

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten! See the following pages for more information!



Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

Zuschläge

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- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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RFX5 (Human) IP-WB Antibody Pair

Catalog #: H00005993-PW2 規格:[1 Set]

List All

Specification		Application Image
Product Description:	This IP-WB antibody pair set comes with one antibody for immunoprecipitation and another to detect the precipitated protein in western blot.	Immunoprecipitation-Westerr Blot
Reactivity:	Human	
Quality Control Testing:	Immunoprecipitation-Western Blot (IP-WB)	
	250 = 150 =	
	100-	
	75 - 50 -	
	37 -	
	25=	
	20 - Immunoprecipitation of RFX5 transfected lysate using rabbit polyclonal	
	anti-RFX5 and Protein A Magnetic Bead (<u>U0007</u>), and immunoblotted with mouse purified polyclonal anti-RFX5.	
Supplied Product:	Antibody pair set content: 1. Antibody pair for IP: rabbit polyclonal anti-RFX5 (300 ul) 2. Antibody pair for WB: mouse purified polyclonal anti-RFX5 (50 ug)	
Storage Instruction:	Store reagents of the antibody pair set at -20°C or lower. Please aliquot to avoid repeated freeze thaw cycle. Reagents should be returned to - 20°C storage immediately after use.	
MSDS:	<u>Download</u>	
Applications		
Immunoprecipit	tation-Western Blot	
Gene Information	on	
Entrez GeneID:	<u>5993</u>	
Gene Name:	RFX5	
Gene Alias:	-	
Gene Description:	regulatory factor X, 5 (influences HLA class II expression)	
Omim ID:	<u>209920, 601863</u>	
Gene Ontology:	: Hyperlink	

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Gene 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 fulllength natures of only two have been determined. [provided by RefSeq

Other

OTTHUMP00000082795,OTTHUMP00000196318,regulatory factor X, 5

Designations:

Gene Pathway

Antigen processing and presentation Primary immunodeficiency

Related Disease

Macular Degeneration

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