abcam

Product datasheet

Anti-EHMT2/G9A antibody ab40542

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Overview

Product name Anti-EHMT2/G9A antibody

Description Rabbit polyclonal to EHMT2/G9A

Host species Rabbit

Tested applications Suitable for: IHC-P, WB

Species reactivity Reacts with: Human

Immunogen Fusion protein corresponding to Human EHMT2/ G9A aa 831-1001.

Database link: **Q96KQ7**

General notes

The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -

80°C. Avoid freeze / thaw cycle.

Purity Whole antiserum

Clonality Polyclonal

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab40542 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

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Application	Abreviews	Notes
IHC-P	★★★★☆ (1)	1/2000. Perform heat mediated antigen retrieval before commencing with IHC staining protocol.
WB	* * * * * * * (5)	1/2000 - 1/2500. Predicted molecular weight: 132 kDa. For endogeneous detection (without G9a over expression): use nuclear extracts and block overnight or perform IP prior to WB.

Target

Function

Histone methyltransferase that specifically mono- and dimethylates 'Lys-9' of histone H3 (H3K9me1 and H3K9me2, respectively) in euchromatin. H3K9me represents a specific tag for epigenetic transcriptional repression by recruiting HP1 proteins to methylated histones. Also mediates monomethylation of 'Lys-56' of histone H3 (H3K56me1) in G1 phase, leading to promote interaction between histone H3 and PCNA and regulating DNA replication. Also weakly methylates 'Lys-27' of histone H3 (H3K27me). Also required for DNA methylation, the histone methyltransferase activity is not required for DNA methylation, suggesting that these 2 activities function independently. Probably targeted to histone H3 by different DNA-binding proteins like E2F6, MGA, MAX and/or DP1. May also methylate histone H1. In addition to the histone methyltransferase activity, also methylates non-histone proteins: mediates dimethylation of 'Lys-373' of p53/TP53. Also methylates CDYL, WIZ, ACIN1, DNMT1, HDAC1, ERCC6, KLF12 and itself.

Tissue specificity

Expressed in all tissues examined, with high levels in fetal liver, thymus, lymph node, spleen and peripheral blood leukocytes and lower level in bone marrow.

Sequence similarities

Belongs to the class V-like SAM-binding methyltransferase superfamily. Histone-lysine

methyltransferase family. Suvar3-9 subfamily.

Contains 7 ANK repeats.
Contains 1 post-SET domain.
Contains 1 pre-SET domain.
Contains 1 SET domain.

Domain

The SET domain mediates interaction with WIZ.

The ANK repeats bind H3K9me1 and H3K9me2.

Post-translational

modifications

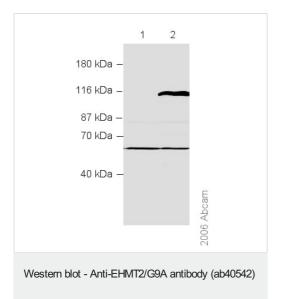
Methylated at Lys-185; automethylated.

Cellular localization

Nucleus. Chromosome. Associates with euchromatic regions. Does not associate with

heterochromatin.

Images



All lanes: Anti-EHMT2/G9A antibody (ab40542) at 1/2500 dilution

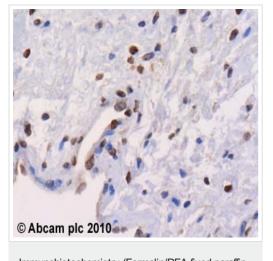
Lane 1 : Untransfected HeLa cell extract

Lane 2 : G9a transfected HeLa cell extract

Predicted band size: 132 kDa **Observed band size:** 132 kDa

Additional bands at: 60 kDa. We are unsure as to the identity of

these extra bands.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-EHMT2/G9A antibody (ab40542)

ab40542 (1:2000) staining EHMT2/ G9A in human prostate using an automated system (DAKO Autostainer Plus). Using this protocol there is strong nuclear staining.

Sections were rehydrated and antigen retrieved with the Dako 3 in 1 AR buffer citrate pH6.1 in a DAKO PT link. Slides were peroxidase blocked in 3% H2O2 in methanol for 10 mins. They were then blocked with Dako Protein block for 10 minutes (containing casein 0.25% in PBS) then incubated with primary antibody for 20 min and detected with Dako envision flex amplification kit for 30 minutes. Colorimetric detection was completed with Diaminobenzidine for 5 minutes. Slides were counterstained with Haematoxylin and coverslipped under DePeX. Please note that, for manual staining, optimization of primary antibody concentration and incubation time is recommended. Signal amplification may be required.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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