For Research Use Only

FOXL2-Specific Polyclonal antibody

Catalog Number:19672-1-AP

Featured Product 14 Publications

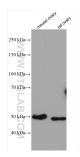
Antibodies | ELISA kits | Proteins www.ptglab.com

Basic Information	Catalog Number: 19672-1-AP	GenBank Accession Number: NM_023067	Purification Method: Antigen affinity purification	
	Size:	GenelD (NCBI):	Recommended Dilutions:	
	150ul , Concentration: 260 ug/ml by	668	WB 1:500-1:2000	
	Nanodrop and 233 ug/ml by Bradford	UNIPROT ID:		
	method using BSA as the standard;	P58012		
	Source:	Full Name:		
	Rabbit	forkhead box L2		
	Isotype: IgG	Calculated MW: 39 kDa		
		Observed MW:		
		49 kDa		
Applications	Tested Applications:	Positive Controls:		
	WB, ELISA	WB : mouse ovary tissue, rat ovary tissue		
	Cited Applications:			
	WB, IHC, IF			
	Species Specificity: human, mouse, rat			
	Cited Species:			
	human, mouse, rat, pig, chicken			
Background Information	FOXL2 is probable a transcriptional re	egulator. It is required to maintain	the female phenotype throughout adulth	
Background Information	and may provide clues to remate mile	rtility. Defects in FOXL2 are a caus Defects in FOXL2 are a cause of p	e of blepharophimosis, ptosis, and	
	epicanthus inversus syndrome (BPES) antibody is specific to the C-term of F	rtility. Defects in FOXL2 are a caus Defects in FOXL2 are a cause of p	e of blepharophimosis, ptosis, and	
	epicanthus inversus syndrome (BPES) antibody is specific to the C-term of F Author Put	rtility. Defects in FOXL2 are a caus Defects in FOXL2 are a cause of pr OXL2.	se of blepharophimosis, ptosis, and remature ovarian failure type 3 (POF3). Th	
	and may provide clues to remate mile epicanthus inversus syndrome (BPES) antibody is specific to the C-term of F Author Put Xian Zhang 346	rtility. Defects in FOXL2 are a caus Defects in FOXL2 are a cause of pr OXL2.	remature ovarian failure type 3 (POF3). The second se	
	Author Put Xian Zhang 344 Siwen Zhang 337	rtility. Defects in FOXL2 are a caus Defects in FOXL2 are a cause of pr OXL2. Demed ID Journal 500579 Chin Med	e of blepharophinosis, ptosis, and remature ovarian failure type 3 (POF3). The Application IF	
	Author Put Xian Zhang 344 Siwen Zhang 337	rtility. Defects in FOXL2 are a cause Defects in FOXL2 are a cause of pr OXL2. med ID Journal 500579 Chin Med 712079 Stem Cell Res Th	e of blepharophimosis, ptosis, and remature ovarian failure type 3 (POF3). The Application IF er IF,IHC	
Notable Publications	Author Put Xian Zhang 344 Siwen Zhang 337	rtility. Defects in FOXL2 are a cause Defects in FOXL2 are a cause of pr OXL2. med ID Journal 500579 Chin Med 712079 Stem Cell Res Th 119367 Clin Cancer Res er shipment.	e of blepharophimosis, ptosis, and remature ovarian failure type 3 (POF3). T Application IF er IF,IHC	
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Background Information Notable Publications Storage	Author Put Xian Zhang 344 Siwen Zhang 333 Jiaqiang Dong 28: Storage: Storage Buffer: PBS with 0.02% sodium azide and 50	rtility. Defects in FOXL2 are a cause Defects in FOXL2 are a cause of pr OXL2. omed ID Journal 500579 Chin Med 712079 Stem Cell Res Th 119367 Clin Cancer Res er shipment. % glycerol	e of blepharophimosis, ptosis, and remature ovarian failure type 3 (POF3). The Application IF er IF,IHC	
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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 freeE: proteintech@ptglab.comin USA), or 1(312) 455-8498 (outside USA)W: ptglab.com

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Selected Validation Data



Various lysates were subjected to SDS PAGE followed by western blot with 19672-1-AP (FOXL2-Specific antibody) at dilution of 1:1000 incubated at room temperature for 1.5 hours.