

LONM rabbit pAb

Catalog_no :	<u>AN3293</u>
Applications :	<u>WB</u>
Reactivity :	<u>Human, Mouse,Rat</u>
Category :	<u>抗原抗体</u>
Size :	<u>100µg/50µg/20µg</u>
Gene_name :	<u>LONP1 PRSS15</u>
Protein_name :	<u>LONM</u>
Humangene_id	<u>9361</u>
:	<u></u>
Humanswissprot	<u>P36776</u>
_no :	<u></u>
Mousegene_id :	<u>74142</u>
Mouseswissprot	<u>Q8CGK3</u>
_no :	<u></u>
Ratgene_id :	<u>170916</u>
Ratswissprot_no	<u>Q92455</u>
:	<u></u>
Immunogen :	<u>Synthesized peptide derived from human LONM</u>
Specificity :	<u>This antibody detects endogenous levels of LONM at Human/Mouse/Rat</u>
Formulation :	<u>Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.251% sodium azide.</u>
Source :	<u>Rabbit</u>
Dilution :	<u>WB 1:500-2000</u>
Purification :	<u>The antibody was affinity-purified from rabbit serum by affinity-chromatography using specific immunogen.</u>
Concentration :	<u>1 mg/ml</u>
Storage_stability	<u>-20°C/1 year</u>
:	<u></u>
Other_name :	<u>Lon protease homolog, mitochondrial (EC 3.4.21.-) (LONHs) (Lon protease-like protein) (LONP) (Mitochondrial ATP-dependent protease Lon) (Serine protease 15)</u>



Molecular Weight : 105KD

Background : This gene encodes a mitochondrial matrix protein that belongs to the Lon family of ATP-dependent proteases. This protein mediates the selective degradation of misfolded, unassembled or oxidatively damaged polypeptides in the mitochondrial matrix. It may also have a chaperone function in the assembly of inner membrane protein complexes, and participate in the regulation of mitochondrial gene expression and maintenance of the integrity of the mitochondrial genome. Decreased expression of this gene has been noted in a patient with hereditary spastic paraplegia (PMID:18378094). Alternatively spliced transcript variants have been found for this gene. [provided by RefSeq, Feb 2013],
