

Product datasheet for PH312448

RFX5 (NM_001025603) Human Mass Spec Standard

Product data:

Product Type: Mass Spec Standards RFX5 MS Standard C13 and N15-labeled recombinant protein (NP_001020774) **Description:** Species: Human **HEK293 Expression Host:** Expression cDNA Clone RC212448 or AA Sequence: Predicted MW: 65.1 kDa >RC212448 representing NM_001025603 **Protein Sequence:** Red=Cloning site Green=Tags(s) MAEDEPDAKSPKTGGRAPPGGAEAGEPTTLLQRLRGTISKAVQNKVEGILQDVQKFSDNDKLYLYLQLPS GPTTGDKSSEPSTLSNEEYMYAYRWIRNHLEEHTDTCLPKQSVYDAYRKYCESLACCRPLSTANFGKIIR EIFPDIKARRLGGRGQSKYCYSGIRRKTLVSMPPLPGLDLKGSESPEMGPEVTPAPRDELVEAACALTCD WAERILKRSFSSIVEVARFLLQQHLISARSAHAHVLKAMGLAEEDEHAPRERSSKPKNGLENPEGGAHKK PERLAOPPKDLEARTGAGPLARGERKKSVVESSAPGANNLQVNALVARLPLLLPRAPRSLIPPIPVSPPI LAPRLSSGALKVATLPLSSRAGAPPAAVPIINMILPTVPALPGPGPGPGPGRAPPGGLTQPRGTENREVGIG GDQGPHDKGVKRTAEVPVSEASGQAPPAKAAKQDIEDTASDAKRKRGRPRKKSGGSGERNSTPLKSAAAM ESAQSSRLPWETWGSGGEGNSAGGAERPGPMGEAEKGAVLAQGQGDGTVSKGGRGPGSQHTKEAEDKIPL VPSKVSVIKGSRSQKEAFPLAKGEVDTAPQGNKDLKEHVLQSSLSQEHKDPKATPP TRTRPLEQKLISEEDLAANDILDYKDDDDKV Tag: C-Myc/DDK **Purity:** > 80% as determined by SDS-PAGE and Coomassie blue staining **Concentration:** >0.05 µg/µL as determined by microplate BCA method Labeling Method: Labeled with [U-13C6, 15N4]-L-Arginine and [U-13C6, 15N2]-L-Lysine Buffer: 25 mM Tris-HCl, 100 mM glycine, pH 7.3 Store at -80°C. Avoid repeated freeze-thaw cycles. Storage: Stability: Stable for 3 months from receipt of products under proper storage and handling conditions. **RefSeq:** NP 001020774 **RefSeq Size:** 3611 **RefSeq ORF:** 1848



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	RFX5 (NM_001025603) Human Mass Spec Standard – PH312448
Locus ID:	5993
UniProt ID:	<u>P48382</u>
Cytogenetics:	1q21.3
Summary:	A lack of MHC-II expression results in a severe immunodeficiency syndrome called MHC-II deficiency, or the bare lymphocyte syndrome (BLS; MIM 209920). At least 4 complementation groups have 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., 1995). 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 transcript variants have been found but the full-length natures of only two have been determined. [provided by RefSeq, Jul 2008]
Protein Families:	Transcription Factors
Protein Pathways	Antigen processing and presentation, Primary immunodeficiency
Product image	25:

116 — 66 — 45 — 35 — 25 — 18 — 14 —

Coomassie blue staining of purified RFX5 protein (Cat# [TP312448]). The protein was produced from HEK293T cells transfected with RFX5 cDNA clone (Cat# [RC212448]) using MegaTran 2.0 (Cat# [TT210002]).

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