

Product Data Sheet

Anti-ICER Antibody

Catalog #	Source	Reactivity	Applications
CPA3875	Rabbit	H, M, R, B, D, P	WB, IH
Description	F	Rabbit polyclonal antibody to	DICER
Immunogen	K	Characteristic Contracteristic Contracteris	otide encompassing a sequence within the center
	r	egion of human ICER. The e	act sequence is proprietary.
Purification	Т	The antibody was purified by	immunogen affinity chromatography.
Specificity	F	Recognizes endogenous leve	ls of ICER protein.
Clonality	P	Polyclonal	
Conjugation			
Form	L	iquid in 0.42% Potassium pl	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	а	and 0.01% sodium azide.	
Dilution	V	WB (1/500 - 1/1000), IH (1/10) - 1/200)
Gene Symbol	C	CREM	
Alternative Na	a <mark>mes</mark> c	CAMP-responsive element m	odulator; Inducible cAMP early repressor; ICER
Entrez Gene	1	1390 (Human); 12916 (Mous	e); 25620 (Rat)
SwissProt	C	Q03060 (Human); P27699 (N	louse); Q03061 (Rat)
Storage/Stabi	lity S	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid
	f	reeze/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

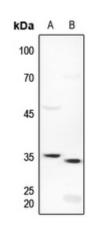
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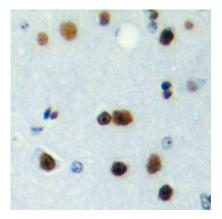
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Western blot analysis of ICER expression in mouse kidney (A), rat lung (B) whole cell lysates. (Predicted band size: 37 kD; Observed band size: 35 kD)



Immunohistochemical analysis of ICER staining in human brain 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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