

PTPN22 Antibody (monoclonal) (M01)

Mouse monoclonal antibody raised against a full length recombinant PTPN22.

Catalog # AT3493a

Product Information

Application	WB, E
Primary Accession	Q9Y2R2
Other Accession	BC017785
Reactivity	Human
Host	mouse
Clonality	monoclonal
Isotype	IgG2a Kappa
Clone Names	4F6
Calculated MW	91705

Additional Information

Gene ID	26191
Other Names	Tyrosine-protein phosphatase non-receptor type 22, Hematopoietic cell protein-tyrosine phosphatase 70Z-PEP, Lymphoid phosphatase, LyP, PEST-domain phosphatase, PEP, PTPN22, PTPN8
Target/Specificity	PTPN22 (AAH17785, 1 a.a. ~ 179 a.a) full-length 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	PTPN22 Antibody (monoclonal) (M01) is for research use only and not for use in diagnostic or therapeutic procedures.

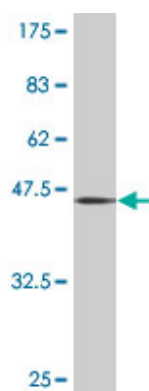
Background

This gene encodes a member of the non-receptor class 4 subfamily of the protein-tyrosine phosphatase family. The encoded protein is a lymphoid-specific intracellular phosphatase that associates with the molecular adapter protein CBL and may be involved in regulating CBL function in the T-cell receptor signaling pathway. Mutations in this gene may be associated with a range of autoimmune disorders including Type 1 Diabetes, rheumatoid arthritis, systemic lupus erythematosus and Graves' disease. Alternatively spliced transcript variants encoding distinct isoforms have been described.

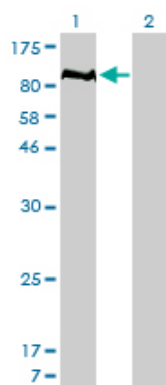
References

Latent autoimmune diabetes in adults differs genetically from classical type 1 diabetes diagnosed after the age of 35 years. Andersen MK, et al. *Diabetes Care*, 2010 Sep. PMID 20805278. Functional polymorphisms of PTPN22 and FcγR genes in Tunisian patients with rheumatoid arthritis. Sfar I, et al. *Arch Inst Pasteur Tunis*, 2009. PMID 20707220. PTPN22 polymorphism is related to autoimmune disease risk in patients with Turner syndrome. Bianco B, et al. *Scand J Immunol*, 2010 Sep. PMID 20696024. Variation at the NFATC2 Locus Increases the Risk of Thiazolidinedione-Induced Edema in the Diabetes REduction Assessment with ramipril and rosiglitazone Medication (DREAM) Study. Bailey SD, et al. *Diabetes Care*, 2010 Jul 13. PMID 20628086. Association of the protein tyrosine phosphatase nonreceptor 22 haplotypes with autoimmune thyroid disease in the Japanese population. Ban Y, et al. *Thyroid*, 2010 Aug. PMID 20615141.

Images

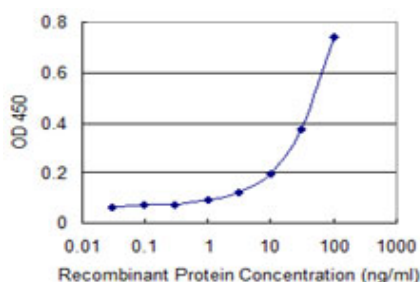


Antibody Reactive Against Recombinant Protein. Western Blot detection against Immunogen (45.43 KDa) .

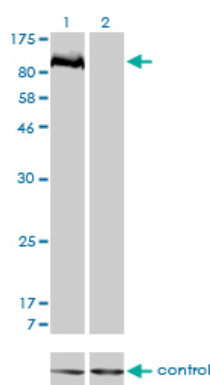


Western Blot analysis of PTPN22 expression in transfected 293T cell line by PTPN22 monoclonal antibody (M01), clone 4F6.

Lane 1: PTPN22 transfected lysate (85.1 KDa).
Lane 2: Non-transfected lysate.



Detection limit for recombinant GST tagged PTPN22 is 1 ng/ml as a capture antibody.



Western blot analysis of PTPN22 over-expressed 293 cell line, cotransfected with PTPN22 Validated Chimera RNAi (Cat # AT3493a)

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.