

Product Data Sheet

Anti-TBP Antibody

Catalog #	Source	Reactivity	Applications
CPA4272	Rabbit	H, M, R, B, C, P, Rb, Z	WB, IH, IF/IC
Description	Ra	abbit polyclonal antibody to	ТВР
Immunogen	KL	H-conjugated synthetic pep	tide encompassing a sequence within the center
	re	gion of human TBP. The exac	ct sequence is proprietary.
Purification	Th	ne antibody was purified by i	immunogen affinity chromatography.
Specificity	Re	ecognizes endogenous levels	s of TBP protein.
Clonality	Рс	olyclonal	
Form	Lic	quid in 0.42% Potassium pho	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	an	nd 0.01% sodium azide.	
Dilution	W	'B (1/500 - 1/1000), IH (1/100	- 1/200), IF/IC (1/100 - 1/500)
Gene Symbol	ТВ	3P	
Alternative N	ames G1	FF2D1; TF2D; TFIID; TATA-bo	x-binding protein; TATA sequence-binding protein;
	TA	ATA-binding factor; TATA-box	factor; Transcription initiation factor TFIID TBP subunit
Entrez Gene	69	908 (Human); 21374 (Mouse)
SwissProt	P2	20226 (Human); P29037 (Mo	ouse)
Storage/Stabi	lity Sh	nipped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid
	fre	eeze/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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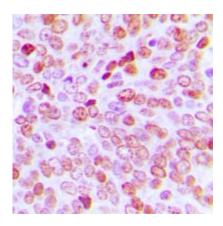
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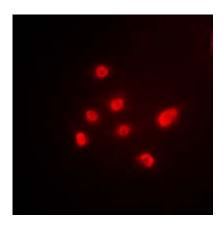
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Western blot analysis of TBP expression in HEK293T (A), NIH3T3 (B) whole cell lysates.



Immunohistochemical analysis of TBP staining in human prostate cancer 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 conjugad compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of TBP staining in HEK293T cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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