

Recombinant Mouse Noggin

Information

Gene ID	18121
Accession #	P97466
Alternate Names	
Source	Escherichia coli.
M.Wt	Approximately 46.4 kDa, a disulfide-linked homodimer consisting of two 206 amino acid polypeptide chains.
AA Sequence	MQHYLHIRPA PSDNLPLVDL IEHPDPIFDP KEKDLNETLL RSLLGGHYDP GFMATSPPED RPGGGGGPAG GAEDLAELDQ LLRQRPSGAM PSEIKGLEFS EGLAQGKKQR LSKKLRRKLQ MWLWSQTFCP VLYAWNDLGS RFWPRYVKVG SCFSKRSCSV PEGMVCKPSK SVHLTVLRWR CQRRGGQRCG WIPIQYPIIS ECKCSC
Appearance	Sterile Filtered White lyophilized (freeze-dried) powder.
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles.
	- 12 months from date of receipt, -20 to -70 °C as supplied.
	 1 month, 2 to 8 °C under sterile conditions after reconstitution. 3 months, -20 to -70 °C under sterile conditions after reconstitution.
Formulation	Lyophilized from a 0.2 μ m filtered concentrated solution in 30 % acetonitrile, 0.1 % TFA.
Reconstitution	We recommend that this vial be briefly centrifuged prior to opening to bring the contents to the bottom. Reconstitute in 10 mM HAc to a concentration less than 0.25 mg/mL. Stock solutions should be apportioned into working aliquots and stored at \leq -20 °C. Further dilutions should be made in appropriate buffered solutions.
Biological Activity	Fully biologically active when compared to standard. The ED ₅₀ as determined by inhibiting BMP-4-induced alkaline phosphatase production of murine ATDC5 cells is less than 2 ng/ml, corresponding to a specific activity of > 5.0×10^5 IU/mg in the presence of 5 ng/ml BMP-4.
Shipping Condition	Gel pack.
Handling	Centrifuge the vial prior to opening.
Usage	For Research Use Only! Not to be used in humans.

Components and Storage

Components	5 µg	100 µg	500 µg
Recombinant Mouse Noggin	5 µg	100 µg	500 µg

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Quality Control	19 million	al Que
Purity	> 95 % by SDS-PAGE and HPLC analyses.	Provintered a
Endotoxin	Less than 1 EU/µg of rMuNoggin as determin	ed by LAL method.

Description

Noggin encoded by the NOG gene, was first isolated from Xenopus, having the function of inducing secondary axis formation in frog embryos. It inhibits TGF-β family ligands and preventing them from binding to their corresponding receptors. Noggin was originally found as a BMP-4 antagonist, and then has been shown to modulate the activities of other BMPs (BMP-2, 7, 13 and 14). Additionally, it has pleiotropic effect, both in early development and later stages. The results of the mouse knockout of noggin suggest that it is involved in numerous developmental processes, such as neural tube fusion and joint formation. In recent report, proximal symphalangism (SYM1) and multiple synostoses syndrome (SYNS1) have relation with the mutant of evolutionarily conserved amino acid residues of Noggin. Mature mouse Noggin shares 99 % and 83 % a.a. sequence identity with human and Xenopus Noggin, respectively.

Reference

- 1. Davis SWandCamper SA. 2007. Dev Biol, 305: 145-60.
- 2. Zhu W, Kim J, Cheng C, et al. 2006. Bone, 39: 61-71.
- 3. Oxley CD, Rashid R, Goudie DR, et al. 2008. Horm Res, 69: 221-6.
- 4. Cooper GM, Usas A, Olshanski A, et al. 2009. Plast Reconstr Surg, 123: 94S-103S.
- 5. Bayramov AV, Eroshkin FM, Martynova NY, et al. 2011. Development, 138: 5345-56.

