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SETD2 Rabbit pAb

Catalog No.: A11757

Basic Information

Observed MW

288kDa

Calculated MW

175kDa/192kDa/287kDa

Category

Primary antibody

Applications

WB

Cross-Reactivity

Human, Mouse, Rat

Background

Huntington's disease (HD), a neurodegenerative disorder characterized by loss of striatal neurons, is caused by an expansion of a polyglutamine tract in the HD protein huntingtin. This gene encodes a protein belonging to a class of huntingtin interacting proteins characterized by WW motifs. This protein is a histone methyltransferase that is specific for lysine-36 of histone H3, and methylation of this residue is associated with active chromatin. This protein also contains a novel transcriptional activation domain and has been found associated with hyperphosphorylated RNA polymerase II.

Recommended Dilutions

WB

1:500 - 1:2000

Immunogen Information

Gene ID 29072

Swiss Prot

Q9BYW2

Immunogen

Recombinant fusion protein containing a sequence corresponding to amino acids 803-1103 of human SETD2 (NP_054878.5).

Synonyms

SETD2;HBP231;HIF-1;HIP-1;HSPC069;HYPB;KMT3A;LLS;SET2;p231HBP

Contact

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Product Information

Source

Isotype

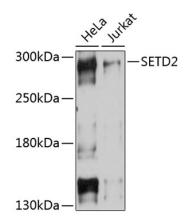
Purification

Rabbit IgG Affinity purification

Storage

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

Buffer: PBS with 0.01% thiomersal,50% glycerol,pH7.3.



Western blot analysis of extracts of various cell lines, using SETD2 Antibody (A11757) at 1:1000

dilution

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

Lysates/proteins: 25ug per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 15s.