

POLR2I Ab

Affinity Biosciences website:www.affbiotech.com order:order@affbiotech.com

Images(2)

Concn.: ~1mg/ml	Mol.Wt.: 15kDa
Source: Rabbit	Clonality: Polyclonal
WB 1:500-1:2000, IHC 1:50-1:200 *The optimal dilutions should be determined by the end user	
Human,Mouse,Rat	
Rabbit IgG in phosphate buffered saline , pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Store at -20 °C. Stable for 12 months from date of receipt.	
The antiserum was purified by peptide affinity chromatography using SulfoLink TM Coupling Resin (Thermo Fisher Scientific).	
A synthesized peptide derived from human POLR2I, corresponding to a region within the internal amino acids.	
P36954	
This gene encodes a subunit of 1 responsible for synthesizing me combination with two other pol- domain of the polymerase, a gro transcribed into RNA. The prod with conserved cysteines and th [provided by RefSeq, Jul 2008]	RNA polymerase II, the polymerase ssenger RNA in eukaryotes. This subunit, in ymerase subunits, forms the DNA binding pove in which the DNA template is luct of this gene has two zinc finger motifs e subunit does possess zinc binding activity.
	Concn.: ~1mg/ml Source: Rabbit WB 1:500-1:2000, IHC 1:50-1 *The optimal dilutions should b Human,Mouse,Rat Rabbit IgG in phosphate bufferd sodium azide and 50% glycerol date of receipt. The antiserum was purified by J SulfoLink™ Coupling Resin (T A synthesized peptide derived f region within the internal amino P36954 This gene encodes a subunit of responsible for synthesizing me combination with two other pol domain of the polymerase, a gro transcribed into RNA. The prod with conserved cysteines and th [provided by RefSeq, Jul 2008]



Western blot analysis of extracts from 293t, using POLR2I Ab. The lane on the left was treated with blocking peptide.



DF6668 at 1/100 staining Rat kidney tissue by IHC-P. The sample was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The sample was then blocked and incubated with the Ab for 1.5 hours at 22°C. An HRP conjugated goat anti-rabbit Ab was used as the secondary Ab.



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IMPORTANT: For western blot, incubate membrane with diluted primary Ab in 5% w/v milk , 1X TBS, 0.1% Tween®20 at 4°C with gentle shaking, overnight.

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