## POLR2A Antibody

Cat.\#: MBS002301
Size: 100ul,200ul

Concn.: $1 \mathrm{mg} / \mathrm{ml}$
Source: Rabbit

Mol.Wt.: 270kDa
Clonality: Polyclonal

Application:

Reactivity:
Purification:

Specificity:

Immunogen:

Uniprot:
Storage Condition and Buffer:

WB 1:500-1:2000, IHC 1:50-1:200, IF/ICC 1:100-1:500, ELISA(peptide) 1:20000-1:40000

Human,Mouse,Rat
The antiserum was purified by peptide affinity chromatography using SulfoLink ${ }^{T M}$ Coupling Resin (Thermo Fisher Scientific).

POLR2A Antibody detects endogenous levels of total POLR2A.

A synthesized peptide derived from human POLR2A, corresponding to a region around the site of serine 1619.

P24928
Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM $\mathrm{NaCl}, 0.02 \%$ sodium azide and $50 \%$ glycerol.Store at $-20^{\circ}$ C.Stable for 12 months from date of receipt.

Western blot analysis of extracts from HeLa, using POLR2A antibody. The lane on the left was treated with the antigenspecific peptide.

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 antibody for 1.5 hours at $22^{\circ} \mathrm{C}$. An HRP conjugated goat anti-rabbit antibody was used as the secondary antibody.


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 $25^{\circ} \mathrm{C}$. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at $37^{\circ} \mathrm{C}$. An Alexa Fluor 594 conjugated goat anti-rabbit IgG $(H+L) A b$, diluted at $1 / 600$, was used as the secondary antibody.

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[^0]:    IMPORTANT: For western blot, incubate membrane with diluted primary Ab in $5 \% \mathrm{w} / \mathrm{v}$ milk , 1 X TBS, $0.1 \%$ Tween ${ }^{\circledR} 20$ at $4^{\circ} \mathrm{C}$ with gentle shaking, overnight.

