

Rabbit Anti-FAM61B antibody

SL11001R

Product Name:	FAM61B
Chinese Name:	FAM61B蛋白抗体 A State of the second sec
Alias:	Protein FAM61B; Putative uncharacterized protein C20orf40; LSM14 homolog B (SCD6, S. cerevisiae); bA11M20.3; C20orf40; Chromosome 20 open reading frame 40; Family with sequence similarity 61 member B; LSM13; LSM14 homolog B; LSM14B SCD6 homolog B (S. cerevisiae); MGC61931; Protein LSM14 homolog B; RNA associated protein 55B; LS14B_HUMAN; LSM14B; bA11M20.3; C20orf40; FAM61B; FT005; LSM13; RAP55B.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human,Mouse,Rat,Dog,Cow,
Applications:	WB=1:500-2000ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800Flow- Cyt=1µg/TestICC=1:100-500IF=1:100-500 (Paraffin sections need antigen repair) not yet tested in other applications. optimal dilutions/concentrations should be determined by the end user.
Molecular weight:	42kDa
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human FAM61B:251-350/385
Lsotype:	IgG
Purification:	affinity purified by Protein A
Storage Buffer:	0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.
Storage:	Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. The lyophilized antibody is stable at room temperature for at least one month and for greater than a year when kept at -20°C. When reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody is stable for at least two weeks at 2-4 °C.
PubMed:	PubMed
Product Detail:	Sm and Sm-like (LSm) proteins form donut-shaped, ubiquitously expressed heptameric complexes that are involved in various steps of RNA metabolism, including RNA-protein interactions and structural changes that are required during ribosomal subunit

assembly. LSm14B, also known as C20orf40, FAM61B or LSM13, is a 385 amino acid protein that exists as multiple alternatively spliced isoforms and may play a role in RNA-related events. The gene encoding LSm14B maps to human chromosome 20. Comprising approximately 2% of the human genome, chromosome 20 contains nearly 63 million bases that encode over 600 genes, some of which are associated with Creutzfeldt-Jakob disease, amyotrophic lateral sclerosis, spinal muscular atrophy, ring chromosome 20 epilepsy syndrome and Alagille syndrome.

Function:

May play a role in control of mRNA translation (By similarity).

Subunit: Component of a ribonucleoprotein (RNP) complex (By similarity).

opiotech Similarity: Belongs to the LSM14 family. Contains 1 DFDF domain.

SWISS: **O9BX40**

Gene ID: 149986

Database links:

Entrez Gene: 149986Human

Entrez Gene: 241846Mouse

SwissProt: Q9BX40Human

SwissProt: Q8CGC4Mouse

Important Note: This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.

