

DLX1 Antibody (monoclonal) (M02)**Mouse monoclonal antibody raised against a partial recombinant DLX1.****Catalog # AT1775a****Specification**

DLX1 Antibody (monoclonal) (M02) - Product Information

Application	WB, E
Primary Accession	P56177
Other Accession	NM_178120
Reactivity	Human
Host	mouse
Clonality	Monoclonal
Isotype	IgG2a Kappa
Calculated MW	27320

DLX1 Antibody (monoclonal) (M02) - Additional Information**Gene ID** 1745**Other Names**

Homeobox protein DLX-1, DLX1

Target/Specificity

DLX1 (NP_835221, 152 a.a. ~ 255 a.a) partial recombinant protein with GST tag. MW of the GST tag alone is 26 KDa.

Dilution

WB~~1:500~1000

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

DLX1 Antibody (monoclonal) (M02) is for research use only and not for use in diagnostic or therapeutic procedures.

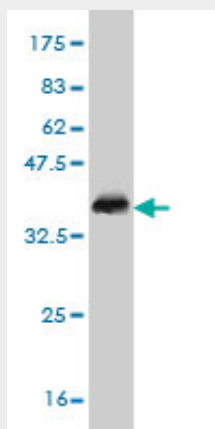
DLX1 Antibody (monoclonal) (M02) - Protocols

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

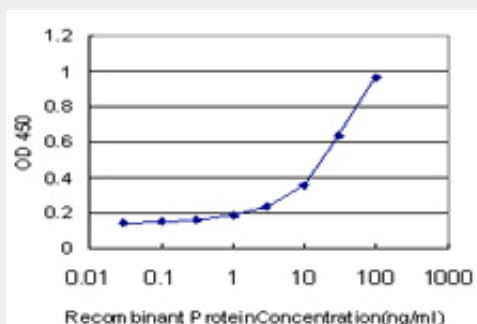
- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)

- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

DLX1 Antibody (monoclonal) (M02) - Images



Antibody Reactive Against Recombinant Protein. Western Blot detection against Immunogen (37.55 KDa) .



Detection limit for recombinant GST tagged DLX1 is approximately 1ng/ml as a capture antibody.

DLX1 Antibody (monoclonal) (M02) - Background

This gene encodes a member of a homeobox transcription factor gene family similar to the Drosophila distal-less gene. The encoded protein is localized to the nucleus where it may function as a transcriptional regulator of signals from multiple TGF- β superfamily members. The encoded protein may play a role in the control of craniofacial patterning and the differentiation and survival of inhibitory neurons in the forebrain. This gene is located in a tail-to-tail configuration with another member of the family on the long arm of chromosome 2. Alternatively spliced transcript variants encoding different isoforms have been described.

DLX1 Antibody (monoclonal) (M02) - References

Maternal genes and facial clefts in offspring: a comprehensive search for genetic associations in two population-based cleft studies from Scandinavia. Jugessur A, et al. PLoS One, 2010 Jul 9. PMID 20634891. High-density association study of 383 candidate genes for volumetric BMD at the femoral neck and lumbar spine among older men. Yerges LM, et al. J Bone Miner Res, 2009 Dec. PMID 19453261. A common variant in DRD3 receptor is associated with autism spectrum disorder. de Krom M, et al. Biol Psychiatry, 2009 Apr 1. PMID 19058789. The DLX1 and DLX2 genes and susceptibility to autism spectrum disorders. Liu X, et al. Eur J Hum Genet, 2009 Feb. PMID

18728693.Association analysis of schizophrenia on 18 genes involved in neuronal migration: MDGA1 as a new susceptibility gene. K?hler AK, et al. Am J Med Genet B Neuropsychiatr Genet, 2008 Oct 5. PMID 18384059.