# [KD Validated] METTL3 Rabbit mAb

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Catalog No.: A26858 Recombinant

## **Basic Information**

## **Observed MW**

75kDa

## **Calculated MW**

64kDa

## Category

Primary antibody

## **Applications**

WB,IHC-P,IP,CHIP,ELISA

#### **Cross-Reactivity**

Human, Mouse, Rat

#### CloneNo number

ARC66900

# **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

200μg-400μg extracts of

whole cells

**ELISA** Recommended starting

concentration is 1 µg/mL.

Please optimize the
concentration based on
your specific assay
requirements.

# **Immunogen Information**

**Gene ID Swiss Prot** 56339 Q86U44

#### **Immunogen**

Recombinant fusion protein containing a sequence corresponding to amino acids 1-156 of human METTL3 (NP\_062826.2).

## **Synonyms**

M6A; IME4; Spo8; MT-A70; hMETTL3

## **Contact**

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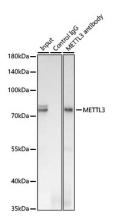
## **Product Information**

SourceIsotypePurificationRabbitIgGAffinity purification

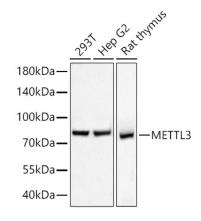
#### Storage

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

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



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.



Western blot analysis of various lysates using [KD Validated] METTL3 Rabbit mAb (A26858) at 1:5000 dilution incubated overnight at  $4^{\circ}$ C.

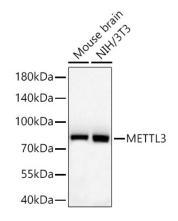
Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25 μg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 1s.



Western blot analysis of various lysates using [KD Validated] METTL3 Rabbit mAb (A26858) at 1:5000 dilution incubated overnight at  $4^{\circ}$ C.

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

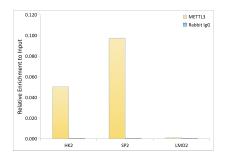
Lysates/proteins: 25 µg per lane.

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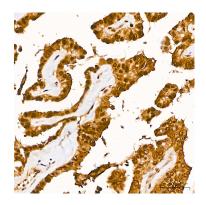
Detection: ECL Basic Kit (RM00020).

Exposure time: 10s.

# **Validation Data**



Chromatin immunoprecipitation was performed with 15  $\mu$ g of cross-linked chromatin from MCF7, using 3  $\mu$ g of [KD Validated] METTL3 Rabbit mAb (A26858) and Rabbit IgG isotype control (AC042). The enrichment of immunoprecipitated DNA at different genomic loci was examined by quantitative PCR. The histogram compares the ratio of the immunoprecipitated DNA to the input at given loci.



Immunohistochemistry analysis of paraffinembedded 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.