





Recombinant Human Rnase T2/Ribonuclease T2(C-6His) Catalog#:P01786 Derived from Human Cells

Catalog#:P01786

Catalog#.F01/80 Delived Holli Hullian Cells	
	Recombinant Human Ribonuclease T2 is produced by our Mammalian
DESCRIPTION	expression system and the target gene encoding Asp25-His256 is expressed with
	a 6His tag at the C-terminus.
	Accession#: O00584
	Known as: Ribonuclease T2;3.1.27;Ribonuclease 6;RNASE6PL
FORMULATION	Supplied as a 0.2 µm filtered solution of 20mM Tris-HCl, 150mM NaCl, 20%
	Glycerol, pH 7.5
	The product is shipped on dry ice/polar packs.
SHIPPING	Upon receipt, store it immediately at the temperature listed below.
	Store at \leq -70°C, stable for 6 months after receipt.
STORAGE	Store at \leq -70°C, stable for 3 months under sterile conditions after opening.
	Please minimize freeze-thaw cycles.
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QUALITY	Mol Mass:28.2kDa AP Mol Mass:38-45kDa, 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.
BACKGROUND	RNASET2 (ribonuclease T2) is an enzyme which belongs to the RNase T2
	family. It is highly expressed in the temporal lobe and fetal brain.
	RNASET2 gene is a novel member of the Rh/T2/S-glycoprotein class
	of extracellular ribonucleases. This protein can be inhibited by Zn2+ and Cu2+. It
	has ribonuclease activity, with higher activity at acidic pH and is
	probably involved in lysosomal degradation of ribosomal
	RNA.Defects in RNASET2 are the cause of leukoencephalopathy cystic without
	megalencephaly. An infantile-onset syndrome of cerebral
	leukoencephalopathy. Affected newborns develop microcephaly
	and neurologic abnormalities including psychomotor impairment,
	seizures and sensorineural hearing impairment. The brain shows
	multifocal white matter lesions, anterior temporal lobe subcortical cysts,
	pericystic abnormal myelination, ventriculomegaly and intracranial
	calcifications.
LDS MAY 0	

