

Anti-Human ZNF410, monoclonal (clone R585.2.2C9)

Recommended name: Zinc finger protein 410

Alternative name(s): another partner for ARF 1; zinc finger protein APA-1; clones 23667 and 23775 zinc finger protein

Cat. No. m14-038
Lot. No. 20150825.IJVR

Quantity: 100 µg
Storage: -20 °C



FOR RESEARCH USE ONLY

NOT FOR USE IN HUMANS

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Uniprot / NCBI Summary

UniProt

Primary Accession # [Q53FM1](#)
Secondary Accession # [Q53FM1](#); [Q86VK4](#)

NCBI

GI # [10863995](#)
GenID [57862](#)
Accession # [NP_067011.1](#)
GenBank Nucleotide # [NM_021188.2](#)

Molecular Weight 52,086 Da (478 aa)

Transcription factor that activates transcription of matrix-remodeling genes such as MMP1 during fibroblast senescence.

Subcellular location: nucleus

General Reference:

Benanti JA, Williams DK, Robinson KL, Ozer HL, Galloway DA (2002) Induction of extracellular matrix-remodeling genes by the senescence-associated protein APA-1. *Mol Cell Biol* **22**:7385-97. [[PubMed](#)]

Physical Characteristics

Quantity: 100 µg

Concentration: 1.0 mg/ml

Host / Isotype: mouse IgG2b

Clonality: monoclonal; ID R585.2.2C9

Immunogen: recombinant protein corresponding to aa residues 129-478 of human ZNF410

Purification: affinity-chromatography using Protein G

Formulation: 30% glycerol, 1x PBS, 0.02% sodium azide

Specificity: monospecific for human ZNF410; see microarray analysis below

Reactivity: human

Stability/Storage: 12 months long term: -20 °C; short term: 4 °C; avoid freeze-thaw cycles; aliquot as required

Handling Notes: small volumes of antibody may occasionally become entrapped in the seal of the product vial during shipment and storage; if necessary, briefly centrifuge the vial on a tabletop centrifuge to dislodge any liquid in the container cap.

Tested Research Applications

Immunoprecipitation: recommended; see below.

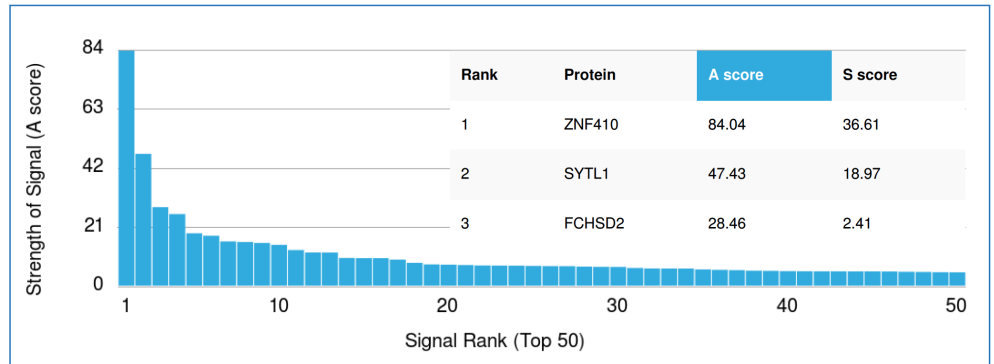
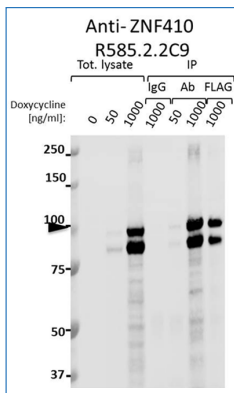
ChIP-Seq: recommended; see page 2

Western Blot: tested on cells transfected with a construct encoding ZNF410; utility on native cells under evaluation

Quality Assurance

IP Analysis:

Tet-ON HeLa cells were transfected with construct encoding ZNF410 (Q86VK4) with an N-terminal fusion of FLAG, YFP (Venus) and V5 tags under a tet-inducible promoter. These cells were stimulated with 0, 50 or 1000 ng/ml doxycycline. Immunoprecipitation (IP) was carried out using 5 µg of either IgG, CDI mAb Anti ZNF410 (cloneID# R585.2.2C9) or 1 µg of FLAG-M2 (Sigma). Immunoblotting was performed using rabbit Anti-FLAG (1:1000, Cell Signaling #2368).



Specificity Analysis with HuProt™ Human Proteome Microarray: Anti Human ZNF410 (clone R585.2.2C9) was analyzed using the CDI HuProt™ Human Proteome Microarray.

For more information on A/S scores and how they relate to specificity, see page 2.

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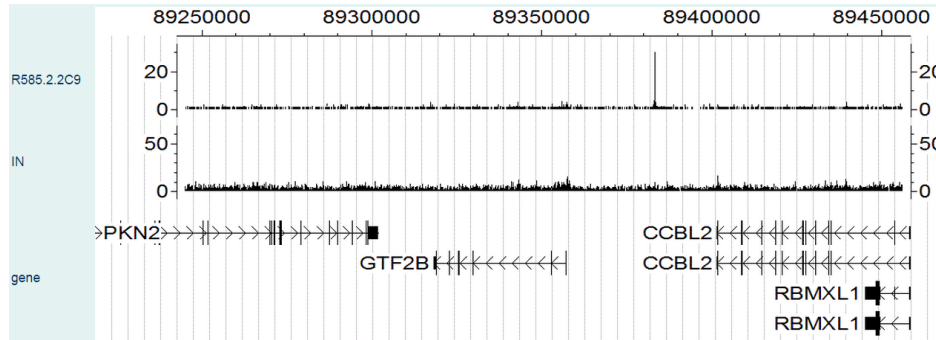
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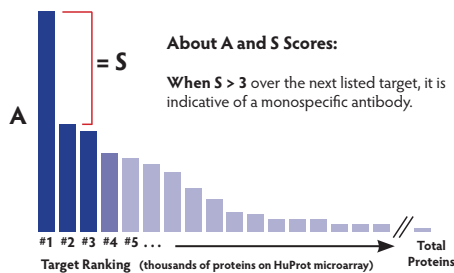
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Tested Research Applications

ChIP-Seq: Recommended



The ChIP was performed with chromatin from 10 million HCT116 cells or HeLa cells and 3 µg of Anti-ZXDC (cloneID # JH98.1.2B5) antibody. The ChIP DNA was sequenced on an Illumina HiSeq platform and read counts were calculated at consecutive 100 bp bins across the human genome hg19. Normalized read-count levels for ChIP-seq of ZXDC (JH98.1.2B5) and control (IN) around the MPRIP loci are displayed in the CisGenome browser.



Statistical Analysis: Thousands of GenePix data points (from the microarray) are analyzed in terms of signal strength and ranked accordingly.

SUMMARY: The A-score indicates the number of standard deviations above background seen for the mean signal bound by the target antigen. The S-score represents the difference between the A-score of the target antigen and the next best hit on the array. S-scores **greater than 3 standard deviations over the next listed target** are deemed statistically significant and indicate **highly specific antibodies**. More info at cdi-lab.com/HighSpec.html

The development of this antibody was supported by the National Institutes of Health Protein Capture Reagent Program under award U54HG06434 to CDI Laboratories and Johns Hopkins University.

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