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Recombinant Human GDNF

Catalog#:P02128 Derived from *E.coli*

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DESCRIPTION	Recombinant Human Glial Cell Line- Derived Neurotrophic Factor is produced by our <i>E.coli</i> expression system and the target gene encoding Ser78- Ile211 is expressed.				
	Accession#: P39905				
	Known as: Glial Cell Line- Derived Neurotrophic Factor; hGDNF; Astrocyte- Derived Trophic Factor; ATF; GDNF				
FORMULATION	Lyophilized from a 0.2 µm filtered solution of 20mM PB, 150mM NaCl, pH 7.4.				
SHIPPING	The product is shipped at ambient temperature.				
	Upon receipt, store it immediately at the temperature listed below.				
STORAGE	Lyophilized protein should be stored at <-20°C, though stable at room				
	temperature for 3 weeks.				
	Reconstituted protein solution can be stored at 4-7°C for 2-7 days.				
	Aliquots of reconstituted samples are stable at < -20°C for 3 months.				
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.				
QUALITY	Mol Mass:15.1kDa AP Mol Mass:17kDa, reducing conditions. Purity: Greater than 95% as determined by reducing SDS-PAGE.				
CONTROL	Endotoxin: Less than 0.1 ng/ μ g (1 EU/ μ g) as determined by LAL test.				
	Glial Cell Line-Derive	.1 11g/p	urotror	or μg) as determined vic Factor (GDNF)	is a disulfide-linked
BACKGROUND	Glial Cell Line-Derived Neurotrophic Factor (GDNF) is a disulfide-linked homodimeric glycoprotein that belongs to the TGF- β superfamily. It has been shown to promote the survival of various neuronal subpopulations in both the central as well as the peripheral nervous systems at different stages of their development. Human GDNF cDNA encodes a 211 amino acid residue prepropeptide that is processed to yield a dimeric protein. Mature human GDNF was predicted to contain two 134 amino acid residue subunits. Cells known to express GDNF include Sertoli cells, type 1 astrocytes, Schwann cells, neurons, pinealocytes and skeletal muscle cells. Mutations in this gene may be associated with Hirschsprung disease.				
		kDa	MK	R	
		120 90		11-1	
		60	-		
		40	-		
	SDS-PAGE	30	Manage		
		20	Name of		
		14	-	10-20	