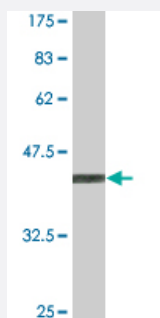


MECP2 monoclonal antibody (M04A), clone 2E8

Catalog # H00004204-M04A

Size 200 uL

Applications



Western Blot detection against Immunogen (35.53 KDa) .

Specification

Product Description	Mouse monoclonal antibody raised against a partial recombinant MECP2.
Immunogen	MECP2 (AAH11612, 81 a.a. ~ 170 a.a) partial recombinant protein with GST tag. MW of the GST tag alone is 26 KDa.
Sequence	PKQRRSIIIRDRGPMYDDPTLPEGWTRKLKQRKSGRSAGKYDVYLINPQGKAFRSKVELIAYFEKVG DTSLDAPNDFDFTVTGRGSPSRREQ
Host	Mouse
Reactivity	Human
Isotype	IgG2a Kappa
Quality Control Testing	Antibody Reactive Against Recombinant Protein. Western Blot detection against Immunogen (35.53 KDa) .
Storage Buffer	In ascites fluid
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

Applications

- Western Blot (Recombinant protein)

[Protocol Download](#)

- ELISA

Gene Info — MECP2

Entrez GeneID [4204](#)

GeneBank Accession# [BC011612](#)

Protein Accession# [AAH11612](#)

Gene Name MECP2

Gene Alias AUTSX3, DKFZp686A24160, MRX16, MRX79, MRXS13, MRXSL, PPMX, RTS, RTT

Gene Description methyl CpG binding protein 2 (Rett syndrome)

Omim ID [105830](#) [300005](#) [300055](#) [300260](#) [300496](#) [300673](#) [312750](#)

Gene Ontology [Hyperlink](#)

Gene Summary

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 dispensable 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. [provided by RefSeq]

Other Designations OTTHUMP00000026021|methyl CpG binding protein 2

Disease

- [Alzheimer disease](#)
- [Angelman syndrome](#)
- [Attention Deficit Disorder with Hyperactivity](#)
- [Autistic Disorder](#)

- [Brain Diseases](#)
- [Cardiovascular Diseases](#)
- [Child Development Disorders](#)
- [Chromosome Aberrations](#)
- [Cognition Disorders](#)
- [Developmental Disabilities](#)
- [Diabetes Mellitus](#)
- [Disease Progression](#)
- [Edema](#)
- [Epilepsy](#)
- [Fractures](#)
- [Fragile X syndrome](#)
- [Genetic Diseases](#)
- [Genetic Predisposition to Disease](#)
- [Lupus Erythematosus](#)
- [Mental Disorders](#)
- [Mental Retardation](#)
- [Microcephaly](#)
- [Motor Skills](#)
- [Movement Disorders](#)
- [NARP](#)
- [Nervous System Diseases](#)
- [Obesity](#)
- [Pain](#)
- [Prader-Willi syndrome](#)

- [Psychomotor Performance](#)
- [Psychotic Disorders](#)
- [Rett syndrome](#)
- [Schizophrenia](#)
- [Scoliosis](#)
- [Seizures](#)
- [Stereotypic Movement Disorder](#)