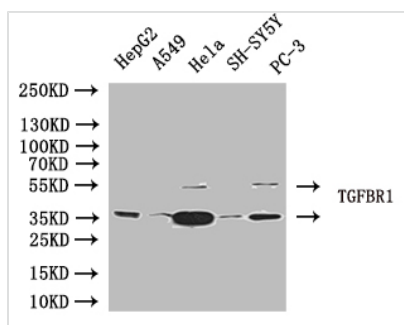




TGFBR1 Antibody

| | |
|----------------------------|---|
| Product Code | CSB-PA023451LA01HU |
| Storage | Upon receipt, store at -20°C or -80°C. Avoid repeated freeze. |
| Uniprot No. | P36897 |
| Immunogen | Synthetic peptide of human TGFBR1 |
| Raised In | Rabbit |
| Species Reactivity | Human |
| Tested Applications | ELISA, WB, IHC, IF; Recommended dilution: WB:1:500-1:2000, IHC:1:100-1:300, IF:1:10-1:100 |
| Form | Liquid |
| Conjugate | Non-conjugated |
| Storage Buffer | pH7.4 PBS, 0.05% NaN ₃ , 40% Glycerol |
| Purification Method | Antigen affinity purification |
| Isotype | IgG |
| Clonality | Polyclonal |
| Product Type | Polyclonal Antibody |
| Immunogen Species | Homo sapiens (Human) |
| Research Area | Cardiovascular |
| Target Names | TGFBR1 |

Image

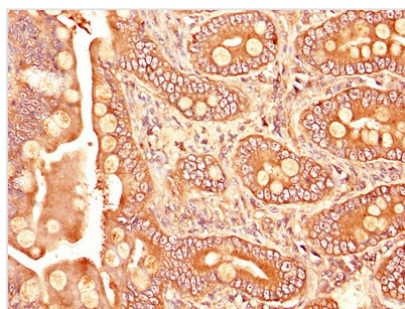


Western Blot

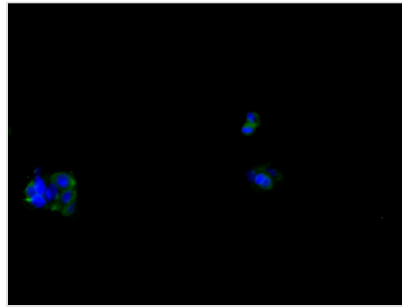
Positive WB detected in: HepG2 whole cell lysate, A549 whole cell lysate, HeLa whole cell lysate, SH-SY5Y, PC-3 whole cell lysate
All lanes: TGFBR1 antibody at 1:1000

Secondary

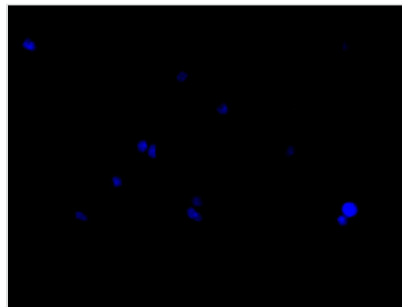
Goat polyclonal to rabbit IgG at 1/50000 dilution
Predicted band size: 56, 57, 48kDa
Observed band size: 58, 40kDa



IHC image of CSB-PA023451LA01HU diluted at 1:1200 and staining in paraffin-embedded human small intestine tissue performed on a Leica Bond™ 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 HepG2 cell with CSB-PA023451LA01HU at 1:40, counter-stained with DAPI. The cells were fixed in 4% formaldehyde and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4C. The secondary antibody was Alexa Fluor 488-conjugated AffiniPure Goat Anti-Rabbit IgG(H+L).



Immunofluorescence staining of HepG2 cell with 5% goat serum, counter-stained with DAPI. The cells were fixed in 4% formaldehyde and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4C. The secondary antibody was Alexa Fluor 488-conjugated AffiniPure Goat Anti-Rabbit IgG(H+L).

Usage

For Research Use Only. Not for use in diagnostic or therapeutic procedures.