiens EGF containing fibulin-like extracellular matrix protein 1 (EFEMP1), transcript variant 3, mRNA

Reactivity: Human

Supplied Product: DEPC water

Target Refseq: NM_001039349.2

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

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: [2202](#)

Gene Name: EFEMP1

Gene Alias: DHRD, DRAD, FBLN3, FBNL, FLJ35535, MGC111353, MLVT, MTLV, S1-5

Gene Description: EGF-containing fibulin-like extracellular matrix protein 1

Omim ID: [126600](#), [601548](#)

Gene Ontology: [Hyperlink](#)

Gene Summary: This gene spans approximately 18 kb of genomic DNA and consists of 12 exons. Alternative splice patterns in the 5' UTR result in three transcript variants encoding the same extracellular matrix protein. Mutations in this gene are associated with Doyme honeycomb retinal dystrophy. [provided by RefSeq]

Other fibrillin-like, fibulin 3

Designations:

Related Disease

[Atherosclerosis](#) [Calcinosis](#) [Coronary Artery Disease](#) [Crohn Disease](#)
[Diabetes Mellitus, Type 2](#) [Genetic Predisposition to Disease](#) [Growth Disorders](#)
[Macular Degeneration](#) [Retinal Degeneration](#) [Retinal Diseases](#) [Retinal Drusen](#)
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