

# **Product Data Sheet**

## **Anti-ETV4 Antibody**

Catalog #	Source	Reactivity	Applications		
CPA3931	Rabbit	H, M, R, D	WB, IH		
Description		Rabbit polyclonal antibody to ETV4			
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the center			
		region of human ETV4. The ε	xact sequence is proprietary.		
Purification		The antibody was purified by immunogen affinity chromatography.			
Specificity		Recognizes endogenous leve	ls of ETV4 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	) - 1/200)		
Gene Symbol		ETV4			
Alternative Na	mes	E1AF; PEA3; ETS translocatio	n variant 4; Adenovirus E1A enhancer-binding protein;		
		E1A-F; Polyomavirus enhanc	er activator 3 homolog; Protein PEA3		
Entrez Gene		2118 (Human); 18612 (Mous	e)		
SwissProt		P43268 (Human); P28322 (N	ouse)		
Storage/Stabili	ity	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
		freeze/thaw cycles.			

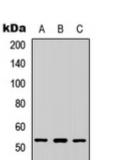
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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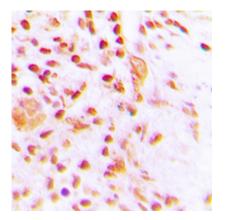
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For research purposes only, not for human use

## **Product Data Sheet**

Western blot analysis of ETV4 expression in HeLa (A), Raw264.7 (B), PC12 (C) whole cell lysates. (Predicted band size: 53 kD; Observed band size: 54 kD)



Immunohistochemical analysis of ETV4 staining in human lung 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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