Revision 1



For Research Use Only. Not for Use in Diagnostic Procedures.

Applications: W	Reactivity: H M R Mk	Sensitivity: Endogenous	MW (kDa): 70	Source/Isotype: Rabbit	UniProt ID: #P35637	Entrez-Gene Id: 2521
Product Usage Information		Application Western Blotting			Dilution 1:1000	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		FUS/TLS Antibody recognizes endogenous levels of total FUS/TLS protein.				
Species predicted to react based on 100% sequence homology		Hamster, Bovine, Horse, Guinea Pig				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Gly272 of human TLS/FUS protein. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		FUS/TLS (fused in sarcoma/translocated in liposarcoma) was initially identified by investigators as a component of fusion proteins found in a variety of cancers, such as myxoid liposarcoma, acute myeloid leukemia, and Ewing's tumor (1). FUS/TLS fusion with the DNA-binding domain of transcription activators, such as CHOP and ERG, leads to aberrant transcription of target genes that is thought by researchers to lead to tumor development (1-5). FUS/TLS is involved in a wide range of RNA processing events, such as pre-mRNA splicing, mRNA transcription, and miRNA processing (1,6). In addition to its role in RNA metabolism, FUS/TLS maintains genomic stability and co-regulates gene expression by interacting with various transcription factors such as nuclear receptors, YB-1, p65 subunit of NF-KB, TFIID, and RUNX2 (1,6,7). More recently, researchers have found several mutations of FUS/TLS in ALS (amyotrophic lateral sclerosis) and FTLD (frontotemporal lobar degeneration) patients that causes cytoplasmic mislocalization of FUS/TLS (6,8-12).				
Background References		 Yang, S. et al. (2010) <i>Int J Biochem Cell Biol</i> 42, 1408-11. Crozat, A. et al. (1993) <i>Nature</i> 363, 640-4. Rabbitts, T.H. et al. (1993) <i>Nat Genet</i> 4, 175-80. Law, W.J. et al. (2006) <i>Brief Funct Genomic Proteomic</i> 5, 8-14. Prasad, D.D. et al. (1994) <i>Oncogene</i> 9, 3717-29. Lagier-Tourenne, C. et al. (2010) <i>Hum Mol Genet</i> 19, R46-64. Baechtold, H. et al. (1999) <i>J Biol Chem</i> 274, 34337-42. Hewitt, C. et al. (2010) <i>Arch Neurol</i> 67, 455-61. Vance, C. et al. (2009) <i>Science</i> 323, 1208-11. Van Langenhove, T. et al. (2010) <i>Neurology</i> 74, 366-71. Da Cruz, S. and Cleveland, D.W. (2011) <i>Curr Opin Neurobiol</i> 21, 904-19. Hock, E.M. et al. (2018) <i>Cell Rep</i> 24, 987-1000.e7. 				
Species Reactiv	vity	Species reactivity is de	termined by testing	g in at least one approve	ed application (e.g.,	western blot).
Western Blot Buffer		IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.				
Applications Key		W: Western Blotting				
Cross-Reactivity Key		H: Human M: Mouse R: Rat Mk: Monkey				
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