

PathHunter[®] eXpress DRD2S CHO-K1 β-Arrestin GPCR Assay

Catalog Number: 93-0351E2 Lot Number: See Vial

1 x 10⁶ cells per vial in 0.1 mL Contents:

Background

PathHunter eXpress β-Arrestin GPCR cells are engineered to co-express the ProLink™ (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

DRD2S **Target GPCR:**

Description: Dopamine Receptor D2 (short isoform)

Receptor Family: Dopamine

Gi/Go Coupling:

Accession Number: NM 016574.3

GPCR Species: Human

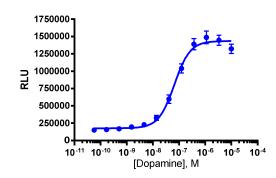
β-Arrestin Isoform: β-Arrestin-2 ProLink™ Tag: ARMS2-PK2

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:	10000
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Control Agonist: Dopamine HCI

AssayComplete™ Cell Plating 0 Reagent **Cell Plating Reagent:**

Cell Incubation Time (Hours): 48 90 Agonist Incubation Time (Minutes): 37 Agonist Incubation Temperature (°C): EC₅₀ for Agonist Stimulation (nM): 65.8 Signal:Background at Agonist E_{max}: 20.9

93-0351E2



Additional Ligand Information

Control Agonist: Dopamine HCl

Vendor: Eurofins DiscoverX® (Catalog No. 92-1161)

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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