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ProductInformation

Anti-MBD2a

Developed in Rabbit Affinity Isolated Antibody

Product Number M 7568

Product Description

Anti-MBD2a is developed in rabbit using as immunogen a synthetic peptide corresponding to amino acids 2-20 of mouse MBD2a conjugated to KLH via a C-terminal added cysteine residue. The sequence is conserved in human, mouse and rat, and is not found in MBD3. Whole antiserum is fractionated and then further purified by ion-exchange chromatography to provide the IgG fraction of antiserum that is essentially free of other rabbit serum proteins.

Anti-MBD2a recognizes MBD2a. Applications include the detection of MBD2a by immunoblotting (approx. 45 kDa). Staining of the MBD2 band in immunoblotting is specifically inhibited by the immunizing peptide.

Chromatin, the physiological packaging structure of histone proteins and DNA, is a key element in the regulation of gene expression. Histones are subjected to post-translational modifications such as acetylation, phosphorylation, and methylation, and play a major role in the regulation of transcription. DNA methylation is the major modification of eukaryotic genomes, which occurs at the fifth position of cytosine in CpG dinucleotide sequences, and is associated with transcriptional repression. Silencing of transcription units have been found to occur in genes located on the inactive X-chromosome, genes silenced by genomic imprinting, and genes silenced in transformed cell lines and tumors. The subject of the protection of the subject of the subject of the protection of the subject of the su

The DNA methylation system is composed of methyl-CpG-binding proteins, as well as of DNA cytosine methyl transferases.^{3, 10} Five methyl-CpG binding proteins were isolated: MeCP2, MBD1, MBD2, MBD3, and MBD4. 10, 11 With the exceptions of MBD2 and MBD3, sequence similarity is limited to the methyl-CpG binding domains themselves (MBD). MBD2 consists of two isoforms, MBD2a and MBD2b, which are generated from a single gene; MBD2a is a 414 amino acids protein, whereas MBD2b lacks the 152 amino acids N-terminal extension. 12 MBD2a is a component of the MeCP1 corepressor complex, which is a 400-800 kDa complex containing as components MBD2, Mi-2, MTA2, MBD3, and HDAC1/2.¹³ Although MBD2 is highly similar to MBD3 and both are components of the MeCP1 complex, there are functional differences between both proteins. MBD2, but not MBD3, binds to methylated DNA in vitro or in vivo, suggesting that MBD2 targets the MeCP1 complex to methylated DNA. 12-14 Likewise MBD2 isoforms have dfferent roles in vivo. Thus, MBD2a, but not MBD2b, interacts with RNA Helicase A (RHA). 15 In connection to the ability of MBD2 to bind to methylated DNA, and its function as part of a corepressor complex, high levels of MBD2 protein appear to protect against some cancers.15, 16

Reagent

Anti-MBD2a is supplied as a solution in 0.01 M phosphate buffered saline, pH 7.4, containing 15 mM sodium azide.

Precautions and Disclaimer

Due to the sodium azide content, a material safety data sheet (MSDS) for this product has been sent to the attention of the safety officer of your institution. Consult the MSDS for information regarding hazardous and safe handling practices.

Storage/Stability

For continuous use, store at 2-8 °C for up to one month. For extended storage, freeze in working aliquots. Repeated freezing and thawing is not recommended. Storage in frost-free freezers is also not recommended. If slight turbidity occurs upon prolonged storage, clarify the solution by centrifugation before use. Working dilutions should be discarded if not used within 12 hours.

Product Profile

By immunoblotting, a working antibody dilution of 1:400-1:800 is recommended using nuclear extracts of HeLa cells.

Recommendation: For immunoblotting, dilute the antibody in phosphate buffered saline containing 0.5% non-fat dry milk and 0.05% TweenTM 20.

Note: In order to obtain the best results using various techniques and preparations, we recommend determining the optimal working dilutions by titration.

References

- Kornberg, R.D., et al., Cell, 98, 285-294 (1999).
- Strahl, B.D., and Allis, C.D., Nature, 403, 41-45 (2000).
- 3. Bird, A., and Wolffe, A.P., Cell, **99**, 451-454 (1999).
- 4. Razin, A., and Szyf, M., Biochim. Biophys. Acta, **782**, 331-342 (1984).
- 5. Nur, I., et al., Nuc. Acids Res., **16**, 9233-9251 (1988).
- Li, M., et al., Gene, 301, 43-51 (2002).
- 7. Razin, A., and Cedar, H., Cell, **77**, 473-476 (1994).
- 8. Riggs, A.D., and Pfeifer, G.P., Trends Genet., **8**, 169-174 (1992).
- Sakai, T., et al., Am. J. Hum. Genet., 48, 880-888 (1991).
- Hendrich, B., and Bird, A., Mol. Cell. Biol., 18, 6538-6547 (1998).
- 11. Hendrich, B., et al., Nature, **401**, 301-304 (2000).
- 12. Hendrich, B., and Bird, A., Mol. Cell. Biol., **18**, 6538--6547 (1998).
- 13. Ng, H.H., et al., Nat. Genet., 23, 58-61 (1999).
- 14. Brackertz, M., et al., J. Biol. Chem., **277**, 40958-40966 (2002).
- 15. Fujita, H., et al., Mol. Cell. Biol., **23**, 2645-2657 (2003).
- 16. Sansom, et al., Nat. Genet., **34**, 145-147 (2003).

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