

TXNDC3 Antibody (monoclonal) (M01)

Mouse monoclonal antibody raised against a partial recombinant TXNDC3. Catalog # AT4416a

Product Information

Application	WB
Primary Accession	<u>Q8N427</u>
Other Accession	<u>NM_016616</u>
Reactivity	Human
Host	mouse
Clonality	monoclonal
Isotype	IgG2a Kappa
Clone Names	1G5
Calculated MW	67270

Additional Information

Gene ID	51314
Other Names	Thioredoxin domain-containing protein 3, NM23-H8, NME/NM23 family member 8, Spermatid-specific thioredoxin-2, Sptrx-2, NME8, SPTRX2, TXNDC3
Target/Specificity	TXNDC3 (NP_057700, 530 a.a. ~ 586 a.a) partial recombinant protein with GST tag. MW of the GST tag alone is 26 KDa.
Dilution	WB~~1:500~1000
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	TXNDC3 Antibody (monoclonal) (M01) is for research use only and not for use in diagnostic or therapeutic procedures.

Background

This gene encodes a protein with an N-terminal thioredoxin domain and three C-terminal nucleoside diphosphate kinase (NDK) domains, but the NDK domains are thought to be catalytically inactive. The sea urchin ortholog of this gene encodes a component of sperm outer dynein arms, and the protein is implicated in ciliary function. Mutations in this gene are implicated in primary ciliary dyskinesia type 6.

References

Personalized smoking cessation: interactions between nicotine dose, dependence and quit-success genotype score. Rose JE, et al. Mol Med, 2010 Jul-Aug. PMID 20379614.Field synopsis and synthesis of genetic

association studies in osteoarthritis: the CUMAGAS-OSTEO information system. Zintzaras E, et al. Am J Epidemiol, 2010 Apr 15. PMID 20237151.Nme protein family evolutionary history, a vertebrate perspective. Desvignes T, et al. BMC Evol Biol, 2009 Oct 23. PMID 19852809.Association of single-nucleotide polymorphisms in RHOB and TXNDC3 with knee osteoarthritis susceptibility: two case-control studies in East Asian populations and a meta-analysis. Shi D, et al. Arthritis Res Ther, 2008. PMID 18471322.A common variant in combination with a nonsense mutation in a member of the thioredoxin family causes primary ciliary dyskinesia. Duriez B, et al. Proc Natl Acad Sci U S A, 2007 Feb 27. PMID 17360648.



Images

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