

PathHunter[®] eXpress HTR1A CHO-K1 β-Arrestin GPCR Assay

Catalog Number: 93-0696E2

Lot Number: See Vial

Contents: 1 x 10⁶ cells per vial in 0.1 mL

Background

PathHunter eXpress β -Arrestin GPCR cells are engineered to co-express the ProLinkTM (PK) tagged GPCR and the Enzyme Acceptor (EA) tagged β -Arrestin. Activation of the GPCR-PK induces β -Arrestin-EA recruitment, forcing complementation of the two β -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.

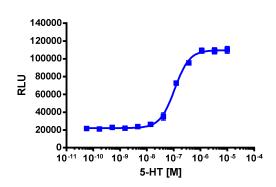
Product Information

Target GPCR:	HTR1A
Description:	5-hydroxytryptamine receptor 1A
Receptor Family:	5-Hydroxytryptamine
Coupling:	Gi/Go
Accession Number:	NM_000524.2
GPCR Species:	Human
β-Arrestin Isoform:	β-Arrestin-2
ProLink™ Tag:	ARMS2-PK2
Cell Type:	CHO-K1
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:



Control Agonist:	Serotonin/5-HT
Cell Plating Reagent:	AssayComplete™ Cell Plating 0 Reagent
Cell Incubation Time (Hours):	24
Agonist Incubation Time (Mine	utes): 90
Agonist Incubation Temperate	ure (°C): 37
EC ₅₀ for Agonist Stimulation (n M): 109.1
Signal:Background at Agonist	t E _{max} : 5.0

10000



Additional Ligand Information

Control Agonist: Serotonin/5-HT **Vendor:** Eurofins DiscoverX[®] (Catalog No. 92-1108)

Additional Prolink[™] Tag Description

PK2 is a slight variant of PK1 and has been shown to enhance EFC. ARMS (Arrestin Recruitment Modulating Sequence) is an 18-21 amino acid spacer between the GPCR and the PK tag and has been shown to enhance β -Arrestin recruitment.

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