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1-866-9 KINASE (54-6273)

Catalog #	Aliquot Size
S351-380G-05	5 µg
S351-380G-10	10 µg

SUV420H1 (KMT5B), Active

Full length recombinant protein expressed in Sf9 cells

Catalog # S351-380G
Lot # P1665-5

Product Description

Recombinant full-length human SUV420H1 (KMT5B) was expressed by baculovirus in Sf9 insect cells using an N-terminal GST tag. The SUV39H2 protein accession number is BC103498.

Gene Aliases

KMT5B, CGI-85, MGC21161, MGC703

Formulation

Recombinant protein stored in 50mM Tris-HCl, pH 7.5, 150mM NaCl, 10mM glutathione, 0.1mM EDTA, 0.25mM DTT, 0.1mM PMSF, 25% glycerol.

Storage and Stability

Store product at –70°C. For optimal storage, aliquot target into smaller quantities after centrifugation and store at recommended temperature. For most favorable performance, avoid repeated handling and multiple freeze/thaw cycles.

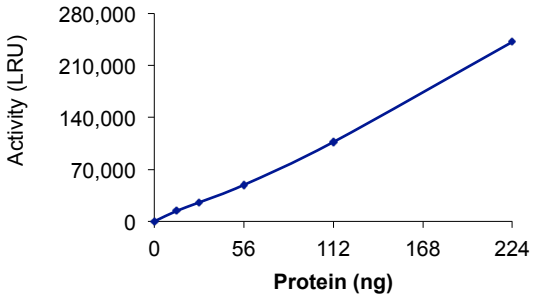
Scientific Background

Histone-lysine N-methyltransferase, Suppressor of variegation 4-20 homolog 1 SUV420H1 specifically trimethylates 'Lys-20' of histone H4, which in turn acts as a specific tag for epigenetic transcriptional repression. Located in pericentric heterochromatin regions, and plays a central role in the establishment of constitutive heterochromatin in these regions. Interacts with a family of proteins that display similarity with dual-specificity phosphatases (dsPTPases) mediated by its SET domains.

References

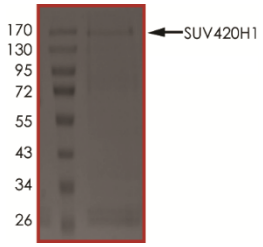
1. Tryndyak VP, et al: Loss of DNA methylation and histone H4 lysine 20 trimethylation in human breast cancer cells is associated with aberrant expression of DNA methyl-transferase 1, Suv4-20h2 histone methyltransferase and methyl-binding proteins. Cancer Biol Ther. 5(1):65-70, 2006.
2. Lai CH, et al: Identification of novel human genes evolutionarily conserved in Caenorhabditis elegans by comparative proteomics. Genome Res. 10(5):703-13, 2000.

Specific Activity



The specific activity of SUV420H1 (KMT5B) was determined to be **300 pmol /min/mg** as per activity assay protocol.

Purity



The purity of SUV420H1 (KMT5B) was determined to be **>70%** by densitometry, approx. MW **160 kDa**.

SUV420H1 (KMT5B), Active

Full length recombinant protein expressed in Sf9 cells

Catalog #	S351-380G
Specific Activity	300 pmol/min/mg
Lot #	P1665-5
Purity	>70%
Concentration	0.05 µg/µl
Stability	1yr at –70°C from date of shipment
Storage & Shipping	Store product at –70°C. For optimal storage, aliquot target into smaller quantities after centrifugation and store at recommended temperature. For most favorable performance, avoid repeated handling and multiple freeze/thaw cycles. Product shipped on dry ice.

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Activity Assay Protocol

Reaction Components

Active Methyltransferase (Catalog #: S351-380G)

Active SUV420H1 (KMT5B) (0.05 μg/μl) diluted with Methyltransferase Reaction Buffer and assayed as outlined in sample activity plot. (Note: these are suggested working dilutions and it is recommended that the researcher perform a serial dilution of Active SUV420H1 (KMT5B) for optimal results).

Methyltransferase Reaction Buffer

Buffer components: 20mM Tris-HCl, pH 8.0, 50 mM NaCl, 1 mM EDTA, 3 mM $MgCl_2$, 0.1 mg/ml BSA. Add 1mM DTT prior to use.

MTase-Glo™ Methyltransferase Assay (Promega, Catalog #: V7601)

S-Adenosyl-Methionine (SAM), 1mM
S-Adenosyl-Homocysteine (SAH), 15 μM
Methyltransferase-Glo™ Reagent, 10X
MTase-Glo™ Detection Solution, 1 bottle

Substrate (Catalog #: H13-58)

Histone H4 Peptide (1-21) diluted in distilled H_2O to a final concentration of 1mg/ml.

Assay Protocol

The SUV420H1 (KMT5B) assay is performed using the Methyltransferase-Glo™ Assays kit (Promega, Catalog #: V7601).

Step 1. Thaw the active SUV420H1 (KMT5B) and all Methyltransferase-Glo™ Assays kit reagents on ice.

Step 2. Prepare the following working solutions with Methyltransferase Reaction Buffer on ice:

- 2X final concentration of Active SUV420H1 (KMT5B) (Catalog # S351-380G)
- 2X Substrate Cocktail: 40 μM of SAM and 100ng/μl of Histone H4 Peptide (1-21) (Catalog # H13-58) in water

Step 3. In a polystyrene 96-well plate, add the following components to bring the initial reaction volume to 20 μl:

Component 1. 10 μl of 2X Substrate Cocktail

Component 2. 10 μl of 2X Active SUV420H1 (KMT5B)

Note: A blank control can be set up as outlined in step 3 by replacing the substrate working solution with an equal volume of Reaction Buffer.

Step 4. Mix the reaction on an orbital shaker for 2 minutes. Seal the plate with a plate seal and incubate at 37°C for 60 minutes

Step 5. Dilute 10X Methyltransferase-Glo™ Reagent with equal volume of nanopure water, and add 5 μl of the 5X Methyltransferase-Glo™ Reagent to all reaction wells

Step 6. Mix on an orbital shaker for 2 minutes and then incubate at room temperature for 30 minutes.

Step 7. Add 25 μl of MTase-GloTM Detection Solution to all reaction wells. Mix for 2 minutes and then incubate at room temperature for 30 minutes

Step 8. Read the plate using the KinaseGlo Luminescence Protocol on a GloMax plate reader (Promega; Cat# E7031)

Step 9. Using the SAH standard curve, determine the concentration of SAH produced (nM) and calculate the methyltransferase specific activity as outlined below. For a detailed protocol of how to determine SAH amount from RLUs, see MTase-Glo™ Methyltransferase Assay protocol at Promega's website: www.promega.com/protocols

Methyltransferase Specific Activity (SA) (nmol/min/mg)

$$= \frac{[SAH](nM) \times Reaction\ Volume(ul)}{Reaction\ Time\ (min) \times Enzyme\ Amount\ (mg)} \times 10^{-6}$$

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