

## **Product Data Sheet**

### Anti-TP53INP1 Antibody

Catalog #	Source	Reactivity	Applications		
CPA2838	Rabbit	H, Mk	WB, IH		
Description		Rabbit polyclonal antibody to TP53INP1			
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the C-term			
	l	region of human TP53INP1.	The exact sequence is proprietary.		
Purification		The antibody was purified by immunogen affinity chromatography.			
Specificity		Recognizes endogenous leve	ls of TP53INP1 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 (1/50 - 1/100)			
Gene Symbol		TP53INP1			
Alternative Names		P53DINP1; SIP; Tumor protein p53-inducible nuclear protein 1; Stress-induced			
		protein; p53-dependent dan	nage-inducible nuclear protein 1; p53DINP1		
Entrez Gene		94241 (Human)			
SwissProt		Q96A56 (Human)			
Storage/Stabil	/Stability Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid				
	Ī	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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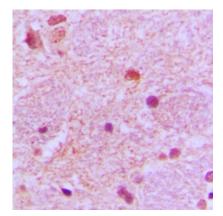
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Western blot analysis of TP53INP1 expression in Hela (A), HepG2 (B) whole cell lysates. (Predicted band size: 27 kD; Observed band size: 27 kD)



Immunohistochemical analysis of TP53INP1 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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