

# 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

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

#### SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

mail@szabo-scandic.com

www.szabo-scandic.com

linkedin.com/company/szaboscandic in





## BBS4 (Human) IP-WB Antibody Pair

**Catalog #**: H00000585-PW1 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-Western Blot
Reactivity:	Human	
Quality Control Testing:	250 =	
	150 <b>-</b> 100 <b>-</b> 75 <b>-</b> 50 <b>-</b>	
	37 <b>-</b> 25 <b>-</b>	
	20-	
	Immunoprecipitation of BBS4 transfected lysate using rabbit polyclonal anti-BBS4 and Protein A Magnetic Bead ( <u>U0007</u> ), and immunoblotted with mouse purified polyclonal anti-BBS4.	
Supplied Product:	Antibody pair set content:  1. Antibody pair for IP: rabbit polyclonal anti-BBS4 (300 ul)  2. Antibody pair for WB: mouse purified polyclonal anti-BBS4 (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	ration-Western Blot	
Gene Information	on	
Entrez GenelD:	<u>585</u>	
Gene Name:	BBS4	
Gene Alias:	-	
Gene Description:	Bardet-Biedl syndrome 4	
Omim ID:	209900, 600374	
Gene Ontology:	: Hyperlink	

Page 1 of 2 2016/5/20

Gene Summary: This gene is a member of the Bardet-Biedl syndrome (BBS) gene family. Bardet-Biedl syndrome is an autosomal recessive disorder characterized by severe pigmentary retinopathy, obesity, polydactyly, renal malformation and mental retardation. The proteins encoded by BBS gene family members are structurally diverse. The similar phenotypes exhibited by mutations in BBS gene family members are likely due to the protein's shared roles in cilia formation and function. Many BBS proteins localize to the basal bodies, ciliary axonemes, and pericentriolar regions of cells. BBS proteins may also be involved in intracellular trafficking via microtubule-related transport. The protein encoded by this gene has sequence similarity to O-linked N-acetylglucosamine (O-GlcNAc) transferases in plants and archaebacteria and in human forms a multi-protein "BBSome" complex with six other BBS proteins. Alternative splice variants have been described but their predicted protein products have not been

Other Designations:

#### **Related Disease**

Bardet-Biedl Syndrome Obesity Obesity Retinal Diseases Tobacco Use Disorder

experimentally verified

服務條款 | 隱私權政策 | 著作及商標 | 網站地圖

©2016 亞諾法生技股份有限公司 Abnova Corporation. 版權所有.

Page 2 of 2 2016/5/20