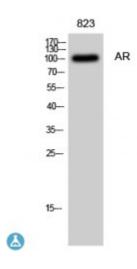


Anti-AR antibody



Description Rabbit polyclonal to AR.

Model STJ91659

Host Rabbit

Reactivity Human

Applications ELISA, IHC, WB

Immunogen Synthesized peptide derived from human AR around the non-phosphorylation

site of S650.

Immunogen Region 590-670 aa

Gene ID <u>367</u>

Gene Symbol AR

Dilution range WB 1:500-1:2000IHC 1:100-1:300ELISA 1:40000

Specificity AR Polyclonal Antibody detects endogenous levels of AR protein.

Tissue Specificity Isoform 2 is mainly expressed in heart and skeletal muscle. Isoform 3 is

expressed by basal and stromal cells of prostate (at protein level).

Purification The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

Note For Research Use Only (RUO).

Protein Name Androgen receptor Dihydrotestosterone receptor Nuclear receptor subfamily 3

group C member 4

Molecular Weight 110/99 kDa

Clonality Polyclonal

Conjugation Unconjugated

Isotype IgG

Modifications

Formulation Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Concentration 1 mg/ml

Storage Instruction Store at -20°C, and avoid repeat freeze-thaw cycles.

Database Links HGNC:644OMIM:300068

Alternative Names Androgen receptor Dihydrotestosterone receptor Nuclear receptor subfamily 3

group C member 4

Function Steroid hormone receptors are ligand-activated transcription factors that

regulate eukaryotic gene expression and affect cellular proliferation and differentiation in target tissues. Transcription factor activity is modulated by bound coactivator and corepressor proteins. Transcription activation is down-regulated by NR0B2. Activated, but not phosphorylated, by HIPK3 and ZIPK/DAPK3. Isoform 3 and isoform 4 lack the C-terminal ligand-binding domain and may therefore constitutively activate the transcription of a specific

set of genes independently of steroid hormones.

Sequence and Domain Family Composed of three domains: a modulating N-terminal domain, a DNA-

binding domain and a C-terminal ligand-binding domain. In the presence of bound steroid the ligand-binding domain interacts with the N-terminal modulating domain, and thereby activates AR transcription factor activity. Agonist binding is required for dimerization and binding to target DNA. The transcription factor activity of the complex formed by ligand-activated AR and DNA is modulated by interactions with coactivator and corepressor proteins. Interaction with RANBP9 is mediated by both the N-terminal domain and the DNA-binding domain. Interaction with EFCAB6/DJBP is

mediated by the DNA-binding domain.

Cellular Localization Nucleus Cytoplasm. 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 RACK1.

Post-translational Sumoylated on Lys-388 (major) and Lys-521. Ubiquitinated. Deubiquitinated

AR transcriptional activity and specificity. Phosphorylated in prostate cancer cells in response to several growth factors including EGF. Phosphorylation is induced by c-Src kinase (CSK). Tyr-535 is one of the major phosphorylation sites and an increase in phosphorylation and Src kinase activity is associated with prostate cancer progression. Phosphorylation by TNK2 enhances the DNA-binding and transcriptional activity and may be responsible for androgen-independent progression of prostate cancer. Phosphorylation at

by USP26. 'Lys-6' and 'Lys-27'-linked polyubiquitination by RNF6 modulates

Phosphorylation by PAK6 leads to AR-mediated transcription inhibition. Palmitoylated by ZDHHC7 and ZDHHC21. Palmitoylation is required for plasma membrane targeting and for rapid intracellular signaling via ERK and

Ser-83 by CDK9 regulates AR promoter selectivity and cell growth.

AKT kinases and cAMP generation.

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