

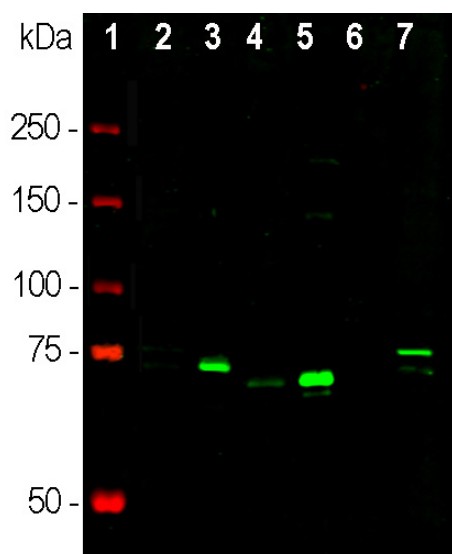
**Ordering Information**  
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**HGNC Name:** MECP2  
**UniProt:** P51608  
**RRID:** AB\_2737436  
**Immunogen:** Full length recombinant human MeCP2 expressed in and purified from *E. coli*.  
**Format:** Concentrated IgY preparation in PBS plus 0.02% Na<sub>2</sub>S<sub>2</sub>O<sub>3</sub>  
**Storage:** Stable at 4°C for several months. For longer term store at -20°C, minimize freeze/thaw cycles.  
**Recommended dilutions:**  
WB: 1:10,000-1:20,000. IF/IHC 1:1,000-1:4,000

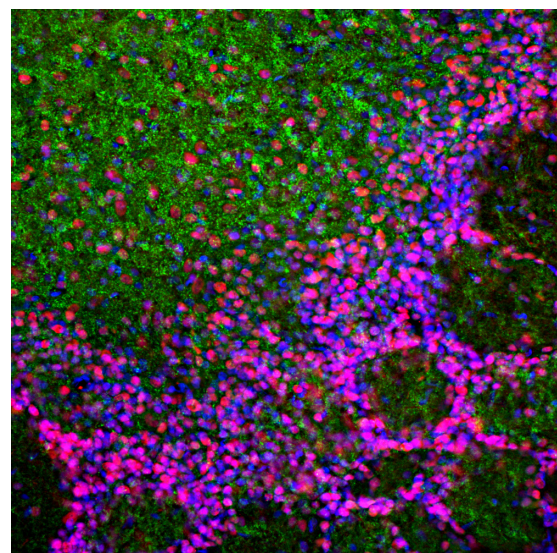
### References:

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* 106:106 (2009).

Applications	Host	Isotype	Molecular Wt.	Species Cross-Reactivity
WB, ICC/IF, IHC	Chicken		70-75 kDa by SDS-PAGE	Hu, Mo, Rt, Ms



Western blot analysis of tissue and cell lysates using chicken pAb to MeCP2, CPCA-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 lysate, [6] C6 cell lysate, and [7] SH-SY5Y cell lysate. The strong band at about 75kDa corresponds to the MeCP2 protein. The MeCP2 proteins of rat and human origin are known to migrate slightly differently on SDS-PAGE gels compared to that of mouse, and as a result bands appear at somewhat different apparent molecular weights.



Immunofluorescent analysis of rat olfactory bulb section stained with chicken pAb to MeCP2, CPCA-MeCP2, dilution 1:2,000 in red and costained with mouse mAb to α-synuclein, MCA-2A7, dilution 1:1,000, in green. The blue is DAPI staining of nuclear DNA. Following transcardial perfusion of rat with 4% paraformaldehyde, brain was post fixed for 24 hours, cut to 45μm, and free-floating sections were stained with the above antibodies. The MeCP2 antibody specifically labels the nuclei of neuronal cells while the MCA-2A7 antibody reveals α-synuclein protein concentrated in presynaptic regions.

### 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 CPCA-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 mouse monoclonal and rabbit polyclonal antibodies to the same protein, MCA-4F11 and RPCA-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.