## **TP63 Antibody**

Catalog No: #32619

Signalway Antibody

Olgitalita) / Ilitaboa)

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## Package Size: #32619-1 50ul #32619-2 100ul

Description		
Product Name	TP63 Antibody	
Host Species	Rabbit	
Clonality	Polyclonal	
Isotype	IgG	
Purification	Affinity purification	
Applications	WB,IHC	
Species Reactivity	Human,Mouse,Rat	
Specificity	The antibody detects endogenous level of total TP63 protein.	
Immunogen Type	Recombinant Protein	
Immunogen Description	Recombinant fusion protein of human p63 (NP_003713.3).	
Target Name	TP63	

	73H;p73L;TP63
Accession No.	Uniprot:Q9H3D4GeneID:8626

Uniprot	Q9H3D4
GeneID	8626
SDS-PAGE MW	75kDa

Concentration	1.0mg/ml	

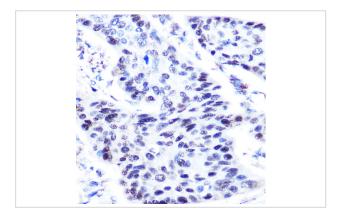
Formulation	PBS with 0.02% sodium azide,50% glycerol,pH7
Storage	Store at -20°C. Avoid freeze / thaw cycles.

# Application Details

WB□1:500 - 1:2000IHC□1:50 - 1:200

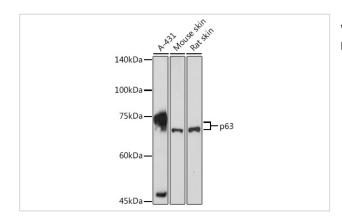
### **Images**

Other Names



Immunohistochemistry of paraffin-embedded human esophageal cancer using p63 antibody.

AIS;B(p51A);B(p51B);EEC3;KET;LMS;NBP;OFC8;RHS;SHFM4;TP53CP;TP53L;TP73L;p40;p51;p53CP;p63;p



Western blot analysis of extracts of various cell lines, using p63 antibody.

#### Background

This gene encodes a member of the p53 family of transcription factors. The functional domains of p53 family proteins include an N-terminal transactivation domain, a central DNA-binding domain and an oligomerization domain. Alternative splicing of this gene and the use of alternative promoters results in multiple transcript variants encoding different isoforms that vary in their functional properties. These isoforms function during skin development and maintenance, adult stem/progenitor cell regulation, heart development and premature aging. Some isoforms have been found to protect the germline by eliminating oocytes or testicular germ cells that have suffered DNA damage. Mutations in this gene are associated with ectodermal dysplasia, and cleft lip/palate syndrome 3 (EEC3); split-hand/foot malformation 4 (SHFM4); ankyloblepharon-ectodermal defects-cleft lip/palate; ADULT syndrome (acro-dermato-ungual-lacrimal-tooth); limb-mammary syndrome; Rap-Hodgkin syndrome (RHS); and orofacial cleft 8.

Note: This product is for in vitro research use only