

# PathHunter® eXpress PTGFR U2OS β-Arrestin GPCR Assay

Catalog Number: 93-0936E3 Lot Number: See Vial

Contents: 1 x 10<sup>6</sup> cells per vial in 0.1 mL

### **Background**

PathHunter eXpress  $\beta$ -Arrestin GPCR cells are engineered to co-express the ProLink<sup>TM</sup> (PK) tagged GPCR and the Enzyme Acceptor (EA) tagged  $\beta$ -Arrestin. Activation of the GPCR-PK induces  $\beta$ -Arrestin-EA recruitment, forcing complementation of the two  $\beta$ -galactosidase enzyme fragments (EA and PK). The resulting functional enzyme hydrolyzes substrate to generate a chemiluminescent signal. These cells have been modified to prevent long term propagation and expansion using a proprietary compound that has no apparent effect on assay performance.

#### **Product Information**

Target GPCR: PTGFR

**Description:** Prostaglandin F receptor (FP)

Receptor Family: Prostanoid

**Coupling**: Gq

Accession Number: NM\_000959.3

GPCR Species: Human

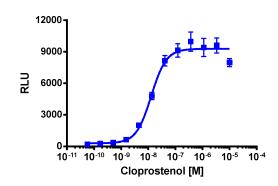
**β-Arrestin Isoform:**  $\beta$ -Arrestin-2

ProLink™ Tag: PK1
Cell Type: U2OS

**Storage:** Short term (<24 h): Store at -80°C; Long term (>24 h): Store in vapor phase of liquid nitrogen.

#### **Functional Performance**

Cells were plated in a 96-well plate and stimulated with a control agonist, using the assay conditions described below. Following stimulation, signal was detected according to the recommended protocol. Please refer below for information on control compounds.



Cell Number/Well: 10000

Control Agonist:(+/-)-Cloprostenol sodium salt hydrateCell Plating Reagent:AssayComplete™ Cell Plating 5 Reagent

Cell Incubation Time (Hours): 24

Agonist Incubation Time (Minutes): 90

Agonist Incubation Temperature (°C): 37

EC<sub>50</sub> for Agonist Stimulation (nM): 12.9

Signal:Background at Agonist E<sub>max</sub>: 49.8



## **Additional Ligand Information**

**Control Agonist:** (+/-)-Cloprostenol sodium salt hydrate

Vendor: Eurofins DiscoverX® (Catalog No. 92-1180)

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