

RAI17/ZMIZ1 (E2X3X) Rabbit mAb

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Applications:	Reactivity:	Sensitivity:	MW (kDa):	Source/Isotype:	UniProt ID:	Entrez-Gene Id:
W, IP, IHC-P	H M Mk	Endogenous	120, 130	Rabbit IgG	#Q9ULJ6	57178

Product Usage Information**Application**

Western Blotting
Immunoprecipitation
Immunohistochemistry (Paraffin)

Dilution

1:1000
1:50
1:100 - 1:400

Storage

Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at -20°C. *Do not aliquot the antibody.*

Specificity/Sensitivity

RAI17/ZMIZ1 (E2X3X) Rabbit mAb recognizes endogenous levels of total RAI17/ZMIZ1 protein.

Source / Purification

Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Pro902 of human RAI17/ZMIZ1 protein.

Background

ZMIZ1, also known as ZIMP10 or RAI17, is a member of the Protein Inhibitor of Activated STAT (PIAS)-like family of transcriptional coregulators (1). ZMIZ1 was initially discovered as a novel coregulator of the androgen receptor (AR) and was shown to augment AR activity by enhancing sumoylation of the receptor (2). Subsequent studies have shown that ZMIZ1 can also act as a transcriptional coactivator of p53 (3) and SMAD3 (4). During thymopoiesis, ZMIZ1 was shown to promote pre-T-cell proliferation through cooperative induction of a subset of NOTCH target genes (5). In T-cell acute lymphoblastic leukemia (T-ALL) cells, ZMIZ1 was shown to engage directly with NOTCH1 through an N-terminal tetratricopeptide repeat (TPR) domain, facilitating its recruitment to a long-range acting enhancer element to promote *MYC* transcription and activity. Moreover, inhibition of ZMIZ1 impaired the initiation and maintenance of NOTCH-induced T-ALL (6). Although NOTCH1 appears to rely on ZMIZ1 to selectively amplify an oncogenic subset of target genes, research studies suggest ZMIZ1 is not involved in NOTCH-mediated intestinal homeostasis or myeloid suppression, suggesting that therapeutic strategies targeting ZMIZ1 may provide an opportunity to indirectly target NOTCH-driven cancers, with reduced adverse effects (7).

Background References

1. Shuai, K. and Liu, B. (2005) *Nat Rev Immunol* 5, 593-605.
2. Sharma, M. et al. (2003) *EMBO J* 22, 6101-14.
3. Lee, J. et al. (2007) *Nucleic Acids Res* 35, 4523-34.
4. Li, X. et al. (2006) *J Biol Chem* 281, 23748-56.
5. Wang, Q. et al. (2018) *Blood* 132, 1279-92.
6. Rakowski, L.A. et al. (2013) *Cancer Res* 73, 930-41.
7. Pinnell, N. et al. (2015) *Immunity* 43, 870-83.

Species Reactivity

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot Buffer

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key

W: Western Blotting **IP:** Immunoprecipitation **IHC-P:** Immunohistochemistry (Paraffin)

Cross-Reactivity Key

H: Human **M:** Mouse **Mk:** Monkey

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