

## **Product Data Sheet**

### **Anti-TXNRD2** Antibody

Catalog #	Source	Reactivity	Applications			
CPA2495	Rabbit	H, M, R, Mk	WB, IH			
Description	R	Rabbit polyclonal antibody to TXNRD2				
Immunogen	K	(LH-conjugated synthetic pep	tide encompassing a sequence within the C-term			
	re	egion of human TXNRD2. The	e exact sequence is proprietary.			
Purification	Т	he antibody was purified by	immunogen affinity chromatography.			
Specificity	R	ecognizes endogenous level	s of TXNRD2 protein.			
Clonality	Clonality Polyclonal					
Conjugation						
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,				
	а	nd 0.01% sodium azide.				
Dilution	V	VB (1/500 - 1/1000), IH (1/100	- 1/200)			
Gene Symbol	Т	XNRD2				
Alternative Na	ames K	(IAA1652; TRXR2; Thioredoxi	n reductase 2 mitochondrial; Selenoprotein Z; SelZ;			
	Т	R-beta; Thioredoxin reducta	se TR3			
Entrez Gene	1	.0587 (Human); 50551 (Rat)				
SwissProt	C	Q9NNW7 (Human); Q9Z0J5 (F	Rat)			
Storage/Stabi	lity S	hipped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid			
	fr	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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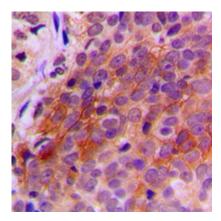


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kDa A B C D E F 130 100 70 45 35 25 20

Western blot analysis of TXNRD2 expression in HEK293T (A), HGC27 (B), mouse lung (C), mouse liver (D), rat lung (E), rat liver (F) whole cell lysates. (Predicted band size: 56 kD; Observed band size: 56 kD)



Immunohistochemical analysis of TXNRD2 staining in human breast 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 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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