abcam

Product datasheet

Anti-FSH-R antibody ab75200

12 References 1 图像

概述

产品名称 Anti-FSH-R抗体

描述 兔多克隆抗体to FSH-R

宿主 Rabbit

经测试应用 适用于: WB

种属反应性 与反应: Human

预测可用于: Mouse, Rat 🔷

免疫原 Synthetic peptide corresponding to Human FSH-R (internal sequence).

阳性对照 Transfected Jurkat and HUVEC cell extracts

常规说明

The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

性能

形式 Liquid

存放说明 Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

存储溶液 pH: 7.40

Preservative: 0.02% Sodium azide

Constituents: 50% Glycerol (glycerin, glycerine), 0.87% Sodium chloride, PBS

Without Mg2+ and Ca2+

纯**度** Immunogen affinity purified

纯**化**说明 Affinity purified from rabbit antiserum by affinity chromatography using epitope specific

immunogen.

克隆 多克隆

同种型 IgG

1

The Abpromise guarantee

Abpromise™承诺保证使用ab75200于以下的经测试应用

"应用说明"部分下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
WB		1/500 - 1/1000. Predicted molecular weight: 78 kDa.

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功能

 $\label{lem:continuity} Receptor for follicle-stimulating hormone. The activity of this receptor is mediated by G proteins$

which activate adenylate cyclase.

组织特异性

Sertoli cells and ovarian granulosa cells.

疾病相关

Defects in FSHR are a cause of ovarian dysgenesis type 1 (ODG1) [MIM:233300]; also known as premature ovarian failure or gonadal dysgenesis XX type or XX gonadal dysgenesis (XXGD) or hereditary hypergonadotropic ovarian failure or hypergonadotropic ovarian dysgenesis with normal karyotype. ODG1 is an autosomal recessive disease characterized by primary amenorrhea, variable development of secondary sex characteristics, and high serum levels of

follicle-stimulating hormone (FSH) and luteinizing hormone (LH).

Defects in FSHR are a cause of ovarian hyperstimulation syndrome (OHSS) [MIM:608115]. OHSS is a disorder which occurs either spontaneously or most often as an iatrogenic complication of ovarian stimulation treatments for in vitro fertilization. The clinical manifestations vary from abdominal distention and discomfort to potentially life-threatening, massive ovarian enlargement and capillary leak with fluid sequestration. Pathologic features of this syndrome include the presence of multiple serous and hemorrhagic follicular cysts lined by luteinized cells, a

condition called hyperreactio luteinalis.

序列相似性 Belongs to the G-protein coupled receptor 1 family. FSH/LSH/TSH subfamily.

Contains 9 LRR (leucine-rich) repeats.

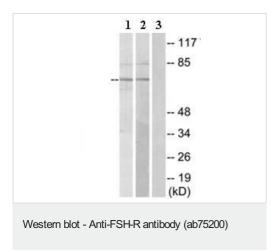
Contains 1 LRRNT domain.

翻译后修饰 N-glycosylated; indirectly required for FSH-binding, possibly via a conformational change that

allows high affinity binding of hormone.

细**胞定位** Cell membrane.

图片



All lanes: Anti-FSH-R antibody (ab75200) at 1/500 dilution

Lane 1 : Transfected Jurkat cell extract

Lane 2: Transfected HUVEC cell extract

Lane 3: Transfected Jurkat cell extract with immunising peptide at

10 µg

Lysates/proteins at 10 µg per lane.

Predicted band size: 78 kDa
Observed band size: 78 kDa

Additional bands at: 85 kDa. We are unsure as to the identity of

these extra bands.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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- Extensive multi-media technical resources to help you
- · We investigate all quality concerns to ensure our products perform to the highest standards

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