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## AR Polyclonal antibody Catalog Number: 19783-1-AP 1 Publications

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	Catalog Number: GenBank Accession Number: 19783-1-AP NM_000044		Purification Method: Antigen affinity purification	
	Concentration: GeneID (NCBI):			······8-·······
	N/A			
	Source:UNIPROTID:RabbitP10275			
	Isotype:Full Name:IgGandrogen receptor			
Applications	Tested Applications: WB, ELISA			
	Cited Applications: WB, IHC			
	Species Specificity: human, mouse, rat			
	Cited Species: rat			
Background Information	AR, also named as DHTR and NR3C 4, belongs to the nuclear hormone receptor family and NR3 subfamily. AR is a ligand-activated transcription factors that regulate eukaryotic gene expression and affect cellular proliferation and differentiation in target tissues. Transcription factor activity is modulated by bound coactivator and corepressor proteins. AR is a activated, but not phosphorylated, by HIPK3. Defects in AR are the cause of androgen insensitivity syndrome (AIS), previously known as testicular feminization syndrome (TFM), which is an X-linked recessive form or pseudohermaphroditism due end-organ resistance to androgen. Defects in AR are the cause of spinal and bulbar muscular atrophy X-linked type 1 (SMAX1) which also known as Kenendy disease. Defects in AR may play a role in metastatic prostate cancer. Defects in AR are the cause of androgen insensitivity syndrome partial (PAIS) which also known as Reifenstein syndrome. This antibody is a rabbit polyclonal antibody raised against a peptide mapping within human AR.			
	known as Reifenstein syndr	ome. This antibody is a ra	abbit polyclonal a	
Notable Publications	known as Reifenstein syndr	ome. This antibody is a ra	abbit polyclonal a	

For technical support and original validation data for this product please contact:T: 4006900926E: Proteintech-CN@ptglab.comW: ptgcn.com

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