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## anti-RAD17 antibody (C-Term)

**Images** 



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Quantity:	100 μL	
Target:	RAD17	
Binding Specificity:	C-Term	
Reactivity:	Human	
Host:	Rabbit	
Clonality:	Polyclonal	
Conjugate:	This RAD17 antibody is un-conjugated	
Application:	Western Blotting (WB), Immunohistochemistry (IHC), Immunofluorescence (IF), Immunochromatography (IC)	

#### **Product Details**

Immunogen:	KLH-conjugated synthetic peptide encompassing a sequence within the C-term region of human RAD17.
Specificity:	Recognizes endogenous levels of RAD17 protein.
Characteristics:	Rabbit polyclonal antibody to RAD17
Purification:	The antibody was purified by immunogen affinity chromatography.

## Target Details

Target:	RAD17
Alternative Name:	RAD17 (RAD17 Products)

### **Target Details**

Background:	R24L, Cell cycle checkpoint protein RAD17, hRad17, RF-C/activator 1 homolog	
Gene ID:	5884	
UniProt:	075943	

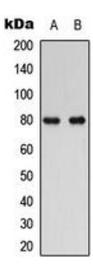
## **Application Details**

Application Notes:	WB (1:500 - 1:1000), IH (1:100 - 1:200), IF/IC (1:100 - 1:500)
Restrictions:	For Research Use only

## Handling

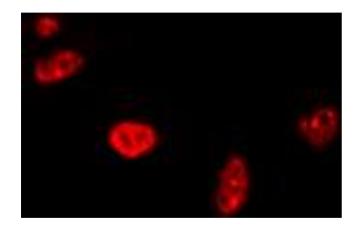
Format:	Liquid
Buffer:	Liquid in 0.42 % Potassium phosphate, 0.87 % Sodium chloride, pH 7.3, 30 % glycerol, and 0.01 % sodium azide.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C
Storage Comment:	Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid freeze/thaw cycles.
Expiry Date:	12 months

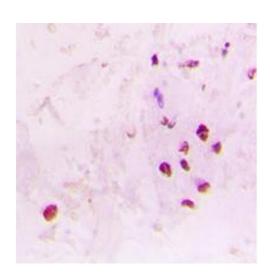
### **Images**



## **Western Blotting**

Image 1. Western blot analysis of RAD17 expression in human Oral cancer (A), HuvEc (B) whole cell lysates.





#### **Immunofluorescence**

Image 2. Immunofluorescent analysis of RAD17 staining in A549 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. DAPI was used to stain the cell nuclei (blue).

#### **Immunohistochemistry**

Image 3. Immunohistochemical analysis of RAD17 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 conjugad compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.