

MULTISCREEN TM β -ARRESTIN2 STABLE CELL LINE HUMAN RECOMBINANT GPR35 RECEPTOR

PRODUCT INFORMATION

Catalog Number: CA1523-1

Lot Number: CA1523-1-032819

Quantity: 1 vial (2 x 10⁶) frozen cells

Freeze Medium: Cellbanker 2 (Amsbio

11891)

Host cell: CHO-K1

Transfection: Expression vector containing full-length human GPR35 cDNA (GenBank Accession Number NM_001195381.1) with FLAG tag sequence at N-terminus and ARRB2 cDNA (GenBank Accession Number NM_004313.3)

Recommended Storage: Liquid nitrogen upon receiving

Propagation Medium: DMEM/F12, 10% FBS, 10 μg/mL puromycin, 800

μg/mL G418

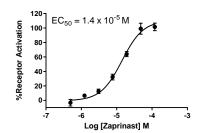
Stability: In progress

Data sheet

Background: GPR35 is an orphan receptor and expressed in various tissues including stomach, gastrointestinal tissues, and several types of immune cells. Upregulation of GPR35 has been found in human mast cells upon stimulation with IgE antibodies, human macrophages treated with the environmental contaminant polycyclic aromatic hydrocarbon benzo[a]pyrene, failing heart cells, and gastric cancer cells. Known agonists of the orphan receptor GPR35 are kynurenic acid, zaprinast, 5-nitro-2-(3-phenylproplyamino) benzoic acid, and lysophosphatidic acids.

Application: Functional assays

Figure 1



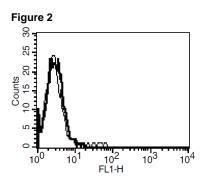


Figure 1. Dose-dependent inhibition of forskolin-stimulated intracellular cAMP accumulation upon treatment with ligand, measured with Multiscreen™ TR-FRET cAMP 1.0 No Wash Assay Kit (Multispan MSCM01). **Figure 2.** Receptor expression on cell surface measured by flow cytometry (FACS) using an anti-FLAG antibody. Thin line: parental cells; thick line: receptor-expressing cells.

References:

Wang JH et al. (2006) Kynurenic Acid as a Ligand for Orphan G Protein-coupled Receptor GPR35. J Biol Chem 281:22021-22028.

MacKenzie1, AE, et al. (2011) GPR35 as a novel therapeutic target. Frontiers in Endocrinology, 2:1-10.

Zhao PW, et al. (2010) Targeting of the Orphan Receptor GPR35 by Pamoic Acid: A Potent Activator of Extracellular Signal-Regulated Kinase and beta-Arrestin2 with Antinociceptive Activity. *Mol Pharm* 78:560–568.

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