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Produktinformation

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- Diagnostik & molekulare Diagnostik
- Laborgeräte & Service

Weitere Information auf den folgenden Seiten!
See the following pages for more information!

Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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E-mail: sales@abnova.com

Datasheet

SOST (Human) Recombinant Protein

Catalog Number: P10578

Regulation Status: For research use only (RUO)

Product Description: Human SOST partial recombinant protein with His tag expressed in CHO cells.

Sequence:
QGWQAFKNDATEIIPELGEYPEPPPELENNKTMNRAE
NGGRPPHHPFETKDVSEYSCRELHFTRYVTDGPCRS
AKPVTELVCSGQCGPARLLPNAIGRGKWWRPSGPDF
RCIPDRYRAQRVQLLCPGGEAPRARKVRLVASCKCKR
LTRFHNQSELKDFGTEAARPQKGRKPRPRARSAKAN
QAELENAY

Host: Mammals

Theoretical MW (kDa): 22.33

Protocols: See our web site at http://www.abnova.com/support/protocols.asp or product page for detailed protocols

Form: Lyophilized

Preparation Method: Mammalian cell (CHO) expression system

Purification: Ni-NTA chromatography

Purity: > 85% by SDS-PAGE

Recommend Usage: SDS-PAGE
The optimal working dilution should be determined by the end user.

Storage Buffer: Lyophilized from filtered PBS, pH 7.4.

Storage Instruction: Store at -20°C for 12 months in lyophilized state.
After reconstitution with deionized water, store at -20 or -80°C for 1 month.
Aliquot to avoid repeated freezing and thawing.

Entrez GeneID: 50964

Gene Symbol: SOST

Gene Alias: VBCH

Gene Summary: Sclerostin is a secreted glycoprotein with a C-terminal cysteine knot-like (CTCK) domain and sequence similarity to the DAN (differential screening-selected gene aberrative in neuroblastoma) family of bone morphogenetic protein (BMP) antagonists. Loss-of-function mutations in this gene are associated with an autosomal-recessive disorder, sclerosteosis, which causes progressive bone overgrowth. A deletion downstream of this gene, which causes reduced sclerostin expression, is associated with a milder form of the disorder called van Buchem disease. [provided by RefSeq]