

**DESCRIPTION**

<b>Source</b>	<i>E. coli</i> -derived	
	Human BMP-2 (Ala284 - Arg396), with an N-terminal Met Accession # P12643	
	Human BMP-6 (Gln382 - His513), with an N-terminal Met Accession # P22004	
	N-terminus	C-terminus

<b>N-terminal Sequence Analysis</b>	Ala284 (BMP-2) & Met (BMP-6)
<b>Structure / Form</b>	Disulfide-linked heterodimer
<b>Predicted Molecular Mass</b>	12.8 kDa (BMP-2) & 15 kDa (BMP-6)

**SPECIFICATIONS**

<b>SDS-PAGE</b>	11 kDa & 14 kDa, reducing conditions
<b>Activity</b>	Measured by its ability to induce alkaline phosphatase production by ATDC5 mouse chondrogenic cells. Binnerts, M.E. <i>et al.</i> (2004) <i>Biochem. Biophys. Res. Commun.</i> <b>315</b> (2):272. The ED <sub>50</sub> for this effect is 4-20 ng/mL.
<b>Endotoxin Level</b>	<0.01 EU per 1 µg of the protein by the LAL method.
<b>Purity</b>	>95%, by SDS-PAGE under reducing conditions and visualized by silver stain.
<b>Formulation</b>	Lyophilized from a 0.2 µm filtered solution in HCl. See Certificate of Analysis for details.

**PREPARATION AND STORAGE**

<b>Reconstitution</b>	Reconstitute at 100 µg/mL in 4 mM HCl.
<b>Shipping</b>	The product is shipped with polar packs. Upon receipt, store it immediately at the temperature recommended below.
<b>Stability &amp; Storage</b>	<b>Use a manual defrost freezer and avoid repeated freeze-thaw cycles.</b> <ul style="list-style-type: none"> <li>● 12 months from date of receipt, -20 to -70 °C as supplied.</li> <li>● 1 month, 2 to 8 °C under sterile conditions after reconstitution.</li> <li>● 3 months, -20 to -70 °C under sterile conditions after reconstitution.</li> </ul>

**BACKGROUND**

Bone Morphogenetic Protein 6 (BMP-6), also known as Vgr-1, and BMP-2 are members of the BMP family of structurally and functionally related proteins and represent a subfamily of the transforming growth factor β (TGF-β) superfamily. BMPs are involved in a wide range of processes including embryogenesis, tissue morphogenesis, cell differentiation and migration, and tumorigenesis. Cellular responses to BMPs are mediated by hetero-oligomeric complexes of type I and type II serine/threonine kinase receptors (1 - 4). Human BMP-2 is synthesized as a 396 amino acid (aa) preproprotein that contains a 23 aa signal sequence, a 259 aa prosegment, and a 114 aa mature region (5). Human BMP-6 is synthesized as a 513 aa precursor protein that contains a 20 aa signal sequence, a 354 aa prosegment, and a 139 aa mature region (6). BMP prosegments are removed by proteolysis, enabling the glycosylated 18 kDa mature BMPs to form active disulfide-linked homodimers or heterodimers (1, 2). Mature human BMP-2 shares 100% aa sequence identity with mouse and rat BMP-2, and mature human BMP-6 shares 96% and 98% aa sequence identity with mouse and rat BMP-6, respectively. They share 48% aa sequence identity with each other. Both BMP-2 and BMP-6 induce osteogenic and chondrogenic differentiation in mesenchymal stem cells (4). Heterodimers of BMP-2 and BMP-6 show increased potency at inducing osteoblastic calcium deposition, chondrogenesis, and *in vivo* bone formation compared to either BMP-2 or BMP-6 homodimers (7, 8). BMP-2/6 heterodimers also show increased activity at inducing trophoectodermal and endodermal differentiation of embryonic stem cells compared to either homodimer (9).

**References:**

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