



Hard-to-Find Antibody

RFX5 DNAxPab

Catalog # H00005993-W01P

Size 200 ug

Specification	
Product Description	Rabbit polyclonal antibody raised against a full-length human RFX5 DNA using DNAx™ Immune tech nology.
Technology	DNAx™ Immune
Immunogen	Full-length human DNA
Sequence	MAEDEPDAKSPKTGGRAPPGGAEAGEPTTLLQRLRGTISKAVQNKVEGILQDVQKFSDNDKLYLY LQLPSGPTTGDKSSEPSTLSNEEYMYAYRWIRNHLEEHTDTCLPKQSVYDAYRKYCESLACCRPL STANFGKIIREIFPDIKARRLGGRGQSKYCYSGIRRKTLVSMPPLPGLDLKGSESPEMGPEVTPAPR DELVEAACALTCDWAERILKRSFSSIVEVARFLLQQHLISARSAHAHVLKAMGLAEEDEHAPRER SSKPKNGLENPEGGAHKKPERLAQPPKDLEARTGAGPLARGERKKSVVESSAPGANNLQVNAL VARLPLLLPRAPRSLIPPIPVSPPILAPRLSSGALKVATLPLSSRAGAPPAAVPIINMILPTVPALPGP GPGPGRAPPGGLTQPRGTENREVGIGGDQGPHDKGVKRTAEVPVSEASGQAPPAKAAKQDIED TASDAKRKRGRPRKKSGGSGERNSTPLKSAAAMESAQSSRLPWETWGSGGEGNSAGGAERP GPMGEAEKGAVLAQGQGDGTVSKGGRGPGSQHTKEAEDKIPLVPSKVSVIKGSRSQKEAFPLA KGEVDTAPQGNKDLKEHVLQSSLSQEHKDPKATPP
Host	Rabbit
Reactivity	Human
Purification	Protein A
Quality Control Testing	Antibody reactive against mammalian transfected lysate.
Storage Buffer	In 1x PBS, pH 7.4
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

Applications



- Western Blot (Transfected lysate)
 - Protocol Download
- Immunofluorescence (Transfected cell)
- Flow Cytometry (Transfected cell)

Gene Info — RFX5	
Entrez GenelD	<u>5993</u>
GeneBank Accession#	NM_000449.3
Protein Accession#	NP_000440.1
Gene Name	RFX5
Gene Alias	-
Gene Description	regulatory factor X, 5 (influences HLA class II expression)
Omim ID	209920 601863
Gene Ontology	<u>Hyperlink</u>
Gene Summary	A lack of MHC-II expression results in a severe immunodeficiency syndrome called MHC-II deficie ncy, or the bare lymphocyte syndrome (BLS; MIM 209920). At least 4 complementation groups ha ve been identified in B-cell lines established from patients with BLS. The molecular defects in complementation groups B, C, and D all lead to a deficiency in RFX, a nuclear protein complex that binds to the X box of MHC-II promoters. The lack of RFX binding activity in complementation group C results from mutations in the RFX5 gene encoding the 75-kD subunit of RFX (Steimle et al., 1 995). RFX5 is the fifth member of the growing family of DNA-binding proteins sharing a novel and highly characteristic DNA-binding domain called the RFX motif. Multiple alternatively spliced trans cript variants have been found but the full-length natures of only two have been determined. [provided by RefSeq
Other Designations	OTTHUMP00000082795 OTTHUMP00000196318 regulatory factor X, 5

Pathway

- Antigen processing and presentation
- Primary immunodeficiency



Disease

Macular Degeneration