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Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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Datasheet

RUNX2 (Human) Recombinant Protein (Q02)

Catalog Number: H00000860-Q02

Regulation Status: For research use only (RUO)

Product Description: Human RUNX2 partial ORF (NP_001019801.1, 311 a.a. - 450 a.a.) recombinant protein with GST-tag at N-terminal.

Sequence:
TSPSIHSTTPLSSTRGTGLPAITDVPRRISDDDTATSDF CLWPSTLSKKSQAGASELGPFSDPRQFPSISSLTESRF SNPRMHYPATFTYTPPVTSGMSLGMSATTHYHTYLPP PYPGSSQSQSGPFQTSSTPYLYYGTS

Host: Wheat Germ (in vitro)

Theoretical MW (kDa): 41.14

Applications: AP, Array, ELISA, WB-Re
(See our web site product page for detailed applications information)

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

Preparation Method: *in vitro* wheat germ expression system

Purification: Glutathione Sepharose 4 Fast Flow

Storage Buffer: 50 mM Tris-HCl, 10 mM reduced Glutathione, pH=8.0 in the elution buffer.

Storage Instruction: Store at -80°C. Aliquot to avoid repeated freezing and thawing.

Entrez GeneID: 860

Gene Symbol: RUNX2

Gene Alias: AML3, CBFA1, CCD, CCD1, MGC120022, MGC120023, OSF2, PEA2aA, PEBP2A1, PEBP2A2, PEBP2aA, PEBP2aA1

Gene Summary: This gene is a member of the RUNX family of transcription factors and encodes a nuclear protein with an Runt DNA-binding domain. This protein is essential for osteoblastic differentiation and skeletal morphogenesis and acts as a scaffold for nucleic acids and regulatory factors involved in skeletal gene expression. The protein can bind DNA both as a monomer or, with more affinity, as a subunit of a heterodimeric complex. Mutations in this gene have been associated with the bone development disorder cleidocranial dysplasia (CCD). Transcript variants that encode different protein isoforms result from the use of alternate promoters as well as alternate splicing. [provided by RefSeq]