For Research Use Only

BBS3 Polyclonal antibody

Catalog Number:12676-1-AP

Featured Product

11 Publications



Basic Information	Catalog Number: 12676-1-AP	GenBank Accession Number: BC024239		Purification Method: Antigen affinity purification	
	Size:	GenelD (NCBI):		Recommended Dilutions:	
	150ul , Concentration: 400 ug/ml by	/ 84100		WB 1:500-1:2000 IP 0.5-4.0 ug for 1.0-3.0 mg of total protein lysate	
	Nanodrop and 393 ug/ml by Bradford UNIPROT ID: method using BSA as the standard; ODHOF 7				
	Source:	Q9H0F7		IHC 1:50-1:500	
	Rabbit	Full Name:		IF/ICC 1:20-1:200	
	lsotype:	Calculated MW:	T the O		
	IgG	186 aa, 21 kDa			
	Immunogen Catalog Number: AG3363	Observed MW: 21 kDa			
Applications	Tested Applications:	Positive Controls:			
				rain tissue, HEK-293T cells, human brain buse brain tissue	
	WB, IHC, IF		IP: rat brain t		
	Species Specificity:			ian kidney tissue,	
	human, mouse, rat, canine			T-RPE1 cells, MDCK cells	
	Cited Species: human, mouse				
	Note-IHC: suggested antigen ra TE buffer pH 9.0; (*) Alternativ retrieval may be performed w buffer pH 6.0	vely, antigen			
Background Information	defects in ARL6 are a cause of Bardet-	iliary organelle. ARL6 is Biedl syndrome type 3 (entary retinopathy, earl n. ARL6 can interact with	s also named BBS3), which i y onset obesi many ARL6 ir	Bardet-Biedl syndrome 3 protein, becau s a genetically heterogeneous disorder ty, polydactyly, hypogenitalism, renal	
	protein trafficking at the base of the c defects in ARL6 are a cause of Bardet- characterized by usually severe pigm malformation and mental retardatior complex with other BBS proteins inclu	iliary organelle. ARL6 is Biedl syndrome type 3 (entary retinopathy, earl n. ARL6 can interact with	s also named BBS3), which i y onset obesi many ARL6 ir and so on.	Bardet-Biedl syndrome 3 protein, becau s a genetically heterogeneous disorder ty, polydactyly, hypogenitalism, renal	
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Background Information Notable Publications	protein trafficking at the base of the c defects in ARL6 are a cause of Bardet- characterized by usually severe pigm malformation and mental retardation complex with other BBS proteins inclu Author Pub Ying Hsu 290	iliary organelle. ARL6 is Biedl syndrome type 3 (entary retinopathy, earl h. ARL6 can interact with Jding BBS1, BBS2, BBS4 a med ID Journa	s also named BBS3), which i y onset obesi many ARL6 ir and so on. I enet	Bardet-Biedl syndrome 3 protein, becau s a genetically heterogeneous disorder ty, polydactyly, hypogenitalism, renal teracting proteins and form BBSome Application	
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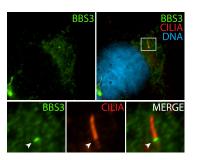
For technical support and original validation data for this product please contact:T: 1 (888) 4PTGLAB (1-888-478-4522) (toll free
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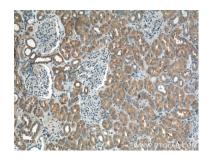
Selected Validation Data



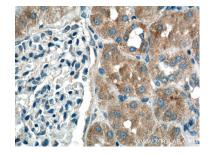
rat brain tissue were subjected to SDS PAGE followed by western blot with 12676-1-AP (BBS3 Antibody) at dilution of 1:1000 incubated at room temperature for 1.5 hours.



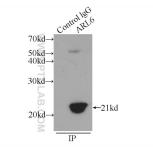
IF result (cytoplasm and the base of cilia stain) of anti-BBS3 (12676-1-AP; 1:50) with hTERT-RPE1 cell (MeOH fixed) by Dr. Moshe Kim.



Immunohistochemical analysis of paraffinembedded human kidney tissue slide using 12676-1-AP (BBS3 Antibody) at dilution of 1:200 (under 10x lens).



Immunohistochemical analysis of paraffinembedded human kidney tissue slide using 12676-1-AP (BBS3 Antibody) at dilution of 1:200 (under 40x lens).



IP result of anti-BBS3 (IP:12676-1-AP, 3ug; Detection:12676-1-AP 1:600) with rat brain tissue lysate 5200ug.