

**bs-15428R****[ Primary Antibody ]****HCP1 Rabbit pAb****BioSS**  
**ANTIBODIES**

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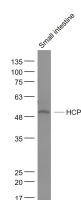
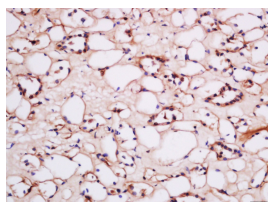
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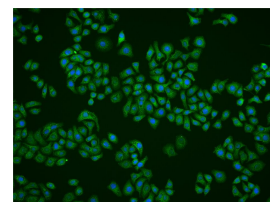
400-901-9800

**— DATASHEET —****Host:** Rabbit**Isotype:** IgG**Clonality:** Polyclonal**GeneID:** 113235**SWISS:** Q96NT5**Target:** HCP1**Immunogen:** KLH conjugated synthetic peptide derived from human HCP1: 341-459/459.**Purification:** affinity purified by Protein A**Concentration:** 1mg/ml**Storage:** 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.

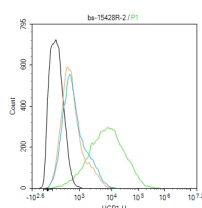
Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.

**Background:** This gene encodes a transmembrane proton-coupled folate transporter protein that facilitates the movement of folate and antifolate substrates across cell membranes, optimally in acidic pH environments. This protein is also expressed in the brain and choroid plexus where it transports folates into the central nervous system. This protein further functions as a heme transporter in duodenal enterocytes, and potentially in other tissues like liver and kidney. Its localization to the apical membrane or cytoplasm of intestinal cells is modulated by dietary iron levels. Mutations in this gene are associated with autosomal recessive hereditary folate malabsorption disease. Alternatively spliced transcript variants encoding different isoforms have been described for this gene. [provided by RefSeq, Aug 2013]**Applications:** WB (1:500-2000)**IHC-P** (1:100-500)**IHC-F** (1:100-500)**IF** (1:100-500)**Flow-Cyt** (2ug/Test)**ICC/IF** (1:100)**Reactivity:** Human, Mouse  
(predicted: Rat, Cow, Chicken)**Predicted MW.:** 50 kDa**Subcellular Location:** Cell membrane**— VALIDATION IMAGES —**Sample: Small intestine (Mouse) Lysate at 40 ug  
Primary: Anti- HCP1 (bs-15428R) at 1/1000  
dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 50 kD Observed band size: 50 kD

Tissue/cell: human kidney tissue; 4% Paraformaldehyde-fixed and paraffin-embedded; Antigen retrieval: citrate buffer (0.01M, pH 6.0), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum, C-0005) at 37°C for 20 min; Incubation: Anti-HCP1 Polyclonal Antibody, Unconjugated(bs-15428R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining



HepG2 cell; 4% Paraformaldehyde-fixed; Triton X-100 at room temperature for 20 min; Blocking buffer (normal goat serum, C-0005) at 37°C for 20 min; Antibody incubation with (HCP1) polyclonal Antibody, Unconjugated (bs-15428R) 1:100, 90 minutes at 37°C; followed by a conjugated Goat Anti-Rabbit IgG antibody at 37°C for 90 minutes, DAPI (blue, C02-04002) was used to stain the cell nuclei.



Blank control: Jurkat. Primary Antibody (green)

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

line): Rabbit Anti-HCP1 antibody (bs-15428R)  
Dilution: 2ug/Test; Secondary Antibody (white  
blue line) : Goat anti-rabbit IgG-FITC Dilution:  
0.5ug/Test. Isotype control (orange  
line) : Normal Rabbit IgG Protocol The cells  
were incubated in 5%BSA to block non-specific  
protein-protein interactions for 30 min at room  
temperature. Cells stained with Primary  
Antibody for 30 min at room temperature. The  
secondary antibody used for 40 min at room  
temperature. Acquisition of 20,000 events was  
performed.

## — SELECTED CITATIONS —

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- **[IF=17.5]** Jun-Jie Zou. et al. Efficient oral insulin delivery with sustained release by folate-conjugated metal-organic framework nanoparticles. MATTER-US. 2025 Mar;8: IF ;Human. 10.1016/j.matt.2024.101948