

androgen receptor Monoclonal antibody

 Catalog Number: 66747-1-Ig 4 Publications

Basic Information

| | | |
|---|---|--|
| Catalog Number: 66747-1-Ig | GenBank Accession Number: BC132975 | Purification Method: Protein A purification |
| Size: 2300 µg/ml | GeneID (NCBI): 367 | CloneNo.: 1F7C12 |
| Source: Mouse | UNIPROT ID: P10275 | Recommended Dilutions: WB 1:600-1:3000 IHC 1:5000-1:20000 IF 1:200-1:800 |
| Isotype: IgG2a | Full Name: androgen receptor | |
| Immunogen Catalog Number: AG17291 | Calculated MW: 914 aa, 99 kDa Observed MW: 110-120 kDa | |

Applications

| | |
|--|---|
| Tested Applications: IF/ICC, IF-P, IHC, WB, ELISA | Positive Controls: WB : LNCaP cells, human testis tissue, NCCIT cells IHC : human prostate cancer tissue, mouse testis tissue, rat testis tissue IF : human prostate cancer tissue, LNCaP cells |
| Cited Applications: IF, IHC, IP, WB | |
| Species Specificity: Human, Mouse, Rat | |
| Cited Species: human, rat, mouse | |
| Note-IHC: suggested antigen retrieval with TE buffer pH 9.0; (*) Alternatively, antigen retrieval may be performed with citrate buffer pH 6.0 | |

Background Information

AR, also named as DHTR and NR3C4, belongs to the nuclear hormone receptor family and NR3 subfamily. AR is a 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. AR is activated, but not phosphorylated, by HIPK3. Defects in AR are the cause of androgen insensitivity syndrome (AIS), previously known as testicular feminization syndrome (TFM), which is an X-linked recessive form of pseudohermaphroditism due end-organ resistance to androgen. Defects in AR are the cause of spinal and bulbar muscular atrophy X-linked type 1 (SMA X1) which also known as Kennedy disease. Defects in AR may play a role in metastatic prostate cancer. Defects in AR are the cause of androgen insensitivity syndrome partial (PAIS) which also known as Reifenstein syndrome. AR exists various isoforms with MW 110-120 kDa and 75-80 kDa. (PMID: 19244107)

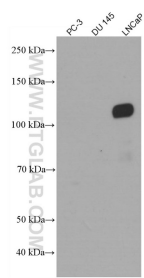
Notable Publications

| Author | Pubmed ID | Journal | Application |
|----------------|-----------|--------------------|-------------|
| Yuan-Xue Jing | 37931646 | Gynecol Endocrinol | WB, IF |
| Kai Song | 37810250 | iScience | WB, IF, IP |
| Parmveer Singh | 37376888 | Development | IHC |

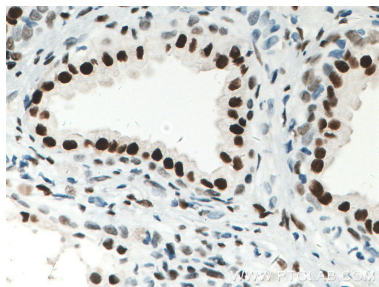
Storage

Storage:
Store at -20°C. Stable for one year after shipment.
Storage Buffer:
PBS with 0.02% sodium azide and 50% glycerol pH 7.3.
Aliquoting is unnecessary for -20°C storage

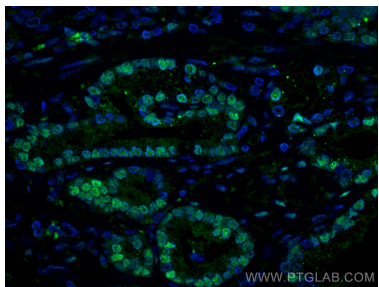
Selected Validation Data



PC-3(AR-), DU 145(AR-) and LNCaP (AR+) cell lysates were subjected to SDS PAGE followed by western blot with 66747-1-Ig (AR antibody) at dilution of 1:3000 incubated at room temperature for 1.5 hours.



Immunohistochemical analysis of paraffin-embedded human prostate cancer tissue slide using 66747-1-Ig (AR antibody) at dilution of 1:20000 (under 40x lens. Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).



Immunofluorescent analysis of (4% PFA) fixed human prostate cancer tissue using AR antibody (66747-1-Ig, Clone: 1F7C12) at dilution of 1:400 and CoraLite®488-Conjugated AffiniPure Goat Anti-Mouse IgG(H+L).