MECP2 Rabbit Polyclonal Antibody



DNA methylation is the major modification of eukaryotic genomes and plays an essential role

in mammalian development. Human proteins MECP2, MBD1, MBD2, MBD3, and MBD4

comprise a family of nuclear proteins related by the presence in each of a methyl-CpG binding domain (MBD). Each of these proteins, with the exception of MBD3, is capable of binding

specifically to methylated DNA. MECP2, MBD1 and MBD2 can also repress transcription from methylated gene promoters. In contrast to other MBD family members, MECP2 is X-linked and

subject to X inactivation. MECP2 is dispensible in stem cells, but is essential for embryonic

development. MECP2 gene mutations are the cause of most cases of Rett syndrome, a progressive neurologic developmental disorder and one of the most common causes of mental

retardation in females. Alternative splicing results in multiple transcript variants encoding

CAB16917

Product Information

Size:

20uL, 50uL, 100uL, 200uL

Observed MW:

Refer to figures

Calculated MW:

52kDa/53kDa

Applications:

IHC

Reactivity:

Human

Immunogen information

Protein Background

Gene ID:

different isoforms.

4204

Uniprot P51608

Antibody Information

Recommended dilutions:

IHC 1:50 - 1:200

Source:

Rabbit

Isotype: IgG

Immunogen:

Synonyms:

A synthetic peptide corresponding to a sequence within amino

MECP2; AUTSX3; MRX16; MRX79; MRXS13; MRXSL; PPMX; RS; RTS;

acids 1-100 of human MECP2 (NP_004983.1).

Storage:

Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.02%

sodium azide, 50% glycerol, pH7.3.

Affinity purification

Purification: