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Catalog No.: A11757 1 Publications



Basic Information

Observed MW 288kDa

Calculated MW 288kDa

Category Primary antibody

Applications ELISA,WB

Cross-Reactivity Human

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

1:500 - 1:2000

Immunogen Information

WB

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

LLS; HYPB; SET2; HIF-1; HIP-1; KMT3A; MRD70; RAPAS; HBP231; HSPC069; p231HBP; SETD2

Product Information

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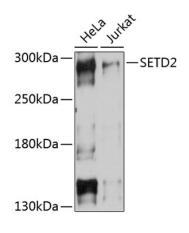
Source

Isotype IgG Purification Affinity purification

Storage

Rabbit

Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.01% thimerosal,50% glycerol,pH7.3.



Western blot analysis of various lysates using SETD2 Rabbit pAb (A11757) at 1:1000 dilution. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution. Lysates/proteins: 25µg per lane. Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit (RM00020). Exposure time: 15s.