



LHR Polyclonal Antibody

| Catalog No | BYab-13394 |
|--------------------|--|
| Isotype | lgG |
| Reactivity | Human;Mouse;Rat |
| Applications | WB;ELISA;IHC |
| Gene Name | LHCGR |
| Protein Name | Lutropin-choriogonadotropic hormone receptor |
| Immunogen | The antiserum was produced against synthesized peptide derived from human LSHR. AA range:621-670 |
| Specificity | LHR Polyclonal Antibody detects endogenous levels of LHR protein. |
| Formulation | Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide. |
| Source | Polyclonal, Rabbit,IgG |
| Purification | The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen. |
| Dilution | WB 1:500-2000;IHC-p 1:50-300; ELISA 2000-20000 |
| Concentration | 1 mg/ml |
| Purity | ≥90% |
| Storage Stability | -20°C/1 year |
| Synonyms | LHCGR; LCGR; LGR2; LHRHR; Lutropin-choriogonadotropic hormone receptor; LH/CG-R; Luteinizing hormone receptor; LHR; LSH-R |
| Observed Band | 80kD |
| Cell Pathway | Cell membrane ; Multi-pass membrane protein . |
| Tissue Specificity | Gonadal and thyroid cells. |
| Function | alternative products:Additional isoforms seem to exist,disease:Defects in LHCGR are a cause of familial male precocious puberty (FMPP) [MIM:176410]; also known as testotoxicosis. In FMPP the receptor is constitutively activated.,disease:Defects in LHCGR are a cause of Leydig cell hypoplasia (LCH) [MIM:152790]. LCH is an autosomal recessive disease characterized by male pseudohermaphroditism. In LCH the testes are small with marked immaturity of the Leydig cells which correlates with undetectable plasma testosterone levels and elevated gonadotropins.,function:Receptor for lutropin-choriogonadotropic hormone. The activity of this receptor is mediated by G proteins which activate adenylate cyclase.,online information:Glycoprotein-hormone Receptors |

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| | Information System, similarity:Belongs to the G-protein coupled receptor 1 family., similarity:Belongs to the G-protein coupled receptor 1 family. | |
| Background | This gene encodes the receptor for both luteinizing hormone and choriogonadotropin. This receptor belongs to the G-protein coupled recept family, and its activity is mediated by G proteins which activate adenylate of Mutations in this gene result in disorders of male secondary sexual charact development, including familial male precocious puberty, also known as testotoxicosis, hypogonadotropic hypogonadism, Leydig cell adenoma with precocious puberty, and male pseudohermaphtoditism with Leydig cell hypoplasia. [provided by RefSeq, Jul 2008], | cyclase. cter |
| matters needing attention | Avoid repeated freezing and thawing! | |
| Usage suggestions | This product can be used in immunological reaction related experiments. I more information, please consult technical personnel. | For |
| | | |

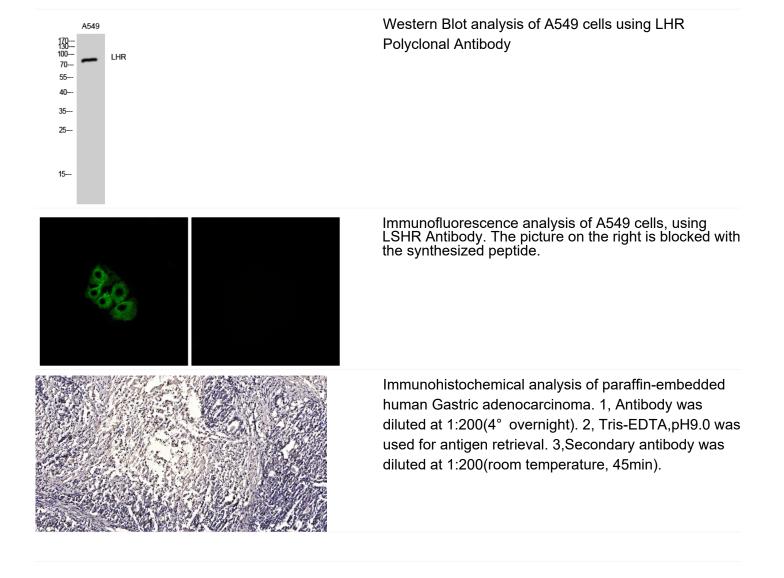
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