

**bs-4562R****[ Primary Antibody ]****CYP2E1 Rabbit pAb****BioSS**  
**ANTIBODIES**

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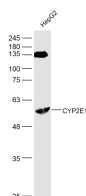
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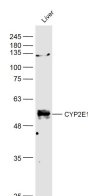
400-901-9800

**DATASHEET****Host:** Rabbit**Isotype:** IgG**Clonality:** Polyclonal**GeneID:** 1571**SWISS:** P05181**Target:** CYP2E1**Immunogen:** KLH conjugated synthetic peptide derived from human CYP2E1: 401-493/493.**Purification:** affinity purified by Protein A**Concentration:** 1mg/ml**Storage:** 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.

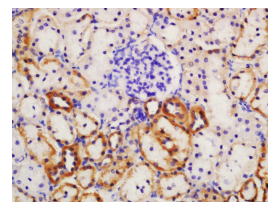
Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.

**Background:** The cytochrome P450 proteins are monooxygenases which catalyze many reactions involved in drug metabolism and biosynthesis. Cytochrome P450 2E1 is induced by ethanol, diabetes and starvation. The enzyme metabolizes both endogenous substrates, such as ethanol, acetone, and acetal, and exogenous substrates including benzene, carbon tetrachloride, ethylene glycol, and nitrosamines. Due to its many substrates, this enzyme may be involved in such varied processes as gluconeogenesis, hepatic cirrhosis, diabetes, and cancer.**Applications:** WB (1:500-2000)**IHC-P** (1:100-500)**IHC-F** (1:100-500)**IF** (1:100-500)**Flow-Cyt** (1:100-500)**Reactivity:** Human, Mouse, Rat  
(predicted: Rabbit, Pig, Sheep, Cow, Dog, Horse)**Predicted MW.:** 57 kDa**Subcellular Location:** Cell membrane ,Cytoplasm**VALIDATION IMAGES**

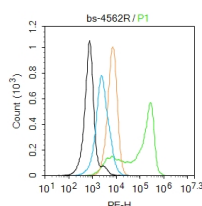
Sample: HepG2(Human) Cell Lysate at 30 ug  
 Primary: Anti-CYP2E1 (bs-4562R) at 1/300 dilution  
 Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution  
 Predicted band size: 57 kD Observed band size: 57 kD



Sample: Liver (Mouse) Lysate at 40 ug  
 Primary: Anti-CYP2E1 (bs-4562R) at 1/300 dilution  
 Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution  
 Predicted band size: 57 kD Observed band size: 57 kD



Tissue/cell: rat 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-CYP2E1 Polyclonal Antibody, Unconjugated(bs-4562R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining



Blank control:U937. Primary Antibody (green line): Rabbit Anti-CYP2E1 antibody (bs-4562R)  
 Dilution: 2µg /10<sup>6</sup> cells; Isotype Control  
 Antibody (orange line): Rabbit IgG. Secondary  
 Antibody : Goat anti-rabbit IgG-PE Dilution: 1µg

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

/test. Protocol The cells were fixed with 4% PFA (10min at room temperature) and then permeabilized with 0.1% PBST for 20 min at room temperature. The cells were then 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.

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## — SELECTED CITATIONS —

- **[IF=6.8]** Ye Jiazhou. et al. Single cell-spatial transcriptomics and bulk multi-omics analysis of heterogeneity and ecosystems in hepatocellular carcinoma. NPJ PRECIS ONCOL. 2024 Nov;8(1):1-18 IHC ;Human. 39548284
- **[IF=4.505]** Wang H et al. Regulatory T cells suppress excessive lipid accumulation in alcoholic liver disease. J Lipid Res. 2019 Feb 21. IHC ;Mouse. 30792182
- **[IF=4.784]** Wang C et al. A novel acidic polysaccharide from the residue of Panax notoginseng and its hepatoprotective effect on alcoholic liver damage in mice. Int J Biol Macromol. 2020 Feb 6;149:1084-1097. WB,IF ;Mouse. 32035151
- **[IF=5.4]** Shilin Sun. et al. The therapeutic effect of wine-processed Corni Fructus on chronic renal failure in rats through the interference with the LPS/IL-1-mediated inhibition of RXR function. J ETHNOPHARMACOL. 2024 Mar;321:117511 WB ;Rat. 38036016
- **[IF=4.2]** Qian Zhu. et al. Chronic alcohol intake disrupts cytochrome P450 enzyme activity in alcoholic fatty liver disease: insights into metabolic alterations and therapeutic targets. FRONT CHEM. 2025 May;13: WB ;Rat. 40433307