

AR (Acetyl Lys633) rabbit pAb

Cat No.:ES8620

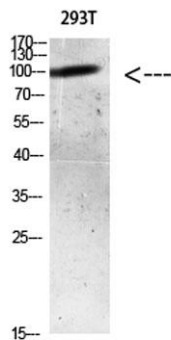
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Overview

| | |
|--------------------------|---|
| Product Name | AR (Acetyl Lys633) rabbit pAb |
| Host species | Rabbit |
| Applications | WB;ELISA |
| Species Cross-Reactivity | Human:K633;Mouse:K613;Rat:K616 |
| Recommended dilutions | WB 1:500-2000, ELISA 1:10000-20000 |
| Immunogen | Synthetic Acetyl peptide from human protein at AA range: 633 |
| Specificity | This antibody detects endogenous levels of AR at Human:K633;Mouse:K613;Rat:K616, It doesn't react with total protein. |
| Formulation | Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide. |
| Storage | Store at -20°C. Avoid repeated freeze-thaw cycles. |
| Protein Name | Androgen receptor (Dihydrotestosterone receptor) (Nuclear receptor subfamily 3 group C member 4) |
| Gene Name | AR DHTR NR3C4 |
| Cellular localization | Nucleus . Cytoplasm . Detected at the promoter of target genes (PubMed:25091737). Predominantly cytoplasmic in unligated form but translocates to the nucleus upon ligand-binding. Can also translocate to the nucleus in unligated form in the presence of RAC |
| Purification | The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen. |
| Clonality | Polyclonal |
| Concentration | 1 mg/ml |
| Observed band | 100kD |
| Human Gene ID | 367 |
| Human Swiss-Prot Number | P10275 |
| Alternative Names | Androgen receptor (Dihydrotestosterone receptor) (Nuclear receptor subfamily 3 group C member 4) |
| 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 from the normal 9-34 repeats to the pathogenic 38-62 repeats 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 isoform



Western blot analysis of mouse-lung lysate, antibody was diluted at 500. Secondary antibody(catalog#:RS0002) was diluted at 1:20000

