

**Fx1A Rabbit pAb**  
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**Catalog # AP93952****Specification**

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**Fx1A Rabbit pAb - Product Information**Application  
Host  
Clonality**IF, IHC-P**  
**Rabbit**  
**Polyclonal****Fx1A Rabbit pAb - Additional Information****Format**

0.01M TBS(pH7.4), 0.09% (W/V) sodium azide and 50% Glyce

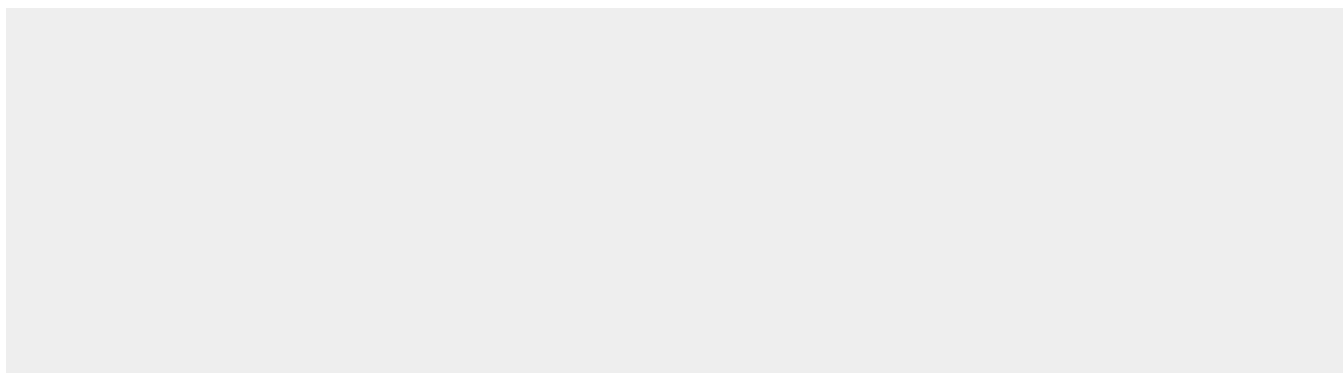
**Storage**

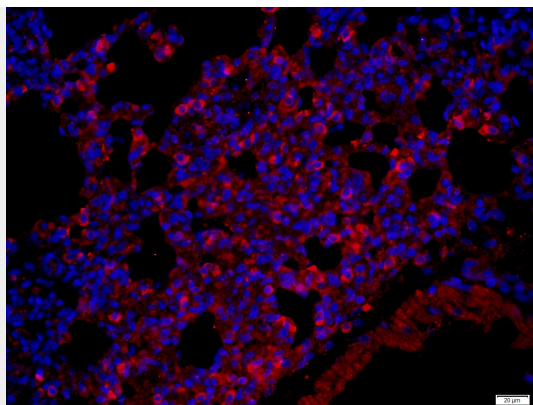
Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. When reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody is stable for at least two weeks at 2-4 °C.

**Fx1A Rabbit pAb - Protein Information****Fx1A Rabbit pAb - Protocols**

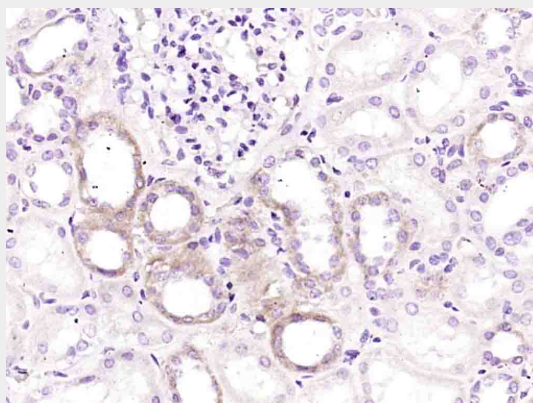
Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

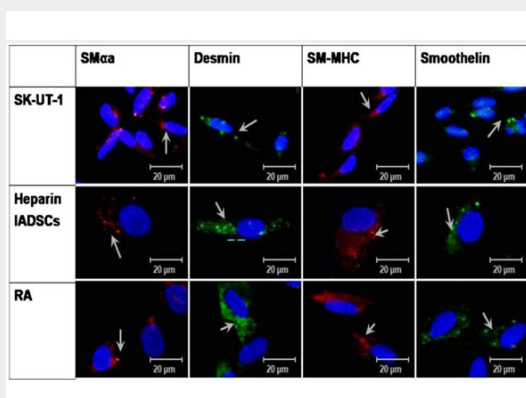
**Fx1A Rabbit pAb - Images**



Paraformaldehyde-fixed, paraffin embedded (rat lung); Antigen retrieval by boiling in sodium citrate buffer (pH6) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (Fx1A) Polyclonal Antibody, Unconjugated (AP93952) at 1:400 overnight at 4°C, followed by a conjugated secondary (Goat Anti-rabbit IgG/Bio) for 20 minutes at 37°C, followed by a conjugated streptavidin (bs-0437P-Cy3) at [1:500] for 40 minutes and DAPI staining of the nuclei.



Paraformaldehyde-fixed, paraffin embedded (Human kidney); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (Fx1A) Polyclonal Antibody, Unconjugated (AP93952) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Isolated hADSCs (IADSCs) were differentiated into SMCs using retinoic acid (RA), heparin was used as a positive control, while SK-UT-1 cells was used as a SMC control. Expression of SMC markers smooth muscle alpha actin (SM-αa, red, Texas Red Conjugated anti-Mouse IgG2, γ2a chain specific, p/n 610-4941), desmin (green, Fluorescein Conjugated anti-Mouse IgG1, γ1 chain

specific, p/n 610-4240), smooth muscle myosin heavy chain (SM-MHC, red, Texas Red Conjugated anti-Mouse  $\kappa$ , kappa chain specific, p/n 610-4910), and smoothelin (green, Fluorescein Conjugated anti-Mouse IgG1,  $\gamma$ 1 chain specific, p/n 610-4240) in differentiated SMCs was determined by indirect immunofluorescence. Nuclei were counter stained with DAPI (blue). Expression of all four markers can be seen in all the cells, particularly in RA differentiated SMCs. Fig. 5. PMID: 21373882.

**Fx1A Rabbit pAb - Background**

This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.