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# anti-KDM4C antibody (C-Term)

3 Images



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| Overview             |  |  |
|----------------------|--|--|
| Quantity:            | 400 μL   |  |
| Target:              | KDM4C  |  |
| Binding Specificity: | AA 1023-1056, C-Term   |  |
| Reactivity:          | Human, Mouse   |  |
| Host:                | Rabbit   |  |
| Clonality:           | Polyclonal   |  |
| Conjugate:           | This KDM4C antibody is un-conjugated   |  |
| Application:         | Western Blotting (WB), Flow Cytometry (FACS)   |  |
| Product Details      |  |  |
| Immunogen:           | This JMJD2C antibody is generated from rabbits immunized with a KLH conjugated synthetic         |  |
|                      | peptide between 1023-1056 amino acids from the C-terminal region of human JMJD2C.                |  |
| Clone:               | RB10796  |  |
| Isotype:             | lg Fraction  |  |
| Purification:        | This antibody is purified through a protein A column, followed by peptide affinity purification. |  |
| T                    |  |  |
| Target Details       |  |  |
| Target:              | KDM4C  |  |
| Alternative Name:    | JMJD2C (KDM4C Products)  |  |
| Background:          | This gene is a member of the Jumonji domain 2 (JMJD2) family and encodes a protein with          |  |

### **Target Details**

one JmjC domain, one JmjN domain, two PHD-type zinc fingers, and two Tudor domains. This nuclear protein functions as a trimethylation-specific demethylase, converting specific trimethylated histone residues to the dimethylated form. Chromosomal aberrations and increased transcriptional expression of this gene are associated with esophageal squamous cell carcinoma. Alternative splicing results in multiple transcript variants.

Molecular Weight: 119982

Gene ID: 23081

NCBI Accession: NP\_001140166, NP\_001140167, NP\_001140168, NP\_055876

UniProt: Q9H3R0

Nuclear Hormone Receptor Binding, Warburg Effect

# **Application Details**

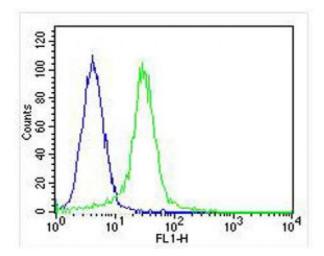
Application Notes: WB: 1:2000. FC: 1:25. FC: 1:25

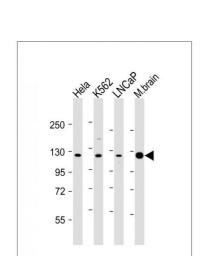
Restrictions: For Research Use only

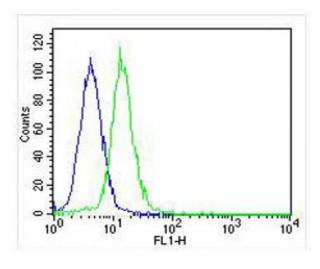
## Handling

Pathways:

| Format:            | Liquid   |  |
|--------------------|--|--|
| Buffer:            | Purified polyclonal antibody supplied in PBS with 0.09 % (W/V) sodium azide.   |  |
| Preservative:      | Sodium azide   |  |
| Precaution of Use: | This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.                     |  |
| Storage:           | 4 °C,-20 °C  |  |
| Storage Comment:   | Maintain refrigerated at 2-8 °C for up to 6 months. For long term storage store at -20 °C in small aliquots to prevent freeze-thaw cycles. |  |
| Expiry Date:       | 6 months   |  |







#### **Flow Cytometry**

**Image 1.** Overlay histogram showing Hela cells stained with (ABIN655391 and ABIN2844939) (green line). The cells were fixed with 2% paraformaldehyde (10 min) and then permeabilized with 90% methanol for 10 min. The cells were then icubated in 2% bovine serum albumin to block non-specific protein-protein interactions followed by the antibody ((ABIN655391 and ABIN2844939), 1:25 dilution) for 60 min at 37 °C. The secondary antibody used was Goat-Anti-Rabbit IgG, DyLight® 488 Conjugated Highly Cross-Adsorbed(OH191631) at 1/400 dilution for 40 min at 37 °C. Isotype control antibody (blue line) was rabbit IgG (1  $\mu$  g/1x10^6 cells) used under the same conditions. Acquisition of >10,000 events was performed.

#### **Western Blotting**

Image 2. All lanes: Anti-JMJD2C Antibody (C-term) at 1:2000 dilution Lane 1: Hela whole cell lysates Lane 2: K562 whole cell lysates Lane 3: LNCaP whole cell lysates Lane 4: mouse brain lysates Lysates/proteins at 20 μg per lane. Secondary Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/10000 dilution. Predicted band size: 120 kDa Blocking/Dilution buffer: 5 % NFDM/TBST.

#### **Flow Cytometry**

Image 3. Overlay histogram showing Hela cells stained with (ABIN655391 and ABIN2844939) (green line). The cells were fixed with 2% paraformaldehyde (10 min) and then permeabilized with 90% methanol for 10 min. The cells were then icubated in 2% bovine serum albumin to block non-specific protein-protein interactions followed by the antibody ((ABIN655391 and ABIN2844939), 1:25 dilution) for 60 min at 37 °C. The secondary antibody used was Goat-Anti-Rabbit IgG, DyLight® 488 Conjugated Highly Cross-

Adsorbed(NA168821)) at 1/400 dilution for 40 min at 37 °C. Isotype control antibody (blue line) was mouse IgG1 (1  $\mu$  g/1x10^6 cells) used under the same conditions. Acquisition of >10, 000 events was performed.