

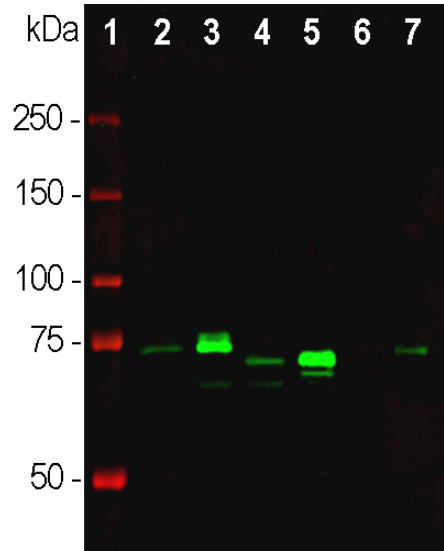
Ordering Information
 Web www.encorbio.com
 Email admin@encorbio.com
 Phone 352-372-7022
 Fax 352-372-7066

HGNC Name: MECP2
UniProt: P51608
RRID: AB_2572345
Immunogen: Full length recombinant human MeCP2
Format: Affinity purified at 1mg/mL in PBS, 50% glycerol, 5mM Na₂S₂O₈
Storage: Stable at 4°C for one year, for longer term store at -20°C
Recommended dilutions:
 WB: 1:10,000-1:20,000 IF/ICC or IHC: 1:2,000

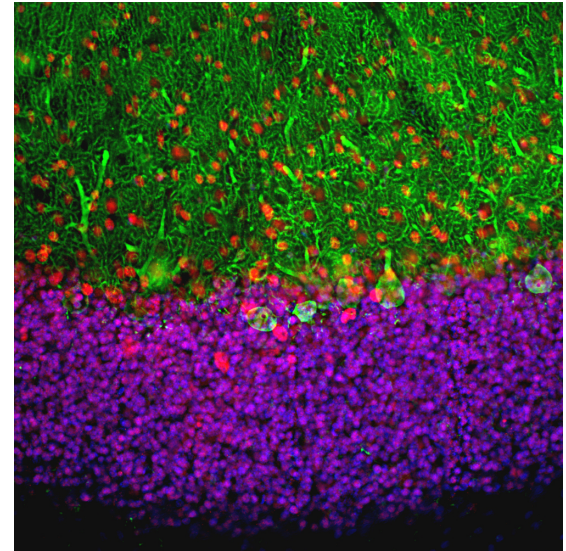
References:

1. Klose RJ, et al. DNA binding selectivity of MeCP2 due to a requirement for A/T sequences adjacent to methyl-CpG. *Mol. Cell* 19:667-78 (2005).
2. Amir RE, et al. Rett syndrome is caused by mutations in X-linked MECP2, encoding methyl-CpG-binding protein 2. *Nat. Genet.* 23:185-8 (1999).
3. Pohodich AE, Zoghbi HY. Rett syndrome: disruption of epigenetic control of postnatal neurological functions. *Hum. Mol. Genet.* 24:R10-6 (2010).
4. Ramocki MB, Tavyev YJ, Peters SU. The MECP2 duplication syndrome. *Am. J. Med. Genet. A.* 152A:1079-88 (2010).
5. Zhou Z, et al. Brain-specific phosphorylation of MeCP2 regulates activity dependent *Bdnf* transcription, dendritic growth, and spine maturation. *Neuron* 52:255-69 (2006).
6. Deng JV, et al. MeCP2 in the nucleus acumbens contributes to neural and behavioral responses to psychostimulants. *Nat. Neurosci.* 13:1128-36 (2010).
7. Tao J, et al. Phosphorylation of MeCP2 at Serine 80 regulates its chromatin association and neurological function. *PNAS* 24:106 (2009).

Applications	Host	Isotype	Molecular Wt.	Species Cross-Reactivity
WB, IF/ICC, IHC	Rabbit		70-75kDa by SDS-PAGE	Hu, mo, rt, ms



Western blot analysis of tissue and cell lysates using rabbit pAb to MeCP2, RPCA-MeCP2, dilution 1:20,000 in green: [1] protein standard (red), [2] Rat whole brain, [3] nuclear fraction of rat brain, [4] mouse whole brain, [5] nuclear fraction of mouse brain, [6] HeLa cells, [7] C6 cells. Major band at about 70-75 kDa corresponds to MeCP2 protein, predominantly detected in the nuclear fraction of the lysates. The apparent molecular weights of MeCP2 based on SDS-PAGE mobility are variable between species and higher than the actual MeCP2 molecular weight which is ~54kDa



Immunofluorescent analysis of rat cerebellum section costained with rabbit pAb to MeCP2, RPCA-MeCP2, dilution 1:5,000, in red, and chicken pAb to calbindin, CPCA-Calb, dilution 1:2,000, in green. Blue is DAPI staining of nuclear DNA. Following transcardial perfusion with 4% paraformaldehyde, brain was post fixed for 24 hours, cut to 45µm, and free-floating sections were stained with above antibodies. The MeCP2 antibody selectively stains nuclei of neuronal cells to a variable degree. Calbindin, often used as Purkinje cell marker, is prominently expressed in dendrites and perikarya of these cells in the cerebellar molecular layer.

Background:

Methyl-CpG Binding Protein 2 (MeCP2) is widely expressed in tissues and particularly heavily in neurons. It is a nuclear protein which was found to bind methylated cytosines in CpG islands in DNA, but which may also bind hydroxymethyl cytosine residues (1-3). Methyl and hydroxymethyl cytosines are generally found in regions of DNA regulating the expression of specific genes and MeCP2 is thought to function as a gene regulator, though it may have other functions. The MeCP2 protein contains one methyl-CpG binding domain and two AT hook domains, highly basic peptides which bind AT rich DNA sequences. The MeCP2 gene, *MECP2* is located on the X-chromosome and loss of function mutations of the gene cause Rett syndrome, a relatively rare neurodevelopmental, autistic disorder which presents almost exclusively in females, since similar mutation of *MECP2* in males is almost invariably lethal (3). Rett patients are apparently normal at birth but develop neurological problems over the period when extensive synaptogenesis is occurring, which coincides with the expression of high levels of MeCP2 in normal brain. The Rett brain shows smaller more densely packed neurons with less extensive dendritic branching (2). Duplication of the *MECP2* gene was found to be the cause of a different neurodevelopmental disorder referred to a *MECP2* duplication syndrome (4). Levels of MeCP2 normally alter dynamically and the protein is regulated by phosphorylation on multiple sites (5-7).

The RPCA-MeCP2 antibody was made against full length recombinant human MeCP2 expressed in and purified from *E. coli* and can be used to identify neurons transiently expressing high levels of this nuclear protein. We also supply a mouse monoclonal and a chicken polyclonal antibody to the same protein, *MCA-4F11* and *CPCA-MeCP2*.

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Abbreviation Key:

mAb—Monoclonal Antibody pAb—Polyclonal Antibody WB—Western Blot IF—Immunofluorescence ICC—Immunocytochemistry IHC—Immunohistochemistry E—ELISA Hu—Human Mo—Monkey Do—Dog Rt—Rat Ms—Mouse Co—Cow Pi—Pig Ho—Horse Ch—Chicken Dr—D. rerio Dm—D. melanogaster Sm—S. mutans Ce—C. elegans Sc—S. cerevisiae Sa—S. aureus Ec—E. coli.