Product Name: Recombinant Human BGN (C-6His)

Catalog #: PHH0153



Summary

Name Biglycan

Purity Greater than 95% as determined by reducing SDS-PAGE

Endotoxin level <1 EU/μg as determined by LAL test.

Construction Recombinant Human Biglycan is produced by our Mammalian expression

system and the target gene encoding Glu20-Lys368 is expressed with a 6His

tag at the C-terminus.

Accession # P21810

Host Human Cells

Species Human

Predicted Molecular Mass 40.47 KDa

Formulation Lyophilized from a 0.2 μm filtered solution of 20mM PB, 150mM NaCl, pH 7.2.

Shipping The product is shipped at ambient temperature. Upon receipt, store it

immediately at the temperature listed below.

Stability&Storage Store at \leq -70°C, stable for 6 months after receipt. Store at \leq -70°C, stable for 3

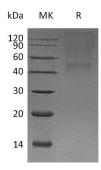
months under sterile conditions after opening. Please minimize freeze-thaw

cycles.

Reconstitution Always centrifuge tubes before opening. Do not mix by vortex or pipetting. It is

not recommended to reconstitute to a concentration less than 100µg/ml. Dissolve the lyophilized protein in distilled water. Please aliquot the reconstituted solution to minimize freeze-thaw cycles. Always centrifuge tubes before opening. Do not mix by vortex or pipetting. It is not recommended to reconstitute to a concentration less than 100µg/ml. Dissolve the lyophilized protein in distilled water. Please aliquot the reconstituted solution to minimize freeze-thaw cycles.

SDS-PAGE image



Background

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SHIS) CENKILIFE

Alternative Names Biglycan; Bone/Cartilage Proteoglycan I; PG-S1; BGN; SLRR1A

Background Biglycan is a 200-350 kD proteoglycan consisting of a 45 kD core protein and two

chrondroitin/dermatan sulfate glycosaminoglycan chains. Biglycan binds to TGF-ß. It also binds to collagen type I in low ionic strength (less than 3 mM phosphate) buffer. At higher ionic strengths, Biglycan does not bind to collagen type I. It enhances the inhibition effect of TGF-ß on osteoclast proliferation at a concentration of 4-20 mg/ml. It also prevents the attachment of CHO cells to

fibronectin, with a 50% inhibition at 17-21 mg/ml.

Note

For Research Use Only, Not for Diagnostic Use.

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