

DESCRIPTION

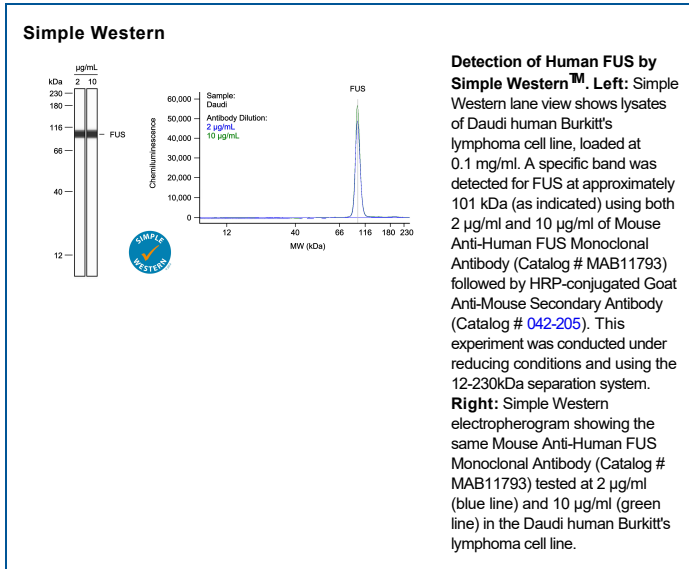
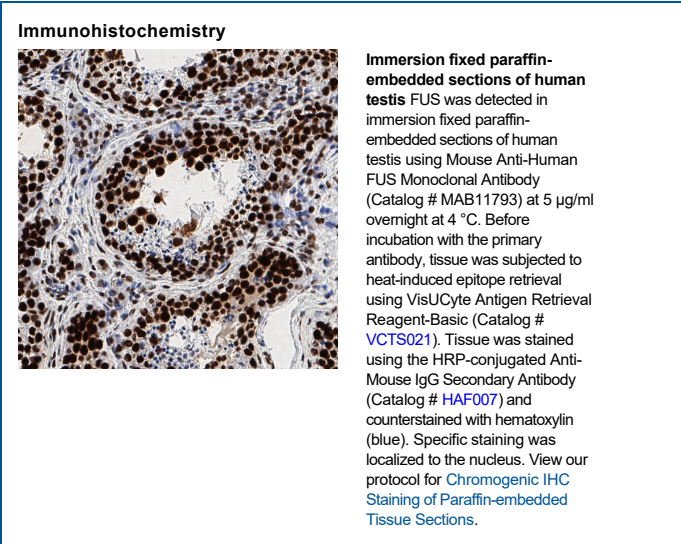
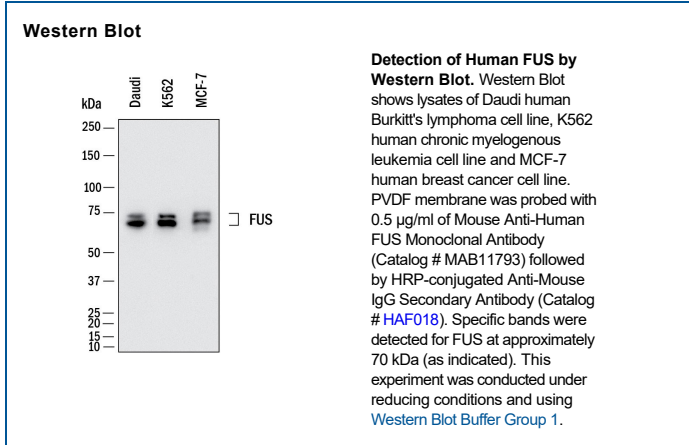
Species Reactivity	Human
Specificity	Detects recombinant human FUS and human TAF-15 in Direct ELISA.
Source	Monoclonal Mouse IgG _{2B} Clone # 1120264)
Purification	Protein A or G purified from hybridoma culture supernatant
Immunogen	<i>E. coli</i> - derived recombinant human FUS Asn263-Gly378 Accession # P35637
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose.

APPLICATIONS

Please Note: Optimal dilutions should be determined by each laboratory for each application. [General Protocols](#) are available in the *Technical Information* section on our website.

	Recommended Concentration	Sample
Western Blot	0.5 µg/mL	Daudi human Burkitt's lymphoma cell line, K562 human chronic myelogenous leukemia cell line and MCF-7 human breast cancer cell line
Immunohistochemistry	0.25-25 µg/mL	Immersion fixed paraffin-embedded sections of human testis
Simple Western	2-10 µg/mL	Daudi human Burkitt's lymphoma cell line

DATA



PREPARATION AND STORAGE

Reconstitution	Reconstitute lyophilized material at 0.2 mg/ml in sterile PBS. For liquid material, refer to CoA for concentration.
Shipping	Lyophilized product is shipped at ambient temperature. Liquid small pack size (-SP) is shipped with polar packs. Upon receipt, store immediately at the temperature recommended below.
Stability & Storage	<p>Use a manual defrost freezer and avoid repeated freeze-thaw cycles.</p> <ul style="list-style-type: none"> • 12 months from date of receipt, -20 to -70 °C as supplied. • 1 month, 2 to 8 °C under sterile conditions after reconstitution. • 6 months, -20 to -70 °C under sterile conditions after reconstitution.

BACKGROUND

Fused in sarcoma (FUS) is a multifunctional RNA-binding protein with a molecular weight of approximately 68 kDa. FUS belongs to the family of heterogeneous nuclear ribonucleoproteins (hnRNPs) and plays critical roles in RNA metabolism, including RNA splicing, transport, translation, and stability. Beyond its functions in RNA handling, FUS contributes to DNA damage repair and transcription regulation, highlighting its importance in cellular homeostasis. FUS is ubiquitously expressed across various tissues, with predominant localization in the nucleus; however, its mislocalization to the cytoplasm is directly implicated in neurodegenerative diseases such as amyotrophic lateral sclerosis (ALS) and frontotemporal dementia (FTD). Mutations in the FUS gene have been identified as causative factors in familial ALS, where altered FUS protein function contributes to protein aggregation and neuronal toxicity. Additionally, FUS is involved in cellular stress responses, forming dynamic assemblies known as stress granules under adverse conditions. Aberrant FUS expression or activity has garnered attention as a key biomarker for neurodegenerative disorders and a promising target for therapeutic interventions.

References:

1. Vance C, Rogelj B, Hortobágyi T, De Vos KJ, Nishimura AL, Sreedharan J, Hu X, Smith B, Ruddy D, Wright P, Ganesalingam J, Williams KL, Tripathi V, Al-Saraj S, Al-Chalabi A, Leigh PN, Blair IP, Nicholson G, de Belleruche J, Gallo JM, Miller CC, Shaw CE. Mutations in FUS, an RNA processing protein, cause familial amyotrophic lateral sclerosis type 6. *Science*. 2009 Feb 27;323(5918):1208-1211. doi: 10.1126/science.1165942. PMID: 19251628; PMCID: PMC4516382.
2. Sama RR, Ward CL, Bosco DA. Functions of FUS/TLS from DNA repair to stress response: implications for ALS. *ASN Neuro*. 2014 Jun 1;6(4):1759091414544472. doi: 10.1177/1759091414544472. PMID: 25289647; PMCID: PMC4189536.
3. Yang S, Warraich ST, Nicholson GA, Blair IP. Fused in sarcoma/translocated in liposarcoma: a multifunctional DNA/RNA binding protein. *Int J Biochem Cell Biol*. 2010 Sep;42(9):1408-11. doi: 10.1016/j.biocel.2010.06.003. Epub 2010 Jun 10. PMID: 20541619.