

bs-1292R**[Primary Antibody]**

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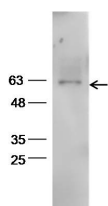
400-901-9800

Aromatase Rabbit pAb

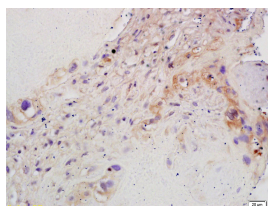
DATASHEET

Host: Rabbit	Isotype: IgG	Applications: WB (1:500-2000) IHC-P (1:100-500) IHC-F (1:100-500) IF (1:100-500) Flow-Cyt (1µg/Test)
Clonality: Polyclonal		
GeneID: 1588	SWISS: P11511	
Target: Aromatase		
Immunogen: KLH conjugated synthetic peptide derived from human Aromatase: 41-140/503.		Reactivity: Human, Rat (predicted: Mouse, Rabbit, Pig, Sheep, Cow, Dog, Horse)
Purification: affinity purified by Protein A		
Concentration: 1mg/ml		Predicted MW.: 58 kDa
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.		Subcellular Location: Cell membrane
Background: Aromatase is a key enzyme in steroidogenesis and plays an important role in sexual differentiation, oestrogen biosynthesis, fertility and carcinogenesis. It is highly conserved amongst mammals, and is highly expressed in placental tissue. Many environmental chemicals may influence aromatase activity and thereby disrupt endocrine function.		

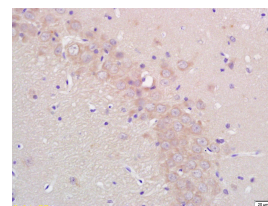
VALIDATION IMAGES



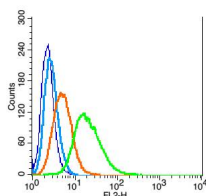
Sample: HepG2 Cell Lysate at 40 µg Primary: Anti-CYP19 (bs-1292R) at 1/300 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 58 kD Observed band size: 60 kD



Tissue/cell: human placenta 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-CYP19/CYP19A1 Polyclonal Antibody, Unconjugated (bs-1292R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody (SP-0023) and DAB (C-0010) staining



Tissue/cell: rat brain 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-CYP19/CYP19A1 Polyclonal Antibody, Unconjugated (bs-1292R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody (SP-0023) and DAB (C-0010) staining



Blank control: RSC96 (blue). Primary Antibody: Rabbit Anti-CYP19 antibody (bs-1292R), Dilution: 1µg in 100 µL 1X PBS containing 0.5% BSA; Isotype Control Antibody: Rabbit IgG (orange), used under the same conditions; Secondary Antibody: Goat anti-rabbit IgG-PE (white blue), Dilution: 1:200 in 1X PBS containing 0.5% BSA. Protocol The cells

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were fixed with 2% paraformaldehyde (10 min). Primary antibody (bs-1292R, 1 μ g /1x10⁶ cells) were incubated for 30 min on the ice, followed by 1 X PBS containing 0.5% BSA + 1 0% goat serum (15 min) to block non-specific protein-protein interactions. Then the Goat Anti-rabbit IgG/PE antibody was added into the blocking buffer mentioned above to react with the primary antibody at 1/200 dilution for 30 min on ice. Acquisition of 20,000 events was performed.

— SELECTED CITATIONS —

- **[IF=8.1]** Chungmo Yang. et al. Injectable Biomimetic Hydrogel Constructs for Cell-Based Menopausal Hormone Therapy with Reduced Breast Cancer Potential. *Biomaterials Research*. 2024 Aug;28 IF ;Rat. 39135549
- **[IF=4.8]** Lizheng Wu. et al. Luoshi Neiyi Prescription inhibits estradiol synthesis and inflammation in endometriosis through the HIF1A/EZH2/SF-1 pathway. *J ETHNOPHARMACOL*. 2024 Dec;335:118659 WB ;Human,Rat. 39098622
- **[IF=5.201]** Zhong Yuyi. et al. MIR143 Inhibits Steroidogenesis and Induces Apoptosis Repressed by H3K27me3 in Granulosa Cells. *Front Cell Dev Biol*. 2020 Oct;8:1159 WB ;Porcine. 33195195
- **[IF=3.688]** Yan Deng. et al. Molecular characterization, expression profile and transcriptional regulation of the CYP19 gene in goose ovarian follicles. *Gene*. 2022 Jan;806:145928 IHC ;Goose. 34455027
- **[IF=3.3]** Arabacı Tamer Sevil. et al. Neuropeptide W protects against ovarian oxidative injury and reinforces ovarian steroidogenic activity via the upregulation of ER α expression. *J PHARM PHARMACOL*. 2024 Mar;: IHC ;Rat. 38457354