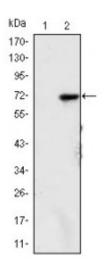


Anti-JMJD2A antibody



Description

Mouse monoclonal to JMJD2A.

Model STJ98191

Host Mouse

Reactivity Human

Applications ELISA, IF, IHC, WB

Immunogen Purified recombinant fragment of human JMJD2A expressed in E. Coli.

Gene ID 9682

Gene Symbol KDM4A

Dilution range WB 1:500-1:2000IHC 1:200-1:1000IF 1:200-1:1000ELISA 1:10000

Specificity JMJD2A Monoclonal Antibody detects endogenous levels of JMJD2A

protein.

Tissue Specificity Ubiquitous.

Purification Affinity purification

Clone ID 5H1

Note For Research Use Only (RUO).

Protein Name Lysine-specific demethylase 4A JmjC domain-containing histone

demethylation protein 3A Jumonji domain-containing protein 2A

Clonality Monoclonal

Conjugation Unconjugated

Isotype IgG1

Formulation Ascitic fluid containing 0.03% sodium azide.

Storage Instruction Store at -20°C, and avoid repeat freeze-thaw cycles.

Database Links HGNC:22978OMIM:609764

Alternative Names Lysine-specific demethylase 4A JmjC domain-containing histone

demethylation protein 3A Jumonji domain-containing protein 2A

Function Histone demethylase that specifically demethylates 'Lys-9' and 'Lys-36'

residues of histone H3, thereby playing a central role in histone code. Does

not demethylate histone H3 'Lys-4', H3 'Lys-27' nor H4 'Lys-20'.

Demethylates trimethylated H3 'Lys-9' and H3 'Lys-36' residue, while it has no activity on mono- and dimethylated residues. Demethylation of Lys residue

generates formaldehyde and succinate. Participates in transcriptional

repression of ASCL2 and E2F-responsive promoters via the recruitment of histone deacetylases and NCOR1, respectively.; Isoform 2: Crucial for muscle differentiation, promotes transcriptional activation of the Myog gene by

directing the removal of repressive chromatin marks at its promoter. Lacks the

N-terminal demethylase domain.

Sequence and Domain Family The 2 Tudor domains recognize and bind methylated histone H3 'Lys-4'

residue (H3K4me). Double Tudor domain has an interdigitated structure and the unusual fold is required for its ability to bind methylated histone tails. Trimethylated H3 'Lys-4' (H3K4me3) is bound in a cage of 3 aromatic residues, 2 of which are from the Tudor domain 2, while the binding specificity is determined by side-chain interactions involving residues from the Tudor domain 1. The Tudor domains are also able to bind trimethylated histone H3 'Lys-9' (H3K9me3), di- and trimethylated H4 'Lys-20' (H4K20me2 and H4K20me3). Has high affinity for H4K20me2, blocking recruitment of

proteins such as TP53BP1.

Cellular Localization Nucleus

Post-translational Modifications

Ubiquitinated by RNF8 and RNF168 following DNA damage, leading to its degradation. Degradation promotes accessibility of H4K20me2 mark for DNA

repair protein TP53BP1, which is then recruited.

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