

Product datasheet for **TA500766M**

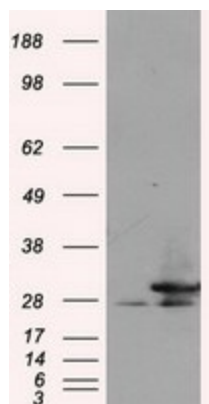
RPA34 (RPA2) Mouse Monoclonal Antibody [Clone ID: OTI4C3]

Product data:

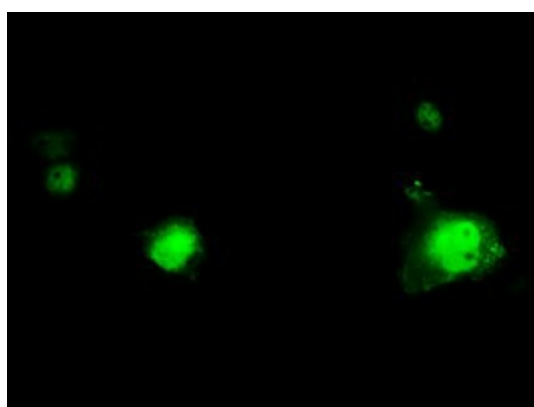
Product Type:	Primary Antibodies
Clone Name:	OTI4C3
Applications:	IF, IP, WB
Recommended Dilution:	WB 1:2000, IF 1:100, IP 2ug/500ul
Reactivity:	Human, Mouse, Rat
Host:	Mouse
Isotype:	IgG1
Clonality:	Monoclonal
Immunogen:	Full length human recombinant protein of human RPA2 (NP_002937) produced in HEK293T cell.
Formulation:	PBS (pH 7.3) containing 1% BSA, 50% glycerol and 0.02% sodium azide.
Concentration:	0.79 mg/ml
Purification:	Purified from mouse ascites fluids or tissue culture supernatant by affinity chromatography (protein A/G)
Conjugation:	Unconjugated
Storage:	Store at -20°C as received.
Stability:	Stable for 12 months from date of receipt.
Predicted Protein Size:	29.1 kDa
Gene Name:	replication protein A2
Database Link:	NP_002937 Entrez Gene 19891 Mouse Entrez Gene 59102 Rat Entrez Gene 6118 Human P15927
Synonyms:	REPA2; RP-A p32; RP-A p34; RPA32
Protein Families:	Druggable Genome, Stem cell - Pluripotency
Protein Pathways:	DNA replication, Homologous recombination, Mismatch repair, Nucleotide excision repair


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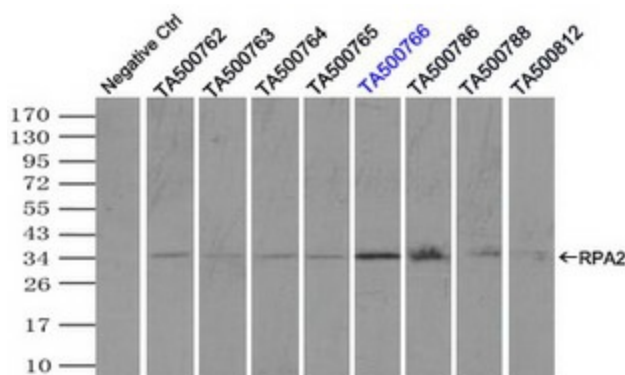
Product images:



HEK293T cells were transfected with the pCMV6-ENTRY control (Left lane) or pCMV6-ENTRY RPA2 ([RC205715], Right lane) cDNA for 48 hrs and lysed. Equivalent amounts of cell lysates (5 ug per lane) were separated by SDS-PAGE and immunoblotted with anti-RPA2. Positive lysates [LY401031] (100ug) and [LC401031] (20ug) can be purchased separately from OriGene.



Anti-RPA2 mouse monoclonal antibody ([TA500766]) immunofluorescent staining of COS7 cells transiently transfected by pCMV6-ENTRY RPA2 ([RC205715]).



Immunoprecipitation (IP) of RPA2 by using TrueMab monoclonal anti-RPA2 antibodies (Negative control: IP without adding anti-RPA2 antibody.). For each experiment, 500ul of DDK tagged RPA2 overexpression lysates (at 1:5 dilution with HEK293T lysate), 2ug of anti-RPA2 antibody and 20ul (0.1mg) of goat anti-mouse conjugated magnetic beads were mixed and incubated overnight. After extensive wash to remove any non-specific binding, the immunoprecipitated products were analyzed with rabbit anti-DDK polyclonal antibody.