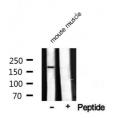


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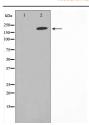
TOP2A Ab

Cat.#: AF0793 Size: 100ul,200ul	Concn.: 1mg/ml Source: Rabbit	Mol.Wt.: 174kDa Clonality: Polyclonal
Application:	WB 1:500-1:2000 IHC 1:50-1:200, IF/ICC 1:100-1:500	
Reactivity:	Human,Mouse	
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink™ Coupling Resin (Thermo Fisher Scientific).	
Specificity:	TOP2A Ab detects endogenous levels of TOP2A.	
Immunogen:	A synthesized peptide derived from human TOP2A.	
Uniprot:	P11388	
Description:	DNA topoisomerase 2-alpha; DNA topoisomerase II, 170 kD; DNA topoisomerase II, alpha isozyme; TOP2; TOP2A; topoisomerase (DNA) II alpha 170kDa; TP2A	
Subcellular Location:	Cytoplasm. Nucleus > nucleopla nucleoplasm.	asm. Generally located in the
Similarity:	The N-terminus has several stru domain (about residues 1-265), (about 266-428) and the toprim (PubMed:25202966). Comparing ATP hydrolysis induces domain are probably part of the mechai rejoining (PubMed:25202966).B topoisomerase family.	the transducer domain domain (455-572) g different structures shows shifts in the N-terminus that nism of DNA cleavage and
Storage Condition and Buffer:	Rabbit IgG in phosphate buffere NaCl, 0.02% sodium azide and s °C.Stable for 12 months from da	50% glycerol.Store at -20

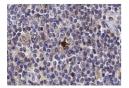


Western blot analysis on mouse muscle lysate using TOP2A Ab





Western blot analysis on Jurkat cell lysate using TOP2A Ab



AF0793 at 1/100 staining human Lymph node tissue sections by IHC-P. The tissue was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The tissue was then blocked and incubated with the Ab for 1.5 hours at 22°C. An HRP conjugated goat antirabbit Ab was used as the secondary.



AF0793 staining Hela by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25°C. The primary Ab was diluted at 1/200 and incubated with the sample for 1 hour at 37°C. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary Ab.

<code>IMPORTANT:</code> 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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