

bs-13175R**[Primary Antibody]****FKBP10 Rabbit pAb****Bioss**
ANTIBODIES

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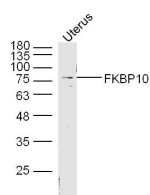
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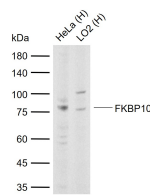
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— DATASHEET —

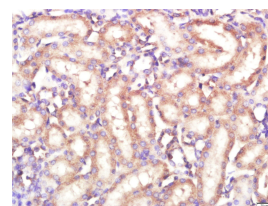
<p>Host: Rabbit</p> <p>Clonality: Polyclonal</p> <p>GeneID: 60681</p> <p>Target: FKBP10</p> <p>Immunogen: KLH conjugated synthetic peptide derived from human FKBP10: 31-130/582.</p> <p>Purification: affinity purified by Protein A</p> <p>Concentration: 1mg/ml</p> <p>Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol. Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.</p> <p>Background: The immunophilins are a highly conserved family of cis-trans peptidyl-prolyl isomerases that bind to and mediate the effects of immunosuppressive drugs, such as cyclosporin, FK506 and rapamycin. Immunophilins have also been implicated in protein folding and trafficking within the endoplasmic reticulum (ER). FKBP10 (FK506-binding protein 10), also known as peptidyl-prolyl cis-trans isomerase, PPIase, Rotamase, 65kDa FK506-binding protein or FKBP65, is a 582 amino acid immunophilin localized to the ER lumen and found in many tissues including heart, spleen, brain, testis and lung. FKBP10 contains two EF-hand calcium-binding domains and four PPIase FKBP-type domains, suggesting an enzymatic role in protein folding by catalyzing the cis-trans isomerization of proline imidic peptide bonds in oligopeptides. FKBP10 also acts as a receptor for the immunosuppressants FK506 and rapamycin, which inhibit FKBP10 activity. FKBP10 is thought to interact with the Raf-1/HSP 90 heterocomplex during signal transduction processes, and may associate with elastin during elastin protein folding and transport. With a valine-24 addition to human FKBP10, human and mouse FKBP10 are almost identical.</p>	<p>Applications: WB (1:500-2000) IHC-P (1:100-500) IHC-F (1:100-500) IF (1:100-500)</p> <p>Reactivity: Human, Mouse, Rat (predicted: Pig, Sheep, Cow, Dog, Horse)</p> <p>Predicted MW.: 61 kDa</p> <p>Subcellular Location: Cytoplasm</p>
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— VALIDATION IMAGES —

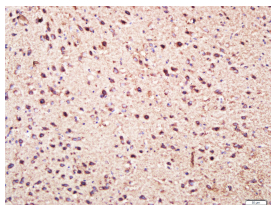
Sample: Uterus (Mouse) Lysate at 40 ug Primary: Anti-FKBP10(bs-13175R) at 1/300 dilution
Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 61 kD
Observed band size: 76 kD



Sample: Lane 1: Human HeLa cell lysates Lane 2: Human LO2 cell lysates Primary: Anti-FKBP10 (bs-13175R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 61 kDa Observed band size: 77 kDa



Paraformaldehyde-fixed, paraffin embedded (rat 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 (FKBP10) Polyclonal Antibody, Unconjugated (bs-13175R) at 1:400 overnight at 4°C, followed by a conjugated secondary (sp-0023) for 20 minutes and DAB staining.



Paraformaldehyde-fixed, paraffin embedded (rat brain); 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 (FKBP10) Polyclonal Antibody, Unconjugated (bs-13175R) at 1:400 overnight at 4°C, followed by a conjugated secondary (sp-0023) for 20 minutes and DAB staining.

— SELECTED CITATIONS —

- **[IF=5.06]** Sun, Zhen, et al. "Identification of Chemoresistance-Related Cell Surface Glycoproteins in Leukemia Cells and Functional Validation of Candidate Glycoproteins." *Journal of Proteome Research* (2014). WB ;="Human". 24467213
- **[IF=4.7]** Haixia Xu. et al. FKBP10 Silencing Alleviates Gluteal Muscle Contracture by Inhibiting Fibrosis and Restoring Autophagy via HSP47/SMAD3 Pathway Inactivation. *AM J PATHOL.* 2025 Apr;; IF ;Rat. 40316212
- **[IF=3.041]** Liang L et al. Comprehensive evaluation of FKBP10 expression and its prognostic potential in gastric cancer. *Oncol Rep.* 2019 Aug;42(2):615-628. IHC ;Human. 31233188
- **[IF=2.6]** Chang Jiale. et al. Screening and expression validation of key proteins for secondary hair follicle growth in cashmere goats based on iTRAQ quantitative proteomics technology. *FRONT VET SCI.* 2024 Oct;11: WB,IHC ;Goat. 39474271