

Rabbit Anti-FAM71A antibody

SL11007R

Product Name:	FAM71A
Chinese Name:	FAM71A蛋白抗体
Alias:	Protein FAM71A; FA71A_HUMAN.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human, Mouse, Rat, Pig, Cow, Rabbit, Sheep,
Applications:	WB=1:500-2000ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800ICC=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:	63kDa
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human FAM71A:101-200/594
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:	<u>PubMed</u>
Product Detail:	Chromosome 1 is the largest human chromosome spanning about 260 million base pairs
	and making up 8% of the human genome. There are about 3,000 genes on chromosome
	1, and considering the great number of genes there are also a large number of diseases
	associated with chromosome 1. Notably, the rare aging disease Hutchinson-Gilford
	progeria is associated with the LMNA gene which encodes lamin A. When defective,
	the LMNA gene product can build up in the nucleus and cause characteristic nuclear
	blebs. The mechanism of rapidly enhanced aging is unclear and is a topic of continuing
	exploration. The MUTYH gene is located on chromosome 1 and is partially responsible

for familial adenomatous polyposis. Stickler syndrome, Parkinsons, Gaucher disease and Usher syndrome are also associated with chromosome 1. A breakpoint has been identified in 1q which disrupts the DISC1 gene and is linked to schizophrenia. Aberrations in chromosome 1 are found in a variety of cancers including head and neck cancer, malignant melanoma and multiple myeloma. The FAM71A gene product has been provisionally designated FAM71A pending further characterization.

Similarity:

Belongs to the FAM71 family.

SWISS:

Q8IYT1

Gene ID:

149647

Database links:

Entrez Gene: 149647Human

SwissProt: Q8IYT1Human

Unigene: 129293Human

Important Note:

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