

Anti-CTCF Picoband Antibody
Catalog # ABO12181**Specification****Anti-CTCF Picoband Antibody - Product Information**

Application	WB, IHC
Primary Accession	P49711
Host	Rabbit
Reactivity	Human, Mouse, Rat
Clonality	Polyclonal
Format	Lyophilized

Description

Rabbit IgG polyclonal antibody for Transcriptional repressor CTCF(CTCF) detection. Tested with WB, IHC-P in Human;Mouse;Rat.

Reconstitution

Add 0.2ml of distilled water will yield a concentration of 500ug/ml.

Anti-CTCF Picoband Antibody - Additional Information

Gene ID 10664

Other Names

Transcriptional repressor CTCF, 11-zinc finger protein, CCCTC-binding factor, CTCFL paralog, CTCF

Calculated MW

82785 MW KDa

Application Details

Immunohistochemistry(Paraffin-embedded Section), 0.5-1 µg/ml, Human, Mouse, Rat, By Heat
Western blot, 0.1-0.5 µg/ml, Human, Rat

Subcellular Localization

Nucleus, nucleoplasm. Chromosome. Chromosome, centromere. May translocate to the nucleolus upon cell differentiation. Associates with both centromeres and chromosomal arms during metaphase. Associates with the H19 ICR in mitotic chromosomes. May be preferentially excluded from heterochromatin during interphase.

Tissue Specificity

Ubiquitous. Absent in primary spermatocytes. .

Protein Name

Transcriptional repressor CTCF

Contents

Each vial contains 5mg BSA, 0.9mg NaCl, 0.2mg Na₂HPO₄, 0.05mg Na₃.

Immunogen

E.coli-derived human CTCF recombinant protein (Position: K521-R727). Human CTCF shares 94%

and 93.1% amino acid (aa) sequence identity with mouse and rat CTCF, respectively.

Purification

Immunogen affinity purified.

Cross Reactivity

No cross reactivity with other proteins

Storage

At -20°C for one year. After r°Constitution, at 4°C for one month. It°Can also be aliquotted and stored frozen at -20°C for a longer time.Avoid repeated freezing and thawing.

Sequence Similarities

Belongs to the CTCF zinc-finger protein family.

Anti-CTCF Picoband Antibody - Protein Information

Name CTCF

Function

Chromatin binding factor that binds to DNA sequence specific sites and regulates the 3D structure of chromatin (PubMed:18347100, PubMed:18654629, PubMed:19322193). Binds together strands of DNA, thus forming chromatin loops, and anchors DNA to cellular structures, such as the nuclear lamina (PubMed:18347100, PubMed:18654629, PubMed:19322193). Defines the boundaries between active and heterochromatic DNA via binding to chromatin insulators, thereby preventing interaction between promoter and nearby enhancers and silencers (PubMed:18347100, PubMed:18654629, PubMed:19322193). Plays a critical role in the epigenetic regulation (PubMed:16949368). Participates in the allele-specific gene expression at the imprinted IGF2/H19 gene locus (PubMed:16107875, PubMed:16815976, PubMed:17827499). On the maternal allele, binding within the H19 imprinting control region (ICR) mediates maternally inherited higher- order chromatin conformation to restrict enhancer access to IGF2 (By similarity). Mediates interchromosomal association between IGF2/H19 and WSB1/NF1 and may direct distant DNA segments to a common transcription factory (By similarity). Regulates asynchronous replication of IGF2/H19 (By similarity). Plays a critical role in gene silencing over considerable distances in the genome (By similarity). Preferentially interacts with unmethylated DNA, preventing spreading of CpG methylation and maintaining methylation-free zones (PubMed:18413740). Inversely, binding to target sites is prevented by CpG methylation (PubMed:18413740). Plays an important role in chromatin remodeling (PubMed:18413740). Can dimerize when it is bound to different DNA sequences, mediating long-range chromatin looping (PubMed:12191639). Causes local

loss of histone acetylation and gain of histone methylation in the beta-globin locus, without affecting transcription (PubMed:[12191639](http://www.uniprot.org/citations/12191639)). When bound to chromatin, it provides an anchor point for nucleosomes positioning (PubMed:[12191639](http://www.uniprot.org/citations/12191639)). Seems to be essential for homologous X-chromosome pairing (By similarity). May participate with Tsix in establishing a regulatable epigenetic switch for X chromosome inactivation (PubMed:[11743158](http://www.uniprot.org/citations/11743158)). May play a role in preventing the propagation of stable methylation at the escape genes from X-inactivation (PubMed:[11743158](http://www.uniprot.org/citations/11743158)). Involved in sister chromatid cohesion (PubMed:[12191639](http://www.uniprot.org/citations/12191639)). Associates with both centromeres and chromosomal arms during metaphase and required for cohesin localization to CTCF sites (PubMed:[18550811](http://www.uniprot.org/citations/18550811)). Plays a role in the recruitment of CENPE to the pericentromeric/centromeric regions of the chromosome during mitosis (PubMed:[26321640](http://www.uniprot.org/citations/26321640)). Acts as a transcriptional repressor binding to promoters of vertebrate MYC gene and BAG1 gene (PubMed:[18413740](http://www.uniprot.org/citations/18413740)), PubMed:[8649389](http://www.uniprot.org/citations/8649389), PubMed:[9591631](http://www.uniprot.org/citations/9591631)). Also binds to the PLK and PIM1 promoters (PubMed:[12191639](http://www.uniprot.org/citations/12191639)). Acts as a transcriptional activator of APP (PubMed:[9407128](http://www.uniprot.org/citations/9407128)). Regulates APOA1/C3/A4/A5 gene cluster and controls MHC class II gene expression (PubMed:[18347100](http://www.uniprot.org/citations/18347100), PubMed:[19322193](http://www.uniprot.org/citations/19322193)). Plays an essential role in oocyte and preimplantation embryo development by activating or repressing transcription (By similarity). Seems to act as tumor suppressor (PubMed:[12191639](http://www.uniprot.org/citations/12191639)).

