

Rabbit Anti-CCDC68 antibody

SL8086R

Product Name:	CCDC68
Chinese Name:	卷曲螺旋结构域蛋白68抗体(皮肤Tlymphocyte淋巴瘤相关抗原)
Alias:	CCD68_HUMAN; se57-1; CCDC 68; CCDC68; Coiled coil domain containing 68; Coiled coil domain containing protein 68; Coiled-coil domain-containing protein 68; CTCL associated antigen se57 1; CTCL tumor antigen se57 1; CTCL-associated antigen se57-1; Cutaneous T cell lymphoma associated antigen; Cutaneous T cell lymphoma associated antigen se57 1; Cutaneous T-cell lymphoma-associated antigen se57-1; FLJ25368; SE57 1.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human,Mouse,Rat,Dog,
Applications:	WB=1:500-2000ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800IF=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:	39kDa
Cellular localization:	The nucleuscytoplasmic
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from Human CCDC68/se57-1:255-335/335
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:	CCDC68, also known as CTCL tumor antigen se57-1, is a 335 amino acid protein expressed in cutaneous T-cell lymphoma (CTCL), bone marrow, colon, small intestine,

spleen, testis and trachea tissues. CTCL is a cancer of the immune system associated with mutation of the T cells. Malignant T cells form lesions on the surface of the skin. se57-1 contains 1 coiled coil domain and is encoded by the CCDC68 gene mapping to human chromosome 18. Chromosome 18 houses over 300 protein-coding genes and contains nearly 76 million bases. There are a variety of diseases associated with defects in chromosome 18-localized genes, some of which include Trisomy 18 (also known as Edwards syndrome), Niemann-Pick disease, hereditary hemorrhagic telangiectasia, erythropoietic protoporphyria and follicular lymphomas.

Tissue Specificity:

Expressed in bone marrow, colon, small intestine, spleen, testis, trachea and cutaneous T-cell lymphoma (CTCL).

P.C.N.CON

SWISS: Q9H2F9

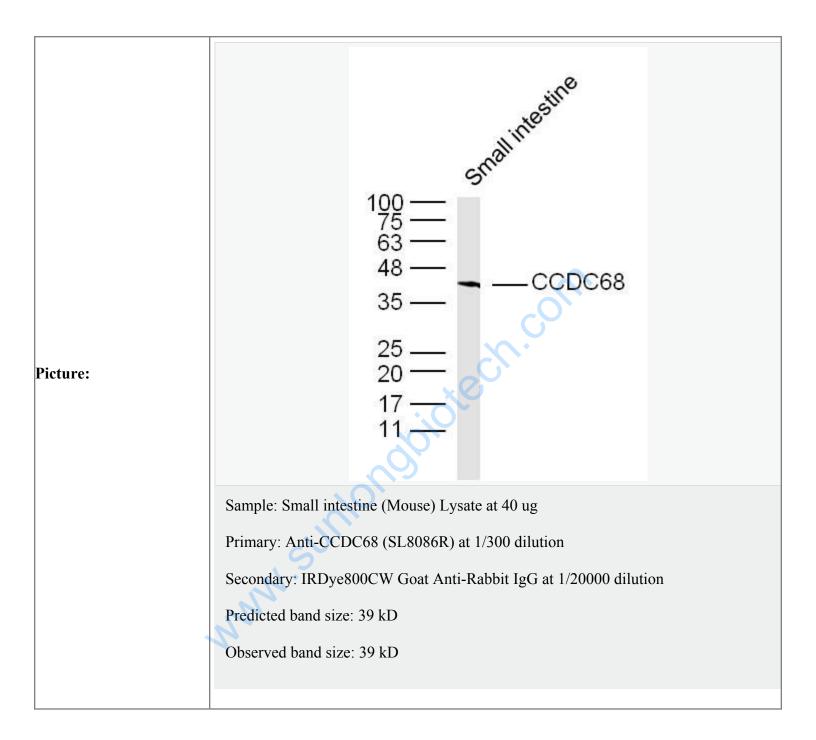
Gene ID: 80323

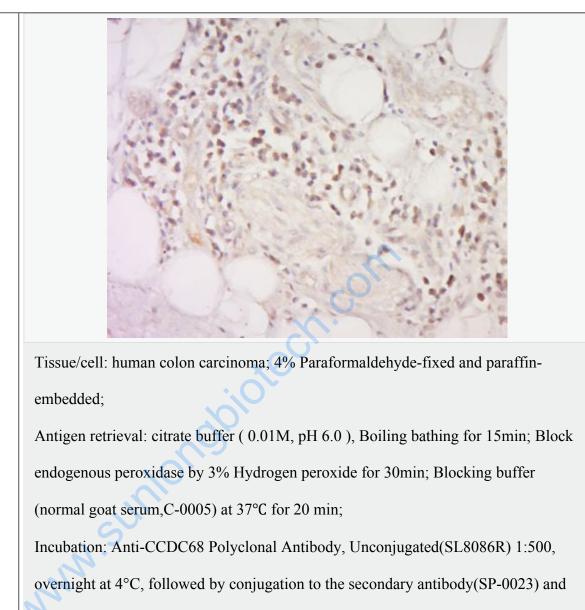
Database links: UniProtKB/Swiss-Prot: Q9H2F9.1

Important Note:

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

MMN.SUI





DAB(C-0010) staining

