## bs-12549R

## [ Primary Antibody ]

## ATP6V1B2 Rabbit pAb



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- DATASHEET		400-90	01-9800	
Host: Rabbit	lsotype: IgG	Applications:	<b>WB</b> (1:500-2000)	
Clonality: Polyclonal	-		<b>IHC-P</b> (1:100-500) <b>IHC-F</b> (1:100-500)	
<b>GeneID:</b> 526	SWISS: P21281		IF (1:100-500)	
Target: ATP6V1B2			ICC/IF (1:100-500) FLISA (1:5000-10000)	
Immunogen: KLH conjugated synthetic peptide derived from human ATP6V1B2: 51-150/511.		: Reactivity:	<b>Reactivity:</b> (predicted: Human, Mouse, Rat, Rabbit, Pig, Sheep, Cow, Chickon, Dog, Horso	
Purification: affinity purified by Protein A				
Concentration: 1mg/ml			Cynomolgus Monkey,	
<b>Storage:</b> 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol. Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.		Predicted MW.:	Orangutan) Predicted MW.: <sup>56 kDa</sup>	
<b>Background:</b> Vacuolar-type H+-ATPase (V-ATPase) is a multisubunit enzyme responsible for acidification of eukaryotic intracellular organelles. V-ATPases pump protons against an electrochemical gradient, while F-ATPases reverse the process, thereby synthesizing ATP. A peripheral V1 domain, which is responsible for ATP hydrolysis, and a integral V0 domain, which is responsible for proton translocation, compose V-ATPase. Nine subunits (A–H) make up the V1 domain and five subunits (a, d, c, c' and c") make up the V0 domain. Like F-ATPase, V-ATPase most likely operates through a rotary mechanism. The V-ATPase B1 isoform functions in proton secretion and is required to maintain proper endolymph pH and normal auditory function. The gene encoding the human V-ATPase B1 isoform maps to chromosome 2cen-q13. Mutations in this gene cause distal renal tubular acidosis associated with sensorineural deafness. The V-ATPase B2 isoform is expressed in kidney and is the only B isoform expressed in osteoclasts. The gene encoding the human V-ATPase B2 isoform maps to chromosome 8p22-p21.		e e	Cell membrane	