Revision 4

Store at -20C

#4334

AML1 Antibody

Cell Signaling TECHNOLOGY®

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For Research Use Only. Not for Use in Diagnostic Procedures.

Applications:	Reactivity:	Sensitivity:	MW (kDa):	Source/Isotype:	UniProt ID:	Entrez-Gene Id:
W, IF-IC, FC-FP	H Mk	Endogenous	55	Rabbit	#Q01196	861

Product Usage Information

Application	Dilution
Western Blotting	1:1000
Immunofluorescence (Immunocytochemistry)	1:50
Flow Cytometry (Fixed/Permeabilized)	1:200

Storage

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

Specificity/Sensitivity

AML1 antibody detects endogenous levels of total AML1 protein and the AML1/ETO fusion protein.

Source / Purification

Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ala36 of human AML1 (RUNX1) protein. Antibodies are purified by protein A and peptide affinity chromatography.

Background

AML1 (also known as Runx1, CBFA2, and PEBP2αB) is a member of the core binding factor (CBF) family of transcription factors (1,2). It is required for normal development of all hematopoietic lineages (3-5). AML1 forms a heterodimeric DNA binding complex with its partner protein CBFβ and regulates the expression of cellular genes by binding to promoter and enhancer elements. AML1 is commonly translocated in hematopoietic cancers: chromosomal translocations include t(8;21) AML1-ETO, t(12;21) TEL-AML, and t(8;21) AML-M2 (6). Phosphorylation of AML1 on several potential serine and threonine sites, including Ser249, is thought to occur in an Erk-dependent manner (7,8).

Background References

1. Wang, S. et al. (1993) *Mol Cell Biol* 13, 3324-39.
2. Ogawa, E. et al. (1993) *Proc Natl Acad Sci U S A* 90, 6859-63.
3. Okuda, T. et al. (1996) *Cell* 84, 321-30.
4. Wang, Q. et al. (1996) *Proc Natl Acad Sci U S A* 93, 3444-9.
5. North, T.E. et al. (2004) *Stem Cells* 22, 158-68.
6. Blyth, K. et al. (2005) *Nat Rev Cancer* 5, 376-87.
7. Tanaka, T. et al. (1996) *Mol Cell Biol* 16, 3967-79.
8. Zhang, Y. et al. (2004) *J Biol Chem* 279, 53116-25.

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 **IF-IC:** Immunofluorescence (Immunocytochemistry) **FC-FP:** Flow Cytometry (Fixed/Permeabilized)

Cross-Reactivity Key

H: Human **Mk:** Monkey

Trademarks and Patents

Limited Uses

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