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10-7600: Monoclonal Antibody to FSH Receptor (Clone: ABM5D29)

Clonality: Monoclonal **Clone Name:** ABM5D29 Application: FACS.WB Reactivity: Human Gene: **FSHR** Gene ID: 2492 **Uniprot ID:** P23945 Format: Purified

Alternative Name: Follicle-stimulating hormone receptor, FSH-R, Follitropin receptor

Isotype: Mouse IgG1, kappa

Immunogen Information: A partial length recombinant protein (a.a 03-200) of FSH Receptor was used as the

immunogen for this antibody.

Description

FSH receptor is a member of the G protein coupled receptor family consisting of 9 introns and 10 exons and a promoter region and is located on chromosome 2p21. FSH receptor was shown to be expressed selectively on the luminal surface of tumor blood vessels. A general characteristic of the blood vessels that express the endothelial FSH receptor is that they are located at the periphery of the tumors. Many studies showed that FSH receptor polymorphisms at positions 307 and 680 may be even more relevant for clinical practice for the reasons of regulating the ovarian reaction to hormone, controlling ovarian hyperstimulation, changing the menstrual cycle, and causing POF (Premature Ovarian Failure) and PCOS (Polycystic Ovary Syndrome).

Product Info

Amount : $25 \mu g / 100 \mu g$

Purification: Protein G Chromatography

Content: 25 μg in 50 μl/100 μg in 200 μl PBS containing 0.05% BSA and 0.05% sodium azide. Sodium

azide is highly toxic.

Storage condition : Store the antibody at 4°C, stable for 6 months. For long-term storage, store at -20°C. Avoid

repeated freeze and thaw cycles.

Application Note

WB: 2-4 μg/ml, FACS: 0.5-1 μg/10⁶



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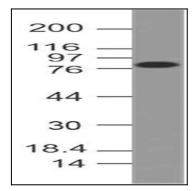


Fig-1: Western blot analysis of FSH Receptor. Anti-FSH Receptor antibody (Clone: ABM5D29) was used at 2 μ g/ml on Hela lysate.

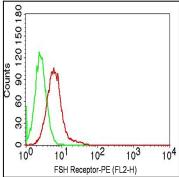


Fig-2: Intracellular flow analysis of FSH receptor antibody on A431 cells using 0.5 μ g/10^6 cells (Clone: ABM5D29). Green represents isotype control; red represents anti-FSH receptor antibody. Goat anti-mouse PE conjugate was used as secondary antibody. (Cells were fixed with 4% paraformaldehyde for 10 min and washed with PBS by centrifuging at 1100 for 5 min followed by permeabilization for 20 min and washed again as mentioned above. Then cell were incubated with primary antibody for 45 min. and after washing the cells twice in PBS, incubated with conjugated secondary antibody for 30 min. Data acquisition was done after washing twice with PBS as mentioned above).

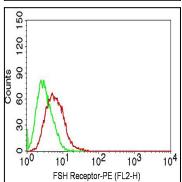


Fig-3: Intracellular flow analysis of FSH receptor antibody on HeLa cells using 0.5 $\mu g/10^{\circ}6$ cells (Clone: ABM5D29). Green represents isotype control; red represents anti-FSH receptor antibody. Goat anti-mouse PE conjugate was used as secondary. (Cells were fixed with 4% paraformaldehyde for 10 min and washed with PBS by centrifuging at 1100 for 5 min followed by permeabilization for 20 min and washed again as mentioned above. Then cell were incubated with primary antibody for 45 min. and after washing the cells twice in PBS, incubated with conjugated secondary antibody for 30 min. Data acquisition was done after washing twice with PBS as mentioned above).