

DYM Antibody (monoclonal) (M01)**Mouse monoclonal antibody raised against a partial recombinant DYM.****Catalog # AT1833a****Specification**

DYM Antibody (monoclonal) (M01) - Product Information

Application	E
Primary Accession	Q7RTS9
Other Accession	NM_017653
Reactivity	Human
Host	mouse
Clonality	Monoclonal
Isotype	IgG1 Kappa
Calculated MW	75935

DYM Antibody (monoclonal) (M01) - Additional Information**Gene ID** 54808**Other Names**

Dymeclin, Dyggve-Melchior-Clausen syndrome protein, DYM

Target/Specificity

DYM (NP_060123, 343 a.a. ~ 430 a.a) partial recombinant protein with GST tag. MW of the GST tag alone is 26 KDa.

Dilution

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

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

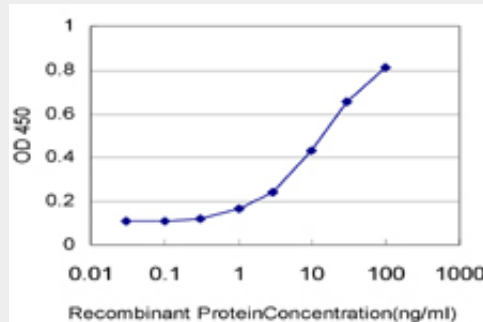
DYM Antibody (monoclonal) (M01) - Protocols

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](#)

DYM Antibody (monoclonal) (M01) - Images



Detection limit for recombinant GST tagged DYM is approximately 1ng/ml as a capture antibody.

DYM Antibody (monoclonal) (M01) - Background

This gene encodes a protein which is necessary for normal skeletal development and brain function. Mutations in this gene are associated with two types of recessive osteochondrodysplasia, Dyggve-Melchior-Clausen (DMC) dysplasia and Smith-McCort (SMC) dysplasia, which involve both skeletal defects and mental retardation.

DYM Antibody (monoclonal) (M01) - References

An association study between the dymeclin gene and schizophrenia in the Japanese population. Yazaki S, et al. J Hum Genet, 2010 Sep. PMID 20555340. The role of height-associated loci identified in genome wide association studies in the determination of pediatric stature. Zhao J, et al. BMC Med Genet, 2010 Jun 14. PMID 20546612. Personalized smoking cessation: interactions between nicotine dose, dependence and quit-success genotype score. Rose JE, et al. Mol Med, 2010 Jul-Aug. PMID 20379614. The gene responsible for Dyggve-Melchior-Clausen syndrome encodes a novel peripheral membrane protein dynamically associated with the Golgi apparatus. Dimitrov A, et al. Hum Mol Genet, 2009 Feb 1. PMID 18996921. Genome-wide association analysis identifies 20 loci that influence adult height. Weedon MN, et al. Nat Genet, 2008 May. PMID 18391952.