

Phospho-Androgen Receptor-S650 Rabbit pAb

货号: B15101

产品信息

| 反应 | Human |
|-------|-----------------------------------------------------------------------------------------------------|
| 宿主 | Rabbit |
| 克隆性 | Polyclonal |
| 预测反应 | |
| 应用 | IHC |
| 推荐浓度 | IHC: 1:50 - 1:100 |
| 理论分子量 | 44kDa/67kDa/99kDa |
| 实测分子量 | 44kDa/67kDa/99kDa |
| 形式 | Liquid |
| 保存条件 | Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.02% sodium azide,50% glycerol,pH7.3. |
| 偶联物 | Unconjugated |
| 阳性对照 | LNCaP cells, MCF-7 cells, K-562 cells |
| 细胞定位 | Cytoplasm,Nucleus |
| 纯化 | Affinity purification |
| | |

抗原信息

| 抗原信息 | A phospho specific peptide corresponding to residues surrounding S650 of human AR |
|------|-----------------------------------------------------------------------------------|
| 序列 | |

靶点信息

| 研究背景 | 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 isoforms have been described. |
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| 基因ID | 367 |
| 基因名 | AR |
| Swiss | P10275 |
| 别名 | AIS;AR8;DHTR;HUMARA;HYSP1;KD;NR3C4;SBMA;SMAX1;TFM;AR |
| | |

产品验证

实验步骤

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