

Rabbit Anti-ZEB1/NIL2A antibody

SL4187R

Product Name:	ZEB1/NIL2A
Chinese Name:	负调 控因子白细胞介素2抗体
Alias:	 AREB 6; AREB6; BZP; Delta crystallin enhancer binding factor 1; DELTA EF1; FECD6; MGC133261; Negative regulator of IL 2; Negative regulator of IL2; NIL 2 A; NIL 2 A zinc finger protein; NIL 2A; NIL-2-A zinc finger protein; NIL2A; Posterior polymorphous corneal dystrophy 3; PPCD3; Represses interleukin 2 expression; TCF 8; TCF-8; TCF8; Transcription factor 8 (represses interleukin 2 expression); Transcription factor 8; ZEB 1; ZEB;ZEB1_HUMAN; ZFHEP; ZFHX 1A; ZFHX1A; Zinc finger E box binding homeobox 1; Zinc finger E-box-binding homeobox 1; Zinc finger
	Specific References(2) SL4187R has been referenced in 2 publications.
	[IF=3.48]Sasaki, Takamitsu, et al. "Significance of epithelial growth factor in the
	epithelial?Cmesenchymal transition of human gallbladder cancer cells."Cancer Science
文献引用	(2012).0WB;Human.
Pub Med	PubMed:22404757
:	[IF=2.06]Xiang, Shuai, et al. "ZEB1 Expression Is Correlated With Tumor Metastasis
	and Reduced Prognosis of Breast Carcinoma in Asian Patients." Cancer Investigation
	(2015).other;
	PubMed:25950745
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human, Mouse, Rat, Chicken, Dog, Pig, Cow, Rabbit,
Applications:	WB=1:500-2000ELISA=1:500-1000IHC-P=1:400-800Flow-Cyt=1µg/Test(Paraffin sections need antigen repair) not yet tested in other applications. optimal dilutions/concentrations should be determined by the end user.

Molecular weight:	124kDa
Cellular localization:	The nucleus
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human ZEB1:211-320/1124
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
PubMed: Product Detail:	This gene encodes a zinc finger transcription factor. The encoded protein likely plays a role in transcriptional repression of interleukin 2. Mutations in this gene have been associated with posterior polymorphous corneal dystrophy-3 and late-onset Fuchs endothelial corneal dystrophy. Alternatively spliced transcript variants encoding different isoforms have been described.[provided by RefSeq, Mar 2010] Function: Acts as a transcriptional repressor. Inhibits interleukin-2 (IL-2) gene expression. Enhances or represses the promoter activity of the ATP1A1 gene depending on the quantity of cDNA and on the cell type. Represses E-cadherin promoter and induces an epithelial-mesenchymal transition (EMT) by recruiting SMARCA4/BRG1. Represses BCL6 transcription in the presence of the corepressor CTBP1. Positively regulates neuronal differentiation. Represses RCOR1 transcription activation during neurogenesis. Represses transcription by binding to the E box (5'-CANNTG-3'). Promotes tumorigenicity by repressing stemness-inhibiting microRNAs. Subunit: Interacts (via N-terminus) with SMARCA4/BRG1. Subcellular Location: Nucleus. Tissue Specificity: Colocalizes with SMARCA4/BRG1 in E-cadherin-negative cells from established lines, and stroma of normal colon as well as in de-differentiated epithelial cells at the invasion front of colorectal carcinomas (at protein level). Expressed in heart and skeletal muscle,
	 but not in liver, spleen, or pancreas. DISEASE: Corneal dystrophy, posterior polymorphous, 3 (PPCD3) [MIM:609141]: A subtype of posterior corneal dystrophy, a disease characterized by alterations of Descemet membrane presenting as vesicles, opacities or band-like lesions on slit-lamp examination and specular microscopy. Affected patient typically are asymptomatic. Note=The

disease is caused by mutations affecting the gene represented in this entry. Corneal dystrophy, Fuchs endothelial, 6 (FECD6) [MIM:613270]: A corneal disease caused by loss of endothelium of the central cornea. It is characterized by focal wart-like guttata that arise from Descemet membrane and develop in the central cornea, epithelial blisters, reduced vision and pain. Descemet membrane is thickened by abnormal collagenous deposition. Note=The disease is caused by mutations affecting the gene represented in this entry.

