

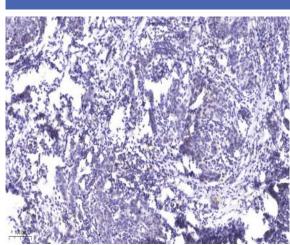
Rb (Phospho Ser807/811) rabbit pAb

Catalog No :	YP1462
Reactivity :	Human;Mouse;Rat
Applications :	WB;IHC
Target :	Rb
Fields :	>>Endocrine resistance;>>Cell cycle;>>Cellular senescence;>>Cushing syndrome;>>Hepatitis C;>>Hepatitis B;>>Human cytomegalovirus infection;>>Human papillomavirus infection;>>Human T-cell leukemia virus 1 infection;>>Kaposi sarcoma-associated herpesvirus infection;>>Epstein-Barr virus infection;>>Pathways in cancer;>>Viral carcinogenesis;>>Chemical carcinogenesis - receptor activation;>>Pancreatic cancer;>>Glioma;>>Prostate cancer;>>Melanoma;>>Bladder cancer;>>Breast cancer;>>Hepatocellular carcinoma;>>Gastric cancer
Gene Name :	RB1
Protein Name :	Rb (Ser807/811)
Human Gene Id :	5925
Human Swiss Prot	P06400
No : Mouse Gene Id :	19645
Mouse Swiss Prot	P13405
No : Rat Gene Id :	24708
Rat Swiss Prot No :	P33568
Immunogen :	Synthesized phosho peptide around human Rb (Ser807 and 811)
Specificity :	This antibody detects endogenous levels of Human Mouse Rat Rb (phospho- Ser807 or 811)



Best Tools for immunolog	gy Research
Formulation :	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source :	Polyclonal, Rabbit,IgG
Dilution :	WB 1:500-2000;IHC 1:50-300
Purification :	The antibody was affinity-purified from rabbit serum by affinity-chromatography
	using specific immunogen.
Concentration :	1 mg/ml
Storage Stability :	-15°C to -25°C/1 year(Do not lower than -25°C)
Observed Band :	106kD
Cell Pathway :	Stem cell pathway; Cell_Cycle_G1S;Cell_Cycle_G2M_DNA;
	Protein_Acetylation
Background :	The protein encoded by this gene is a negative regulator of the cell cycle and
	was the first tumor suppressor gene found. The encoded protein also stabilizes
	constitutive heterochromatin to maintain the overall chromatin structure. The active, hypophosphorylated form of the protein binds transcription factor E2F1.
	Defects in this gene are a cause of childhood cancer retinoblastoma (RB),
	bladder cancer, and osteogenic sarcoma. [provided by RefSeq, Jul 2008],
Function :	disease:Defects in RB1 are a cause of bladder cancer
	[MIM:109800].,disease:Defects in RB1 are a cause of osteogenic sarcoma
	[MIM:259500]., disease: Defects in RB1 are the cause of childhood cancer
	retinoblastoma (RB) [MIM:180200]. RB is a congenital malignant tumor that
	arises from the nuclear layers of the retina. It occurs in about 1:20'000 live births and represents about 2% of childhood malignancies. It is bilateral in about 30% of
	cases. Although most RB appear sporadically, about 20% are transmitted as an
	autosomal dominant trait with incomplete penetrance. The diagnosis is usually
	made before the age of 2 years when strabismus or a gray to yellow reflex from
	pupil ("cat eye") is investigated.,function:Key regulator of entry into cell division
	that acts as a tumor suppressor. Acts as a transcription repressor of E2F1 target
	genes. The underphosphorylated, active form of RB1 interacts
Outro allestar	Nucleus During konstingents differentiation and dation by KATOD/DOAE'
Subcellular	Nucleus . During keratinocyte differentiation, acetylation by KAT2B/PCAF is required for nuclear localization
Location :	
Expression	Expressed in the retina. Expressed in foreskin keratinocytes (at protein level)
Expression :	(PubMed:20940255).





Products Images

Immunohistochemical analysis of paraffin-embedded human Breast cancer. 1, Antibody was diluted at 1:200(4° overnight). 2, Tris-EDTA,pH9.0 was used for antigen retrieval. 3,Secondary antibody was diluted at 1:200(room temperature, 45min).

Western Blot analysis of 1 Jurkat treated with LPS, 2 Jurkat, using primary antibody at 1:1000 dilution. Secondary antibody(catalog#:RS23920) was diluted at 1:10000

