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## [KD Validated] METTL3 Rabbit mAb

Catalog No.: A26858 Recombinant

## **Basic Information**

#### **Observed MW**

75kDa

#### **Calculated MW**

64kDa

## **Category**

SMab Recombinant Monoclonal Antibody

## **Applications**

WB,IHC-P,IP,ChIP,ELISA

## **Cross-Reactivity**

Human, Mouse, Rat

## **Background**

This gene encodes the 70 kDa subunit of MT-A which is part of N6-adenosine-methyltransferase. This enzyme is involved in the posttranscriptional methylation of internal adenosine residues in eukaryotic mRNAs, forming N6-methyladenosine.

## **Recommended Dilutions**

WB 1:5000 - 1:10000

**IHC-P** 1:200 - 1:2000

**IP** 0.5μg-4μg antibody for

200µg-400µg extracts

of whole cells

ChIP 5µg antibody for

10μg-15μg of Chromatin

**ELISA** Recommended starting

concentration is 1 µg/mL. Please optimize the concentration based on your specific assay requirements.

## **Contact**

www.abclonal.com

## **Immunogen Information**

**Gene ID**Swiss Prot
Q86U44

## **Immunogen**

Recombinant protein (or fragment). This information is considered to be commercially sensitive.

## **Synonyms**

M6A; IME4; Spo8; MT-A70; hMETTL3

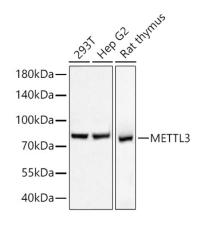
## **Product Information**

SourceIsotypePurificationRabbitIgGAffinity purification

#### Storage

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.09% Sodium azide, 0.05% BSA, 50% glycerol, pH7.3.



Western blot analysis of various lysates using [KD Validated] METTL3 Rabbit mAb (A26858) at 1:5000 dilution incubated overnight at 4°C.

Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000

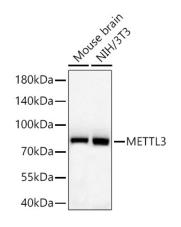
dilution.

Lysates/proteins: 25 µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 1s.



Western blot analysis of various lysates using [KD Validated] METTL3 Rabbit mAb (A26858) at 1:5000 dilution incubated overnight at 4°C.

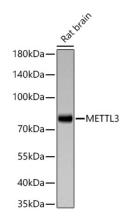
Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25 µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 10s.



Western blot analysis of lysates from Rat brain using [KD Validated] METTL3 Rabbit mAb (A26858) at 1:1000 dilution incubated overnight at 4°C.

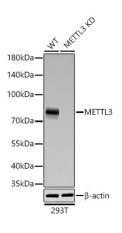
Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution

Lysates/proteins: 25 µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 60s.



Western blot analysis of lysates from wild type (WT) and METTL3 knockdown (KD) 293T cells using [KD Validated] METTL3 Rabbit mAb (A26858) at 1:1000 dilution incubated overnight at  $4^{\circ}$ C.

Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000

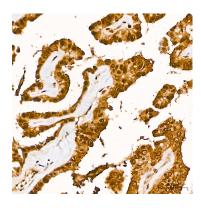
dilution.

Lysates/proteins: 25 µg per lane.

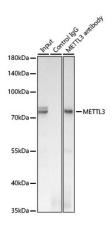
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 15s.



Immunohistochemistry analysis of paraffin-embedded Human thyroid cancer tissue using [KD Validated] METTL3 Rabbit mAb (A26858) at a dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate Buffer (pH 6.0) prior to IHC staining.



Immunoprecipitation of METTL3 from 300  $\mu$ g extracts of HepG2 cells was performed using 0.5  $\mu$ g of [KD Validated] METTL3 Rabbit mAb (A26858). Rabbit IgG isotype control (AC042) was used to precipitate the Control IgG sample. IP samples were eluted with 1X Laemmli Buffer. The Input lane represents 10% of the total input. Western blot analysis of immunoprecipitates was conducted using [KD Validated] METTL3 Rabbit mAb (A26858) at a dilution of 1 : 1000.