Similarity:

Belongs to the delta-EF1/ZFH-1 C2H2-type zinc-finger family. Contains 7 C2H2-type zinc fingers. Contains 1 homeobox DNA-binding domain. piotecn.com

SWISS: P37275

Gene ID: 6935

Database links:

Entrez Gene: 396029Chicken

Entrez Gene: 535183Cow

Entrez Gene: 6935Human

Entrez Gene: 21417 Mouse

Entrez Gene: 25705Rat

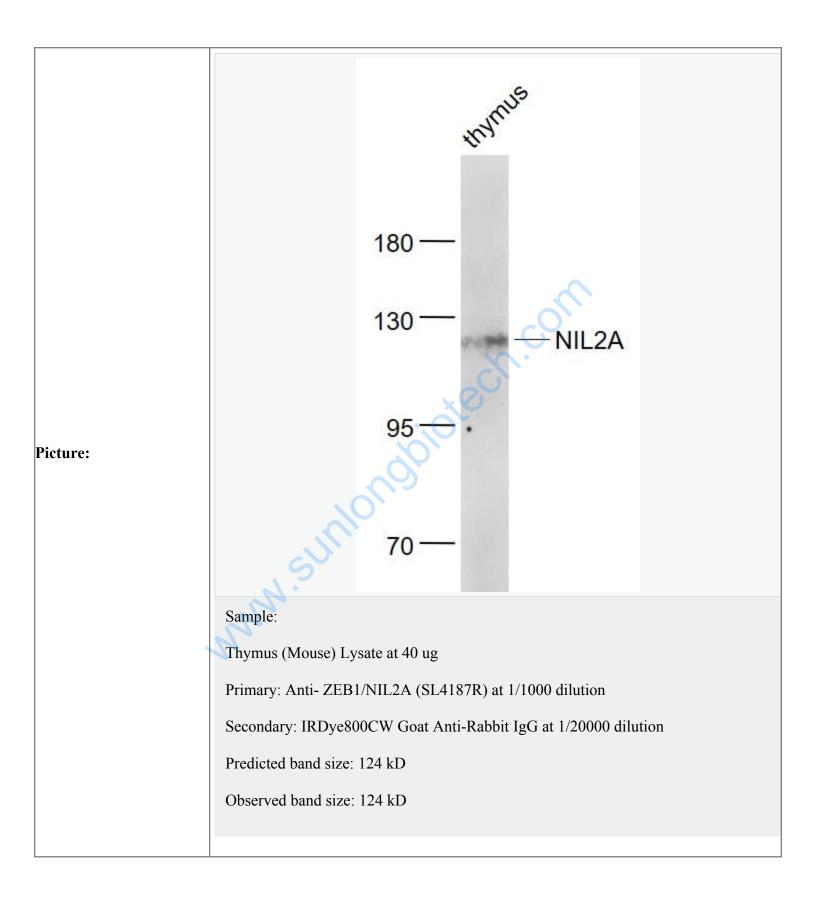
Omim: 189909Human

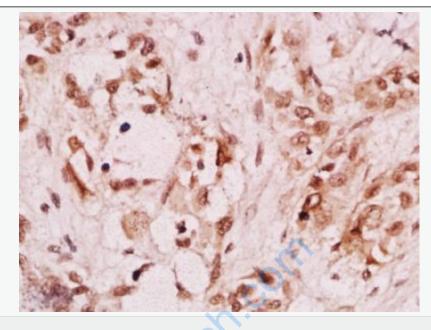
SwissProt: P36197Chicken

SwissProt: P37275Human

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 breast cancer); 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 (ZEB1) Polyclonal Antibody, Unconjugated (SL4187R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.

