## **STRA6 Antibody**



## PACO57456

Reactivity:

Human

## **Product Information**

Size: Protein Background:

50ug May act as a high-affinity cell-surface receptor for the complex retinol-retinol binding protein (RBP/RBP4). Acts by removing retinol from RBP/RBP4 and transports it across

the plasma membrane, where it can be metabolized. This mechanism does not depend on endocytosis. Binds to RBP/RBP4 with high affinity. Increases cellular retinol uptake

from the retinol-RBP complex.

Source: Gene ID:

Rabbit STRA6

Isotype: Uniprot

IgG Q9BX79

ELISA:1:2000-1:10000, IHC:1:200-1:500,

Applications: Synonyms:

ELISA, IHC, IF

Receptor for retinol uptake STRA6 (Retinol-binding protein receptor STRA6) (Stimulated

**Recommended dilutions:** by retinoic acid, gene 6 protein homolog), STRA6

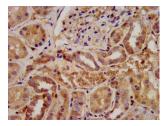
Immunogen:

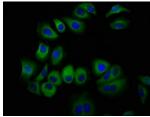
IF:1:50-1:200 Recombinant Human Receptor for retinol uptake STRA6 protein (530-667AA).

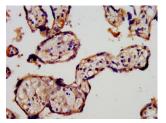
Storage:

Preservative: 0.03% Proclin 300. Constituents: 50% Glycerol, 0.01M PBS, pH 7.4

## **Product Images**







IHC image of PACO57456 diluted at 1:300 and staining in paraffinembedded human kidney tissue performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.

Immunofluorescence staining of A549 cells with PACO57456 at 1:100, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).

IHC image of PACO57456 diluted at 1:300 and staining in paraffinembedded human placenta tissue performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.