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anti-CSNK1G1 antibody (C-Term)

2 Images



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Overview		
Quantity:	400 μL	
Target:	CSNK1G1	
Binding Specificity:	AA 359-387, C-Term	
Reactivity:	Mouse	
Host:	Rabbit	
Clonality:	Polyclonal	
Conjugate:	This CSNK1G1 antibody is un-conjugated	
Application:	Western Blotting (WB), Immunohistochemistry (Paraffin-embedded Sections) (IHC (p))	
Product Details		
Immunogen:	This Mouse Csnk1g1 antibody is generated from rabbits immunized with a KLH conjugated	
	synthetic peptide between 359-387 amino acids from the C-terminal region of mouse Csnk1g1.	
Isotype:	lg Fraction	
Purification:	This antibody is purified through a protein A column, followed by peptide affinity purification.	
Target Details		
Target:	CSNK1G1	
Alternative Name:	Csnk1g1 (CSNK1G1 Products)	
Background:	Casein kinases are operationally defined by their preferential utilization of acidic proteins such	
	as caseins as substrates. It can phosphorylate a large number of proteins. Participates in Wnt signaling (By similarity).	
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Target Details

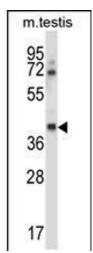
Molecular Weight:	53 kDa
Gene ID:	214897
UniProt:	Q8BTH8
Pathways:	Hedgehog Signaling

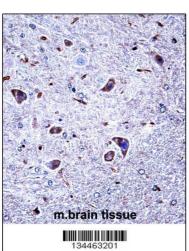
Application Details

Application Notes:	For WB starting dilution is: 1:1000
	For IHC-P starting dilution is: 1:10~50
Restrictions:	For Research Use only

Handling

Format:	Liquid
Concentration:	0.5 mg/mL
Buffer:	Supplied in PBS with 0.09 % (W/V) sodium azide.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	4 °C,-20 °C
Storage Comment:	Store at 4°C for three months and -20°C, stable for up to one year. As with all antibodies care should be taken to avoid repeated freeze thaw cycles. Antibodies should not be exposed to prolonged high temperatures.





Western Blotting

Image 1. Western blot analysis in mouse testis tissue lysates (35ug/lane).

Immunohistochemistry

Image 2. Mouse Csnk1g1 Antibody immunohistochemistry analysis in formalin fixed and paraffin embedded mouse brain tissue followed by peroxidase conjugation of the secondary antibody and DAB staining.