

**Anti-APC (Phospho-S2054) Antibody**

Catalog #	Source	Reactivity	Applications
CPA1056	Rabbit	H, M, R, Mk, P	WB, IH
<b>Description</b>	Rabbit polyclonal antibody to APC (Phospho-S2054)		
<b>Immunogen</b>	KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding S2054 of human APC protein. The exact sequence is proprietary.		
<b>Purification</b>	The antibody was purified by immunogen affinity chromatography.		
<b>Specificity</b>	Recognizes endogenous levels of APC protein only when phosphorylated at S2054.		
<b>Clonality</b>	Polyclonal		
<b>Conjugation</b>			
<b>Form</b>	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide.		
<b>Dilution</b>	WB (1/500 - 1/1000), IH (1/50 - 1/100)		
<b>Gene Symbol</b>	APC		
<b>Alternative Names</b>	DP2.5; Adenomatous polyposis coli protein; Protein APC; Deleted in polyposis 2.5		
<b>Entrez Gene</b>	324 (Human); 24205 (Rat)		
<b>SwissProt</b>	P25054 (Human); Q61315 (Mouse); P70478 (Rat)		
<b>Storage/Stability</b>	Shipped at 4°C. Upon delivery 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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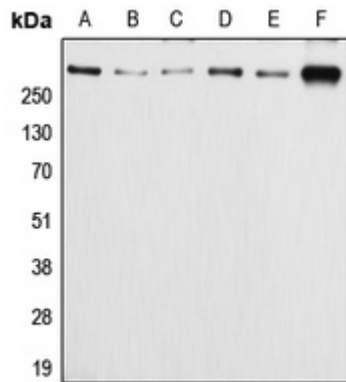
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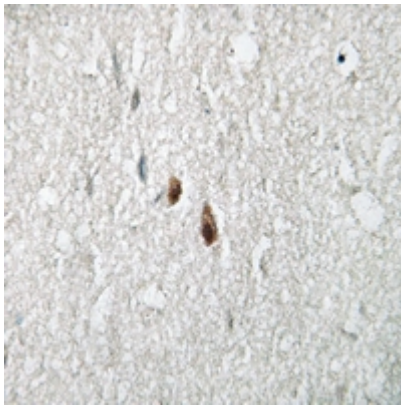
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## Product Data Sheet



Western blot analysis of APC (Phospho-S2054) expression in HeLa (A), Caco2 (B), SW480 (C), MCF7 (D), SP2/0 (E), PC12 (F) whole cell lysates. (Predicted band size: 311 kD; Observed band size: 310 kD)



Immunohistochemical analysis of APC (Phospho-S2054) 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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