Cellular Location

Nucleus, nucleoplasm. Chromosome. Chromosome, centromere. Note=May translocate to the nucleolus upon cell differentiation. Associates with both centromeres and chromosomal arms during metaphase. Associates with the H19 ICR in mitotic chromosomes. May be preferentially excluded from heterochromatin during interphase

Tissue Location

Ubiquitous. Absent in primary spermatocytes.

Anti-CTCF Picoband Antibody - Protocols

Provided below are standard protocols that you may find useful for product applications.

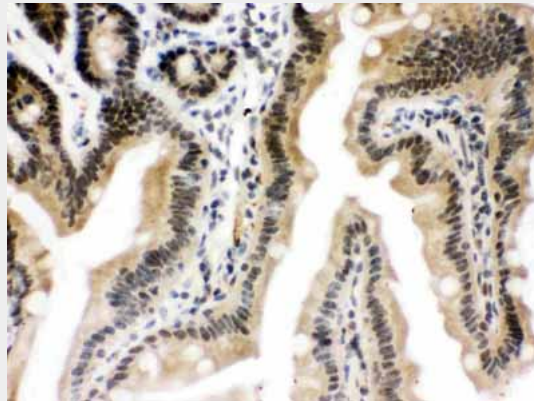
- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

Anti-CTCF Picoband Antibody - Images

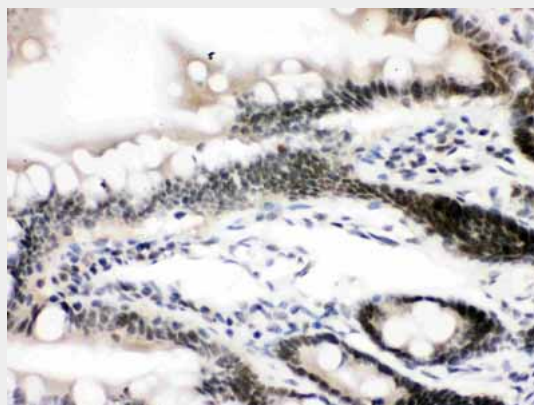




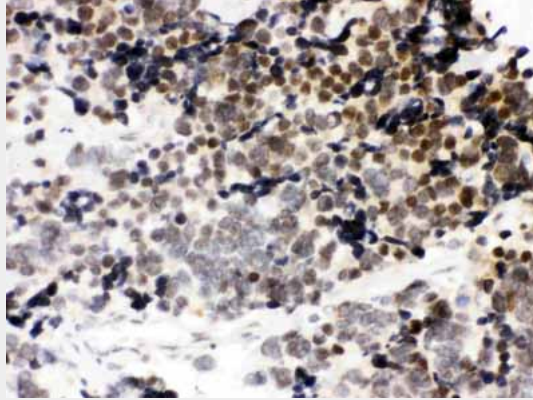
Anti-CTCF Picoband antibody, ABO12181, Western blotting All lanes: Anti CTCF (ABO12181) at 0.5ug/ml Lane 1: Rat Thymus Tissue Lysate at 50ug Lane 2: Rat Gaster Tissue Lysate at 50ug Lane 3: Human Placenta Tissue Lysate at 50ug Predicted bind size: 88KD Observed bind size: 88KD



Anti-CTCF Picoband antibody, ABO12181, IHC(P) IHC(P): Mouse Intestine Tissue



Anti-CTCF Picoband antibody, ABO12181, IHC(P) IHC(P): Rat Intestine Tissue



Anti-CTCF Picoband antibody, ABO12181, IHC(P)IHC(P): Human Lung Cancer Tissue

Anti-CTCF Picoband Antibody - Background

Transcriptional repressor CTCF also known as 11-zinc finger protein or CCCTC-binding factor is a transcription factor that in humans is encoded by the CTCF gene. This gene is a member of the BORIS + CTCF gene family and encodes a transcriptional regulator protein with 11 highly conserved zinc finger (ZF) domains. And this nuclear protein is able to use different combinations of the ZF domains to bind different DNA target sequences and proteins. Depending upon the context of the site, the protein can bind a histone acetyltransferase (HAT)-containing complex and function as a transcriptional activator or bind a histone deacetylase (HDAC)-containing complex and function as a transcriptional repressor. If the protein is bound to a transcriptional insulator element, it can block communication between enhancers and upstream promoters, thereby regulating imprinted expression. Mutations in this gene have been associated with invasive breast cancers, prostate cancers, and Wilms' tumors. Alternatively spliced transcript variants encoding different isoforms have been found for this gene.