

Rabbit Anti-ZNF687 antibody

SL16514R

Product Name:	ZNF687
Chinese Name:	Zinc finger protein687抗体
Alias:	4931408L03Rik; DKFZp78111719; KIAA1441; mKIAA1441; RP11-126K1.3; Zfp687; Zinc finger protein 687; ZN687 HUMAN; ZNF687.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human,
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:	129kDa
Cellular localization:	The nucleus 🤍
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human ZNF687:1001-1100/1237
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:	Zinc-finger proteins contain DNA-binding domains and have a wide variety of
	functions, most of which encompass some form of transcriptional activation or
	repression. The majority of zinc-finger proteins contain a kruppel-type DNA binding
	domain and a KRAB domain, which is thought to interact with KAP1, thereby recruiting
	histone modifying proteins. ZNF687 (zinc finger protein 687) is a 1,237 amino acid
	nuclear protein that is involved in transcriptional regulation. A member of the Krüppel
	nuclear protein that is involved in transcriptional regulation. A member of the Krupper

C2H2-type zinc-finger protein family, ZNF687 contains ten C2H2-type zinc fingers and exists as two alternatively spliced isoforms. The gene encoding ZNF687 maps to human chromosome 1, which comprises nearly 8% of the human genome and houses a large number of disease-associated genes, including those that are involved in familial adenomatous polyposis, Stickler syndrome, Parkinson's disease, Gaucher disease, schizophrenia and Usher syndrome.

Function: May be involved in transcriptional regulation.

Subcellular Location: Nucleus.

Similarity: Belongs to the krueppel C2H2-type zinc-finger protein family. Contains 10 C2H2-type zinc fingers. joiotect

SWISS: Q8N1G0

Gene ID: 57592

Database links:

Entrez Gene: 57592 Human

Entrez Gene: 78266 Mouse

Omim: 610568 Human

SwissProt: Q8N1G0 Human

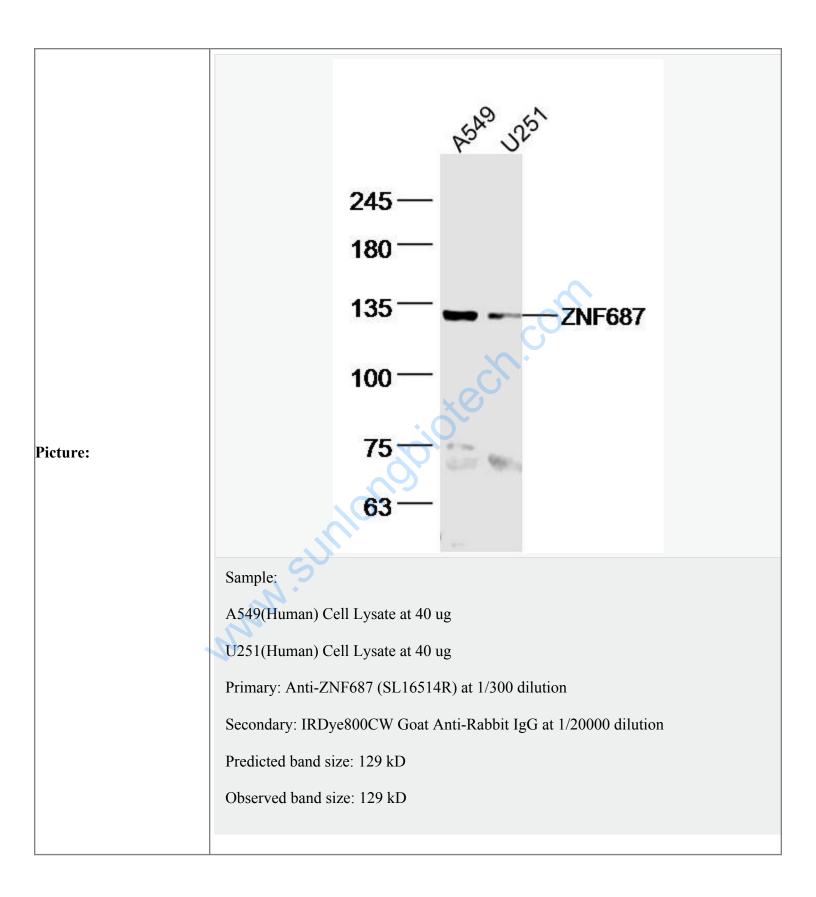
SwissProt: Q9D2D7 Mouse

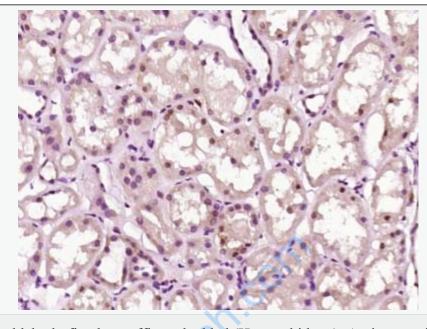
Unigene: 186756 Human

Unigene: 389478 Mouse

Important Note:

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





Paraformaldehyde-fixed, paraffin embedded (Human kidney); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (ZNF687) Polyclonal Antibody, Unconjugated (SL16514R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.