# **MECP2** Rabbit pAb

Catalog No.: A0706



### **Basic Information**

### **Observed MW**

52kDa

### **Calculated MW**

52kDa

#### Category

Polyclonal Antibody

### **Applications**

WB,FC

### **Cross-Reactivity**

Human

## **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. 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 cognitive disability in females. Alternative splicing results in multiple transcript variants encoding different isoforms.

# **Recommended Dilutions**

WB 1:500 - 1:1000

FC 1:20 - 1:50

### **Immunogen Information**

**Gene ID Swiss Prot** 4204 P51608

### **Immunogen**

Synthetic peptide. This information is considered to be commercially sensitive.

### **Synonyms**

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

### Contact

www.abclonal.com

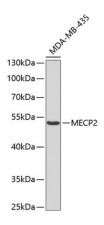
### **Product Information**

SourceIsotypePurificationRabbitIgGAffinity purification

### Storage

Store at 4°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.02% sodium azide,pH7.3.

# Validation Data



Western blot analysis of lysates from MDA-MB-435 cells, using MECP2 Rabbit pAb (A0706).

Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.