## bs-1509R

## [ Primary Antibody ]

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# BMPR1A Rabbit pAb

DATASHEET -

Host: Rabbit Isotype: IgG

Clonality: Polyclonal

GeneID: 657 **SWISS:** P36894

Target: BMPR1A

**Immunogen:** KLH conjugated synthetic peptide derived from human BMPR-1A:

101-200/532.

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 bone morphogenetic protein (BMP) receptors are a family of transmembrane serine/threonine kinases that include the type I receptors BMPR1A and BMPR1B and the type II receptor BMPR2. These receptors are also closely related to the activin receptors, ACVR1 and ACVR2. The ligands of these receptors are members of the TGF-beta superfamily. TGF-betas and activins transduce their signals through the formation of heteromeric complexes with 2 different types of serine (threonine) kinase receptors: type I receptors of about 50-55 kD and type II receptors of about 70-80 kD. Type II receptors bind ligands in the absence of type I receptors, but they require their respective type I receptors for signaling, whereas type I receptors require their respective type II receptors for ligand binding. [provided by RefSeq].

Applications: WB (1:500-2000)

**IHC-P** (1:100-500) IHC-F (1:100-500) **IF** (1:100-500) Flow-Cyt (1ug/Test)

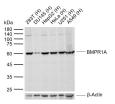
Reactivity: Human, Rat

(predicted: Mouse)

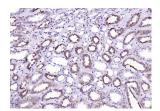
**Predicted** 60 kDa MW.:

Subcellular Cell membrane

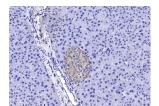
### VALIDATION IMAGES



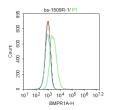
Sample: Lane 1: Human 293T cell lysates Lane 2: Human DU145 cell lysates Lane 3: Human HepG2 cell lysates Lane 4: Human HeLa cell lysates Lane 5: Human U251 cell lysates Lane 6: Human A549 cell lysates Primary: Anti-BMPR1A (bs-1509R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 60 kDa Observed band size: 60 kDa



Paraformaldehyde-fixed, paraffin embedded (Human kidney); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (BMPR1A) Polyclonal Antibody, Unconjugated (bs-1509R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Paraformaldehyde-fixed, paraffin embedded (rat pancreas); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (BMPR1A) Polyclonal Antibody, Unconjugated (bs-1509R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Blank control: A549. Primary Antibody (green

line): Rabbit Anti-BMPR1A antibody (bs-1509R)
Dilution: 1ug/Test; Secondary Antibody: Goat
anti-rabbit IgG-FITC Dilution: 0.5ug/Test.
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 -

- [IF=2.4] Tao Liu. et al. Dapagliflozin attenuates cardiac remodeling and dysfunction in rats with β-adrenergic receptor overactivation through restoring calcium handling and suppressing cardiomyocyte apoptosis. DIABETES VASC DIS RE.;(): WB;Rat. 37589258
- [IF=2] Yuki Koizumi. et al. Possible roles of bone morphogenetic protein 4 in regulating endometrial function in cows. ANIM SCI J. 2023 Aug;94(1):e13866 IHC; Bovine. 37632404