

Product Data Sheet

Anti-TECR Antibody

Catalog #	Source	Reactivity	Applications		
CPA7228	Rabbit	H, M, R, B	WB, IH		
Description		Rabbit polyclonal antibody to	D TECR		
Immunogen		KLH-conjugated synthetic pe	ptide encompassing a sequence within the C-term		
		region of human TECR. The e	xact sequence is proprietary.		
Purification		The antibody was purified by	immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	ls of TECR protein.		
Clonality		Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
		and 0.01% sodium azide.			
Dilution		WB (1/500 - 1/1000), IH (1/10	0 - 1/200)		
Gene Symbol		TECR			
Alternative Na	ames	GPSN2; SC2; Very-long-chain	enoyl-CoA reductase; Synaptic glycoprotein SC2;		
		Trans-2,3-enoyl-CoA reducta	se; TER		
Entrez Gene		9524 (Human); 106529 (Mou	ise); 191576 (Rat)		
SwissProt		Q9NZ01 (Human); Q9CY27 (I	Mouse); Q64232 (Rat)		
Storage/Stabi	lity	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
		freeze/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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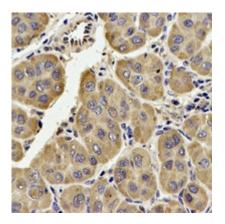
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140

For research purposes only, not for human use

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Western blot analysis of TECR expression in HEK293T (A) whole cell lysates. (Predicted band size: 36 kD; Observed band size: 35 kD)



Immunohistochemical analysis of TECR staining in human liver 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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