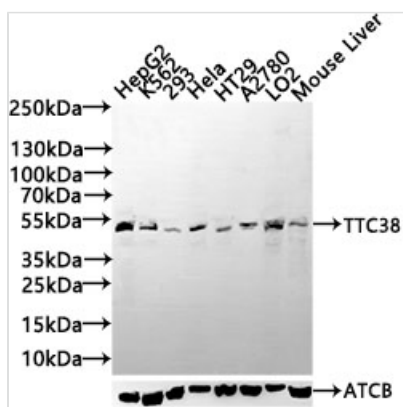




# TTC38 Antibody

<b>Product Code</b>	CSB-PA719094LA01HU
<b>Storage</b>	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
<b>Uniprot No.</b>	Q5R3I4
<b>Immunogen</b>	Fusion protein of human TTC38
<b>Raised In</b>	Rabbit
<b>Species Reactivity</b>	Human, Mouse
<b>Tested Applications</b>	ELISA, WB, IHC; Recommended dilution: WB:1:500-1:2000, IHC:1:20-1:200
<b>Form</b>	Liquid
<b>Storage Buffer</b>	pH7.4 PBS, 0.05% NaN <sub>3</sub> , 40% Glycerol
<b>Purification Method</b>	Antigen Affinity Purified
<b>Isotype</b>	IgG
<b>Clonality</b>	Polyclonal
<b>Product Type</b>	Polyclonal Antibody
<b>Immunogen Species</b>	Homo sapiens (Human)
<b>Research Area</b>	Cell Biology
<b>Target Names</b>	TTC38

## Image

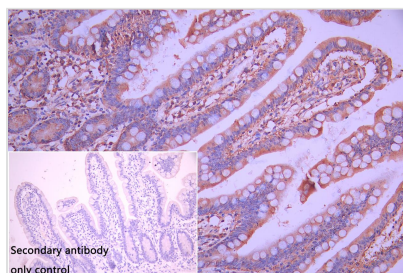


### Western Blot

Positive WB detected in: HepG2 whole cell lysate(30µg), K562 whole cell lysate(30µg), 293 whole cell lysate(30µg), HeLa whole cell lysate(30µg), HT29 whole cell lysate(30µg), A2780 whole cell lysate(30µg),LO2 whole cell lysate(30µg),Mouse Liver tissue lysate(30µg)  
All lanes: TTC38 antibody at 1:1000

### Secondary

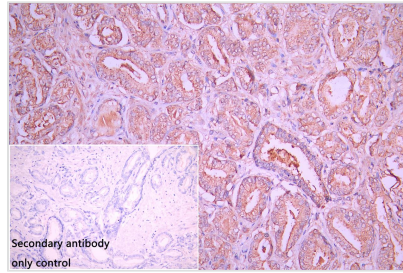
Goat polyclonal to rabbit IgG at 1/40000 dilution  
Predicted band size: 53 kDa  
Observed band size: 53 kDa  
Exposure time: 120s



IHC image of CSB-PA719094LA01HU diluted at 1:100 and staining in paraffin-embedded human small intestine tissue section 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 Goat anti-rabbit polymer IgG labeled by HRP and visualized using 0.05% DAB. Secondary antibody only control: uses 1% BSA instead of



primary antibody



IHC image of CSB-PA719094LA01HU diluted at 1:100 and staining in paraffin-embedded human prostate cancer 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 Goat anti-rabbit polymer IgG labeled by HRP and visualized using 0.05% DAB. Secondary antibody only control: uses 1% BSA instead of primary antibody

## Usage

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