

EMD Antibody (monoclonal) (M01)**Mouse monoclonal antibody raised against a partial recombinant EMD.****Catalog # AT1898a****Specification**

EMD Antibody (monoclonal) (M01) - Product Information

Application	WB, IHC, IF, E
Primary Accession	P50402
Other Accession	BC000738
Reactivity	Human
Host	mouse
Clonality	Monoclonal
Isotype	IgG2a Kappa
Calculated MW	28994

EMD Antibody (monoclonal) (M01) - Additional Information**Gene ID** 2010**Other Names**

Emerin, EMD, EDMD, STA

Target/Specificity

EMD (AAH00738, 1 a.a. ~ 110 a.a) partial recombinant protein with GST tag. MW of the GST tag alone is 26 KDa.

Dilution

WB~~1:500~1000

IHC~~1:100~500

IF~~1:50~200

E~~N/A

Format

Clear, colorless solution in phosphate buffered saline, pH 7.2 .

Storage

Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

Precautions

EMD Antibody (monoclonal) (M01) is for research use only and not for use in diagnostic or therapeutic procedures.

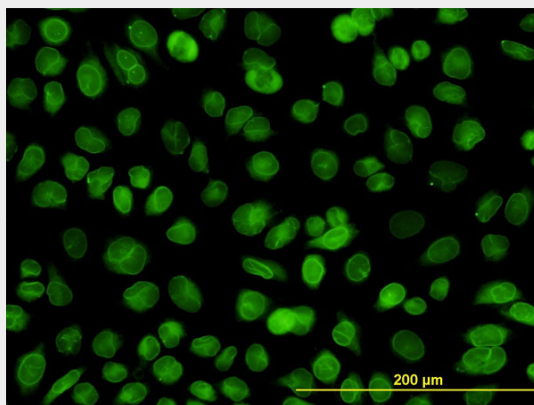
EMD Antibody (monoclonal) (M01) - Protocols

Provided below are standard protocols that you may find useful for product applications.

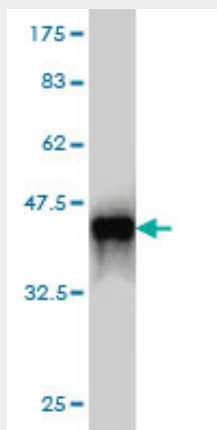
- [Western Blot](#)

- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

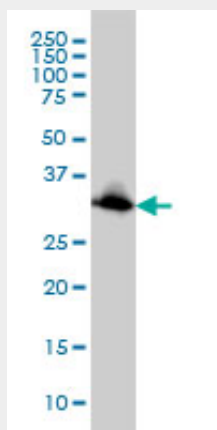
EMD Antibody (monoclonal) (M01) - Images



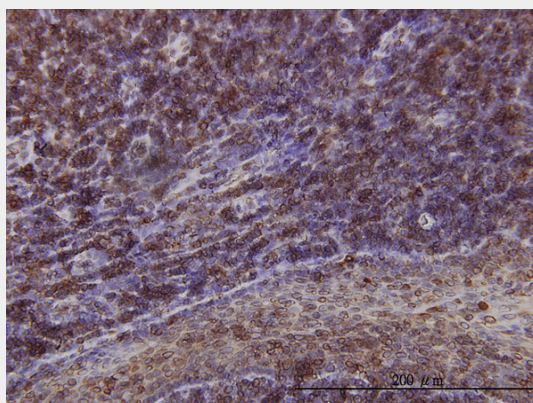
Immunofluorescence of monoclonal antibody to EMD on HeLa cell. [antibody concentration 10 ug/ml]



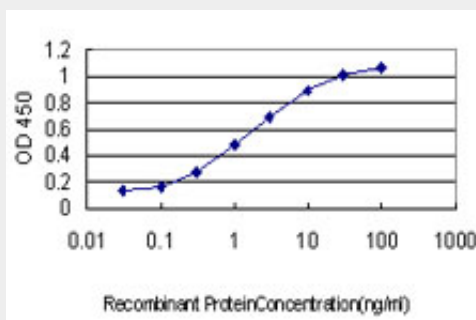
Antibody Reactive Against Recombinant Protein. Western Blot detection against Immunogen (37.84 KDa) .



EMD monoclonal antibody (M01), clone 3B9 Western Blot analysis of EMD expression in HeLa (Cat # AT1898a)



Immunoperoxidase of monoclonal antibody to EMD on formalin-fixed paraffin-embedded human tonsil. [antibody concentration 1 ug/ml]



Detection limit for recombinant GST tagged EMD is approximately 0.03ng/ml as a capture antibody.

EMD Antibody (monoclonal) (M01) - Background

Emerin is a serine-rich nuclear membrane protein and a member of the nuclear lamina-associated protein family. It mediates membrane anchorage to the cytoskeleton. Dreifuss-Emery muscular dystrophy is an X-linked inherited degenerative myopathy resulting from mutation in the emerin gene.

EMD Antibody (monoclonal) (M01) - References

Variation at the NFATC2 Locus Increases the Risk of Thiazolinedione-Induced Edema in the Diabetes REduction Assessment with ramipril and rosiglitazone Medication (DREAM) Study. Bailey SD, et al. Diabetes Care, 2010 Jul 13. PMID 20628086. A novel custom resequencing array for dilated cardiomyopathy. Zimmerman RS, et al. Genet Med, 2010 May. PMID 20474083. Emery-Dreifuss dystrophy: a 4-year follow-up on a laminopathy of special interest. Hausmanowa-Petrusewicz I, et al. Neurol Neurochir Pol, 2009 Sep-Oct. PMID 20054742. Gene-centric association signals for lipids and apolipoproteins identified via the HumanCVD BeadChip. Talmud PJ, et al. Am J Hum Genet, 2009 Nov. PMID 19913121. Tyrosine phosphorylation of nuclear-membrane protein emerin by Src, Abl and other kinases. Tiffert KE, et al. J Cell Sci, 2009 Oct 15. PMID 19789182.