

**p73 (Phospho Thr86) Rabbit pAb**

<b>Catalog No :</b>	YP1850
<b>Reactivity :</b>	Human;Mouse
<b>Applications :</b>	IHC;WB
<b>Target :</b>	p73
<b>Gene Name :</b>	TP73 P73
<b>Protein Name :</b>	Tumor protein p73 (p53-like transcription factor) (p53-related protein)
<b>Sequence :</b>	O15350
<b>Human Gene Id :</b>	7161
<b>Human Swiss Prot No :</b>	O15350
<b>Mouse Gene Id :</b>	22062
<b>Mouse Swiss Prot No :</b>	Q9JJP2
<b>Immunogen :</b>	Synthesized peptide derived from human p73 (Phospho Thr86)
<b>Specificity :</b>	This antibody detects endogenous levels of p73 (Phospho Thr86) Rabbit pAb at Human, Mouse
<b>Formulation :</b>	Liquid in PBS containing 50% glycerol, and 0.02% sodium azide.
<b>Source :</b>	Rabbit,polyclonal
<b>Dilution :</b>	WB 1:500-2000 IHC 1:50-200
<b>Purification :</b>	The antibody was affinity-purified from rabbit serum by affinity-chromatography using specific immunogen.
<b>Concentration :</b>	1 mg/ml

**Storage Stability :** -15°C to -25°C/1 year(Do not lower than -25°C)

**Observed Band :** 70kD

**Background :** tumor protein p73(TP73) Homo sapiens This gene encodes a member of the p53 family of transcription factors involved in cellular responses to stress and development. It maps to a region on chromosome 1p36 that is frequently deleted in neuroblastoma and other tumors, and thought to contain multiple tumor suppressor genes. The demonstration that this gene is monoallelically expressed (likely from the maternal allele), supports the notion that it is a candidate gene for neuroblastoma. Many transcript variants resulting from alternative splicing and/or use of alternate promoters have been found for this gene, but the biological validity and the full-length nature of some variants have not been determined. [provided by RefSeq, Feb 2011],

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**Function :** cofactor: Binds 1 zinc ion per subunit., disease: Maps to a chromosome region frequently mutated in diverse cell lines of human cancer. Appears not to be frequently mutated in human cancers, in contrast to p53. Hemizygoty is observed in neuroblastoma and oligodendroglioma., domain: Possesses an acidic transactivation domain, a central DNA binding domain and a C-terminal oligomerization domain that binds to the ABL tyrosine kinase SH3 domain., domain: The WW-binding motif mediates interaction with WWOX., function: Participates in the apoptotic response to DNA damage. Isoforms containing the transactivation domain are pro-apoptotic, isoforms lacking the domain are anti-apoptotic and block the function of p53 and transactivating p73 isoforms. May be a tumor suppressor protein., induction: Not induced by DNA damage. Isoforms lacking the transactivation domain block gene induction., miscellaneous: Activ

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**Subcellular Location :** Nucleus . Cytoplasm. Accumulates in the nucleus in response to DNA damage.

**Expression :** Expressed in striatal neurons of patients with Huntington disease (at protein level). Brain, kidney, placenta, colon, heart, liver, spleen, skeletal muscle, prostate, thymus and pancreas. Highly expressed in fetal tissue. Expressed in the respiratory epithelium (PubMed:34077761).

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**Sort :** 999

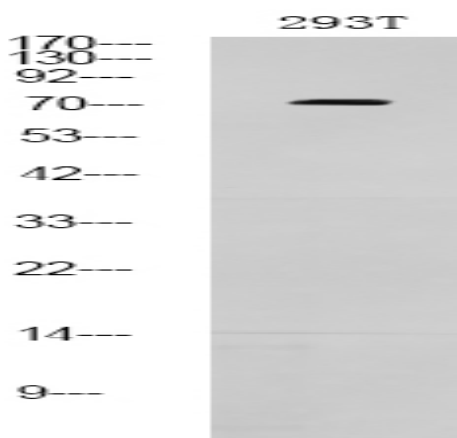
**No4 :** 1

**Host :** Rabbit

**Modifications :** Phospho

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## Products Images



Western Blot analysis of 293T using primary antibody at 1:1000 dilution 4 °C, overnight. Secondary antibody(catalog#:RS23920) was diluted at 1:10000 25 °C 1.5hours