

## **Product datasheet for TA324879**

## **SAR1 (SAR1A) Rabbit Polyclonal Antibody**

**Product data:** 

**Product Type:** Primary Antibodies

**Applications:** FC, IHC, WB

**Recommended Dilution:** WB: 1:1000, IHC: 1:10~50, FC: 1:10~50 **Reactivity:** Human (Predicted: Mouse, Bovine, Pig)

Host: Rabbit Isotype: IgG

Clonality: Polyclonal

**Immunogen:** This SAR1A antibody is generated from rabbits immunized with a KLH conjugated synthetic

peptide between 122-149 amino acids from the Central region of human SAR1A.

**Formulation:** Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide.

**Concentration:** lot specific

**Purification:** This antibody is purified through a protein A column, followed by peptide affinity purification.

**Conjugation:** Unconjugated

**Storage:** Store at -20°C as received.

**Stability:** Stable for 12 months from date of receipt.

Predicted Protein Size: 22367 Da

**Gene Name:** secretion associated Ras related GTPase 1A

Database Link: NP 001136120

Entrez Gene 20224 MouseEntrez Gene 56681 Human

Q9NR31

**Synonyms:** masra2; SAR1; Sara; SARA1



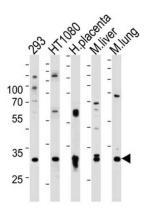
**OriGene Technologies, Inc.** 9620 Medical Center Drive, Ste 200

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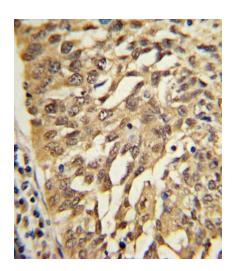
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## **Product images:**

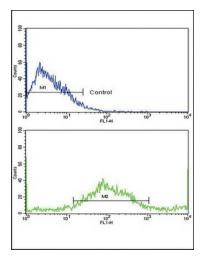


Western blot analysis of lysates from 293, HT1080 cell line, human placenta, mouse liver and lung tissue (from left to right), using SAR1A Antibody (Center) (Cat. #TA324879).TA324879 was diluted at 1:1000 at each lane. A goat anti-rabbit IgG H&L (HRP) at 1:5000 dilution was used as the secondary antibody.Lysates at 35ug per lane.



Formalin-fixed and paraffin-embedded human lung carcinoma with SAR1A Antibody (Center), which was peroxidase-conjugated to the secondary antibody, followed by DAB staining. This data demonstrates the use of this antibody for immunohistochemistry; clinical relevance has not been evaluated.





Flow cytometric analysis of NCI-H292 cells using SAR1A Antibody (Center) (bottom histogram) compared to a negative control cell (top histogram). FITC-conjugated goat-anti-rabbit secondary antibodies were used for the analysis.