

bs-0943R**[Primary Antibody]****BioSS**
ANTIBODIES

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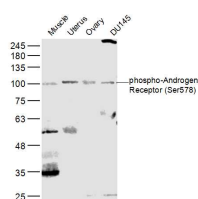
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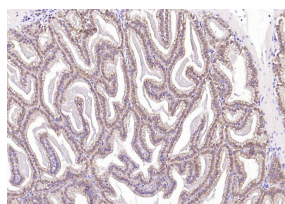
400-901-9800

phospho-AR/Androgen receptor (Ser578) Rabbit pAb**— DATASHEET —**

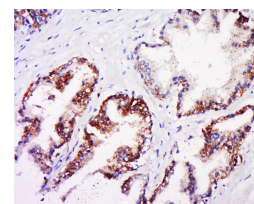
Host: Rabbit	Isotype: IgG	Applications: WB (1:500-2000) IHC-P (1:100-500) IHC-F (1:100-500) IF (1:100-500) Flow-Cyt (3ug/Test)
Clonality: Polyclonal		
GeneID: 367	SWISS: P10275	
Target: AR/Androgen receptor (Ser578)		
Immunogen: KLH conjugated Synthesised phosphopeptide derived from human Androgen Receptor around the phosphorylation site of Ser578: CG(p-S)C.		
Purification: affinity purified by Protein A		Reactivity: Human, Mouse, Rat (predicted: Pig, Cow, Chicken, Dog, Horse)
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 androgen receptor gene is more than 90 kb long and codes for a protein that has 3 major functional domains: the N-terminal domain, DNA-binding domain, and androgen-binding domain. The protein functions as a steroid-hormone activated transcription factor. Upon binding the hormone ligand, the receptor dissociates from accessory proteins, translocates into the nucleus, dimerizes, and then stimulates transcription of androgen responsive genes. This gene contains 2 polymorphic trinucleotide repeat segments that encode polyglutamine and polyglycine tracts in the N-terminal transactivation domain of its protein. Expansion of the polyglutamine tract causes spinal bulbar muscular atrophy (Kennedy disease). Mutations in this gene are also associated with complete androgen insensitivity (CAIS). Two alternatively spliced variants encoding distinct isoforms have been described. [provided by RefSeq, Jul 2008]		Predicted MW.: 43/101 kDa Subcellular Location: Cytoplasm ,Nucleus

— VALIDATION IMAGES —

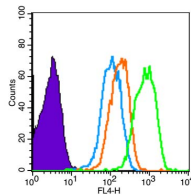
Sample: Muscle (Mouse) Lysate at 40 ug Uterus (Mouse) Lysate at 40 ug Ovary (Rat) Lysate at 40 ug DU145 (Human) Cell Lysate at 40 ug Primary: Anti-phospho-Androgen Receptor (Ser578) (bs-0943R) at 1/300 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 101 kD Observed band size: 101 kD



Paraformaldehyde-fixed, paraffin embedded (mouse prostate); 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; Incubation with (phospho-AR/Androgen receptor (Ser578)) Polyclonal Antibody, Unconjugated (bs-0943R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Tissue/cell: human prostate 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-AR Polyclonal Antibody, Unconjugated (bs-0943R) 1:500, overnight at 4°C, followed by conjugation to the secondary antibody (SP-0023) and DAB (C-0010) staining



Blank control (Black line): U87MG (Black).
 Primary Antibody (green line): Rabbit Anti-phospho-Androgen Receptor (Ser578) antibody (bs-0943R) Dilution: 1 μ g / 10⁶ cells; Isotype Control Antibody (orange line): Rabbit IgG .
 Secondary Antibody (white blue line): Goat anti-rabbit IgG-AF647 Dilution: 1 μ g /test. Protocol
 The cells were fixed with 4% PFA (10min at room temperature) and then permeabilized with 90% ice-cold methanol 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 10,000 events was performed.