

## **Product Data Sheet**

## **Anti-SHCB Antibody**

Catalog #	Source	Reactivity	Applications		
CPA2592	Rabbit	H, M, R, Mk	WB, IH		
Description	Rat	obit polyclonal antibody t	o SHCB		
Immunogen	KLF	I-conjugated synthetic pe	ptide encompassing a sequence within the center		
	reg	ion of human SHCB. The	exact sequence is proprietary.		
Purification	The	antibody was purified b	y immunogen affinity chromatography.		
Specificity	Rec	cognizes endogenous leve	els of SHCB protein.		
Clonality	Pol	Polyclonal			
Conjugation					
Form	Liq	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	and	d 0.01% sodium azide.			
Dilution	WB	8 (1/500 - 1/1000), IH (1/50	- 1/100)		
Gene Symbol	SHO	22			
Alternative Na	imes SCk	<; SHCB; SHC-transformin	g protein 2; Protein Sck; SHC-transforming protein B; Src		
	hor	nology 2 domain-contain	ing-transforming protein C2; SH2 domain protein C2		
Entrez Gene	257	759 (Human); 216148 (Mo	ouse); 314612 (Rat)		
SwissProt	P98	3077 (Human); Q8BMC3 (	(Mouse); 070142 (Rat)		
Storage/Stabil	l <mark>ity</mark> Shi	pped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	free	eze/thaw cycles.			

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

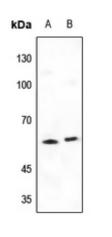
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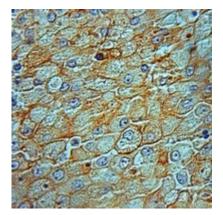
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Western blot analysis of SHCB expression in mouse testis (A), rat testis (B) whole cell lysates. (Predicted band size: 61 kD; Observed band size: 59 kD)



Immunohistochemical analysis of SHCB staining in human prostate formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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