

Product Data Sheet

Animal-Free GDF-5 Protein, Human (His)

Cat. No.: HY-P700023AF

Synonyms: Cartilage-Derived Morphogenetic Protein-1; Growth/Differentiation Factor-5; GDF-5; CDMP-1

Species: Source: E. coli

P43026 (A382-R501) Accession:

Gene ID: 8200

Molecular Weight: Approximately 14.52 kDa

PROPERTIES

AA Sequence

MAPLATRQGK RPSKNLKARC SRKALHVNFK DMGWDDWIIA PLEYEAFHCE GLCEFPLRSH LEPTNHAVIQ TLMNSMDPES TPPTCCVPTR LSPISILFID SANNVVYKQY EDMVVESCGC

Measure by its ability to induce alkaline phosphatase production by ATDC5 cells. The ED₅₀ for this effect is <14 ng/mL **Biological Activity**

Lyophilized powder. **Appearance**

Formulation Lyophilized from a solution containing 20 mM sodium citrate, 0.2 MNaCl, pH 3.5.

Endotoxin Level <0.1 EU per 1 μ g of the protein by the LAL method.

Reconsititution It is not recommended to reconstitute to a concentration less than 100 $\mu g/mL$ in ddH₂O.

Storage & Stability Stored at -20°C for 2 years. After reconstitution, it is stable at 4°C for 1 week or -20°C for longer (with carrier protein). It is recommended to freeze aliquots at -20°C or -80°C for extended storage.

Shipping Room temperature in continental US; may vary elsewhere.

DESCRIPTION

Background

GDF-5 Protein plays a pivotal role in bone and cartilage formation, intricately regulating chondrogenic tissue differentiation through dual pathways. Firstly, it positively influences chondrogenic tissue differentiation by binding with high affinity to BMPR1B and with lower affinity to BMPR1A, leading to the induction of SMAD1-SMAD5-SMAD8 complex phosphorylation and subsequent SMAD protein signaling transduction. Simultaneously, it negatively regulates chondrogenic differentiation through interaction with NOG. This protein is essential for preventing excessive muscle loss upon denervation, a function mediated by phosphorylated SMAD1/5/8. Additionally, GDF-5 binds bacterial lipopolysaccharide (LPS) and mediates LPSinduced inflammatory responses, including TNF secretion by monocytes. Structurally, it forms homodimers linked by

disulfide bonds and interacts with serine proteases HTRA1 and HTRA3. Following LPS binding, GDF-5 may form a complex with CXCR4, HSP90AA1, and HSPA8. Moreover, it interacts with high affinity with NOG to inhibit chondrogenesis and with BMPR1B and BMPR1A to positively regulate chondrocyte differentiation through SMAD-dependent signaling. Additionally, it engages in interactions with FBN1 and FBN2.

Caution: Product has not been fully validated for medical applications. For research use only.

Tel: 609-228-6898 Fax: 609-228-5909 E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA

Page 2 of 2 www.MedChemExpress.com