

## MeCP2 rabbit pAb

Cat No.: ES20671

For research use only

## Overview

Product Name MeCP2 rabbit pAb

Host speciesRabbitApplicationsWBSpecies Cross-ReactivityHumanRecommended dilutionsWB: 1:2000

ImmunogenSynthetic Peptide of MeCP2 AA range: 313-363SpecificityThe antibody detects endogenous MeCP2 proteins.FormulationLiquid in PBS containing 50% glycerol, 0.5% BSA and

0.02% sodium azide.

**Storage** Store at -20°C. Avoid repeated freeze-thaw cycles.

**Protein Name** Methyl-CpG-binding protein 2

Gene Name MECP2

Cellular localization Nucleus . Colocalized with methyl-CpG in the

genome. Colocalized with TBL1X to the

heterochromatin foci. .

**Purification** The antibody was affinity-purified from rabbit

antiserum by affinity-chromatography using

epitope-specific immunogen.

**Clonality** Polyclonal

Concentration

Observed band 53kD Human Gene ID 4204 Human Swiss-Prot Number P51608

Alternative Names Methyl-CpG-binding protein 2 (MeCp-2 protein)

(MeCp2)

**Background** 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.

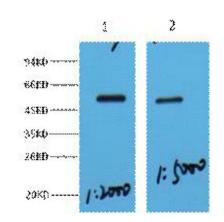


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MECP2, MBD1 and MBD2 can also repress transcription from methylated gene promoters. In contrast to other MBD family members, MECP2 is X-linked and subject to X inactivation. MECP2 is dispensible in stem cells, but is essential for embryonic development. 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 different isofor



Western blot analysis of Hela, diluted at 1) 1:2000 2) 1:5000. Secondary antibody(catalog#:RS0002) was diluted at 1:20000

