

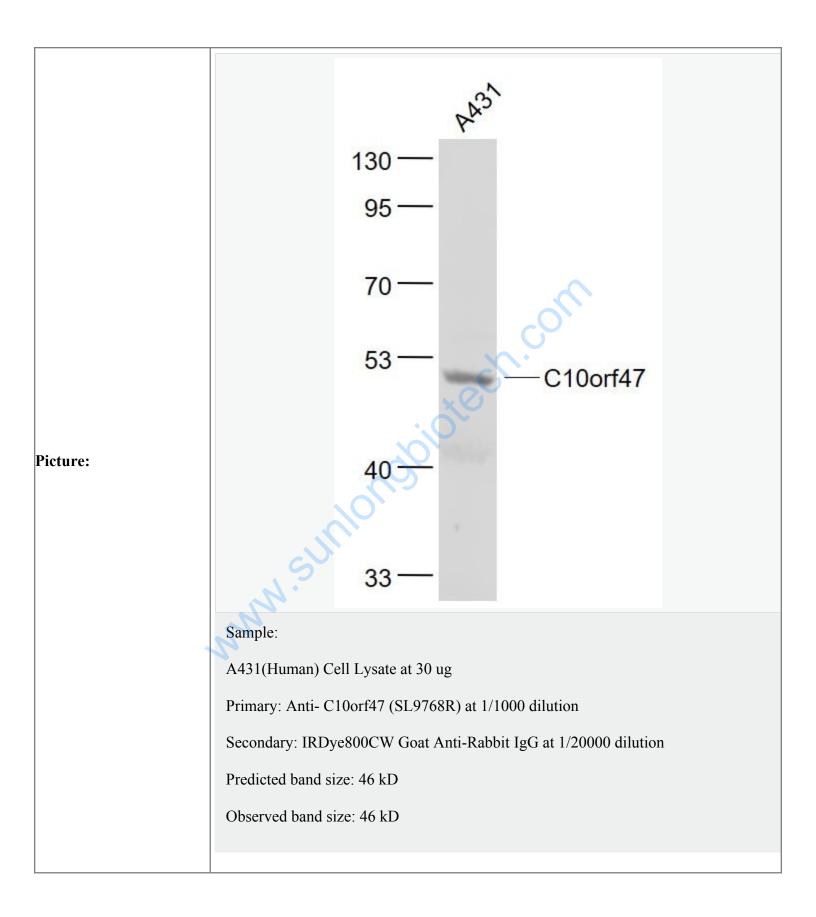
Rabbit Anti-C10orf47 antibody

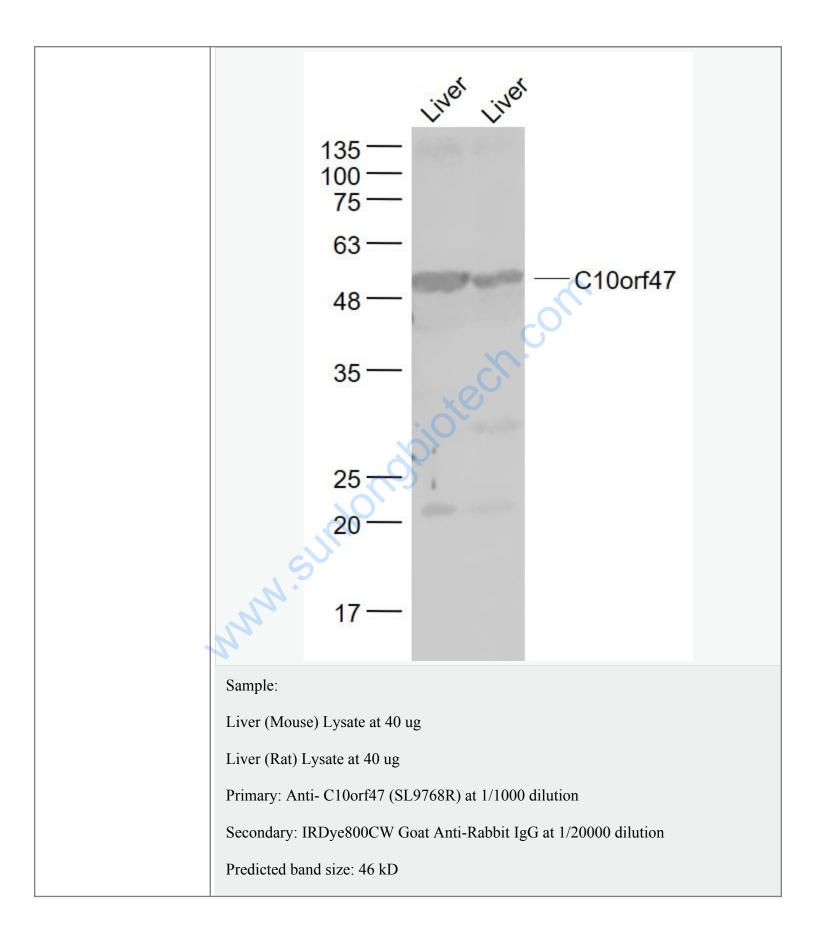
SL9768R

Product Name:	C10orf47
Chinese Name:	10 号染色体开放 阅读框47 抗体
Alias:	C10orf47; Chromosome 10 open reading frame 47; CJ047_HUMAN; Hypothetical
	protein LOC254427; MGC35403; Uncharacterized protein C10orf47.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human, Mouse, Rat, Pig, Horse, Rabbit, Sheep,
	WB=1:500-2000ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800IF=1:50-
Applications:	200 (Paraffin sections need antigen repair)
Applications:	not yet tested in other applications.
	optimal dilutions/concentrations should be determined by the end user.
Molecular weight:	46kDa
Cellular localization:	The nucleuscytoplasmicThe cell membraneExtracellular matrixSecretory protein
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human C10orf47:351-435/435
Lsotype:	IgG
Purification:	affinity purified by Protein A
Storage Buffer:	0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.
	Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. The lyophilized
Storage:	antibody is stable at room temperature for at least one month and for greater than a year
Stor age.	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
	C10orf47 is a 435 amino acid protein that exists as two alternatively spliced isoforms
	that are encoded by a gene located on human chromosome 10. Spanning nearly 135
Product Detail:	million base pairs, chromosome 10 makes up approximately 4.5% of total DNA in cells
I I JULICE D'ULAII.	and encodes nearly 1,200 genes. Several protein-coding genes, including those that
	encode for chemokines, cadherins, excision repair proteins, early growth response
	factors (Egrs) and fibroblast growth receptors (FGFRs), are located on chromosome 10.

Defects in some of the genes that map to chromosome 10 are associated with Charcot- Marie Tooth disease, Jackson-Weiss syndrome, Usher syndrome, nonsyndromatic deafness, Wolman's syndrome, Cowden syndrome, multiple endocrine neoplasia type 2 and porphyria.
SWISS: Q86WR7
Gene ID:
254427
Database links:
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Important Note:
This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.
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	Observed band size: 52 kD

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