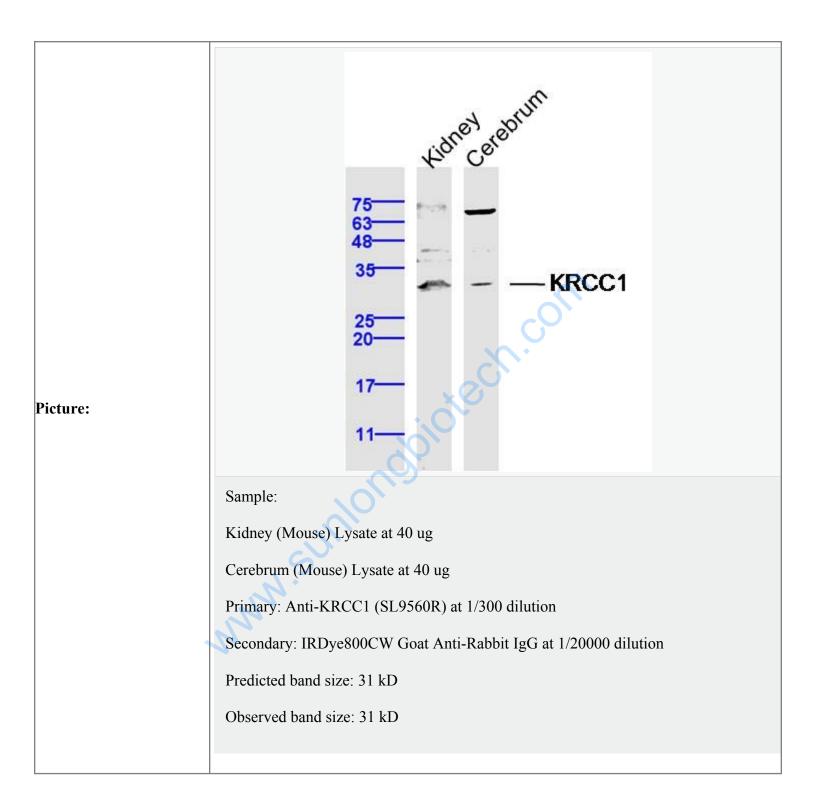


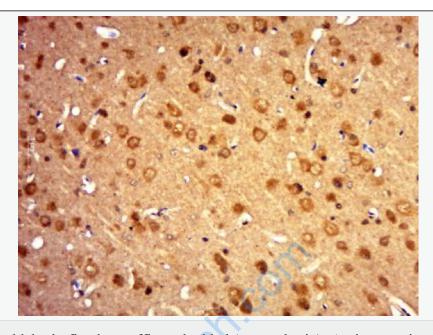
## Rabbit Anti-KRCC1 antibody

SL9560R

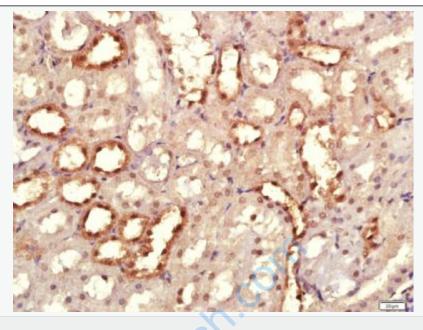
Product Name:	KRCC1
Chinese Name:	含赖氨酸卷曲螺旋蛋白1抗体
Alias:	CHBP2; cryptogenic hepatitis binding protein; Cryptogenic hepatitis-binding protein 2; KRCC1; KRCC1_HUMAN; lysine rich coiled coil 1; Lysine-rich coiled-coil protein 1.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human,Mouse,Rat,Cow,
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:	31kDa
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human KRCC1:161-259/259
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  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 for . When reconstituted in sterile pH 7.4 0.01 M PBS or diluent of antibody the antibody is stable for at least two weeks at 2-4  for .
PubMed:	PubMed
Product Detail:	KRCC1 is a 259 amino acid protein that is encoded by a gene located on human chromosome 2p11.2. Consisting of 237 million bases, chromosome 2 is the second largest human chromosome and encodes over 1,400 genes. A number of genetic diseases are linked to genes on chromosome 2. Harlequin icthyosis, a rare and morbid skin deformity, is associated with mutations in the ABCA12 gene. The lipid metabolic disorder sitosterolemia is associated with ABCG5 and ABCG8. An extremely rare

recessive genetic disorder, Alstr鰉 syndrome, is due to mutations in the ALMS1 gene. Interestingly, chromosome 2 contains what appears to be a vestigial second centromere and vestigial telomeres which gives credence to the hypothesis that human chromosome 2 is the result of an ancient fusion of two ancestral chromosomes seen in modern form today in apes.
SWISS: Q9NPI7
Gene ID: 51315
Database links:
Entrez Gene: 51315Human
SwissProt: Q9NPI7Human
Database links:   Entrez Gene: 51315Human   SwissProt: Q9NPI7Human   Unigene: 469254Human   Important Note:   This product as supplied is intended for research use only, not for use in human,
Important Note:
This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.
MMN.SUMORIS

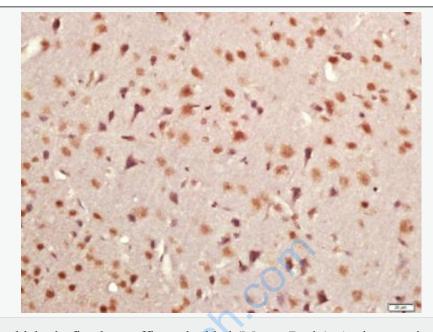




Paraformaldehyde-fixed, paraffin embedded (mouse brain); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (KRCC1) Polyclonal Antibody, Unconjugated (SL9560R) at 1:400 overnight at 4°C, followed by a conjugated secondary (sp-0023) for 20 minutes and DAB staining.



Paraformaldehyde-fixed, paraffin embedded (Mouse 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 (KRCC1) Polyclonal Antibody, Unconjugated (bs9560R) at 1:400 overnight at 4°C, followed by a conjugated secondary antibody (sp-0023) for 20 minutes and DAB staining.



Paraformaldehyde-fixed, paraffin embedded (Mouse Brain); 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 (KRCC1) Polyclonal Antibody, Unconjugated (bs9560R) at 1:400 overnight at 4°C, followed by a conjugated secondary antibody (sp-0023) for 20 minutes and DAB staining.