

## **PRODUCT DATASHEET**

## Ready-to-Assay<sup>™</sup> CCR1 Chemokine Receptor Frozen Cells

### CATALOG NUMBER: HTS005RTA

**CONTENTS**: Pack contains 2 vials of mycoplasma-free cells, 1 ml per vial. Fifty (50) mL of Media Component. **STORAGE**: Vials are to be stored in liquid  $N_2$ . Media Component at 4°C (-20°C for prolonged storage).

## BACKGROUND

Ready-to-Assay<sup>™</sup> GPCR frozen cells are designed for simple, rapid calcium assays with no requirement for intensive cell culturing. Eurofins Discovery Services has optimized the freezing conditions to provide cells with high viability and functionality post-thaw. The user simply thaws the cells and resuspends them in media, dispenses cell suspension into assay plates and, following overnight recovery, assays for calcium response.

CCR1 is a GPCR that binds to a variety of CC ligands, including MIP-1 $\alpha$ , RANTES, MCP-3, HCC-1, HCC-2, HCC-4, and MPIF-1 (Olson and Ley, 2002). Lymphocytes, macrophages, dendritic cells, and GM-CSF-activated neutrophils express CCR1 (Kaufmann *et al.*, 2001; Cheng *et al.*, 2001). Two selective, non-peptide small molecule antagonists of CCR1, BX-471 and CP-481,715, have been synthesized (Gladue *et al.*, 2003; Liang *et al.*, 2000). Pharmacological and genetic targeting of CCR1, either alone or in combination with cyclosporin A, reduces cardiac and renal allograft rejection (Gao *et al.*, 2000; Horuk *et al.*, 2001a; Horuk *et al.*, 2001b), allergic encephalomyelitis (Liang *et al.*, 2000), and renal fibrosis (Anders *et al.*, 2002) in experimental models. Cloned human CCR1-expressing cell line is made in the Chem-1 host, which supports high levels of recombinant CCR1 expression on the cell surface and contains high levels of the promiscuous G protein G $\alpha$ 15 to couple the receptor to the calcium signaling pathway. Thus, the cell line is an ideal tool for screening for agonists, antagonists and modulators at CCR1.

# **USE RESTRICTIONS**

Please see User Agreement (Label License) for further details. One such restriction is that the contents of the supplied vial(s) are limited to a single use and shall not be propagated and/or re-frozen by licensee.

## WARNINGS

For Research Use Only; Not for Use in Diagnostic Procedures Not for Animal or Human Consumption

#### GMO

This product contains genetically modified organisms. Este producto contiene organismos genéticamente modificados. Questo prodotto contiene degli organismi geneticamente modificati. Dieses Produkt enthält genetisch modifizierte Organismen. Ce produit contient organismes génétiquement des modifiés. Dit product bevat genetisch gewijzigde organismen. Tämä tuote sisältää geneettisesti muutettuja organismeja. Denna produkt innehåller genetiskt ändrade organismer.

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## **APPLICATIONS**

Calcium Flux Assays

#### **APPLICATION DATA**

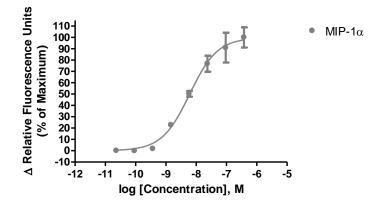


Figure 1. Representative data for activation of CCR1 receptor. Calcium flux in CCR1–expressing Chem-1 cell line induced by MIP-1α. CCR1–expressing Chem-1 cells were loaded with a calcium dye, and calcium flux in response to the indicated ligand, 4-fold serial dilution with each concentration performed in duplicate, was determined on a Molecular Devices FLIPR<sup>TETRA</sup> with ICCD camera. Maximal fluorescence signal obtained in this experiment was 10,000 RLU (Relative Light Units).

Table 1. Comparison of EC<sub>50</sub> values of CCR1-expressing Chem-1 cells with values described in the literature.

| LIGAND | ASSAY        | POTENCY (nM) | REFERENCE              |
|--------|--------------|--------------|------------------------|
| MIP-1α | Calcium Flux | 6            | Eurofins Internal Data |

# ASSAY SETUP

- 1. Immediately upon receipt, thaw cells or place cells in liquid nitrogen.
- 2. Thaw cells rapidly by removing from liquid nitrogen and immediately immersing in a 37°C water bath. Immediately after ice has thawed, sterilize the exterior of the vial with 70% ethanol.
- 3. Add 1mL of pre-warmed Media Component to each vial of cells. Place contents from two vials into a 15 mL conical tube and bring the volume to 10 mL of Media Component.
- 4. Centrifuge the cell suspension at 190 x g for four minutes
- 5. Remove supernatant and add 10.5 mL of pre-warmed Media Component to resuspend the cell pellet.
- Seed cell suspension into appropriate assay microplate (100 μL/well for 96-well plate, 25 μL/well for 384-well plate).
- 7. When seeding is complete, place the assay plate at room temperature for 30 minutes.
- 8. Move assay plate to a humidified 37°C 5% CO2 incubator for 24 hours.
- After 24 hour incubation, remove assay plate from the incubator and wash sufficiently with Hank's Balanced Salt Solution (HBSS) supplemented with 20mM HEPES, 2.5mM Probenecid at pH 7.4 to remove all trace of Media Component.



- Prepare Fluo-8, AM (AAT Bioquest: 21080) Ca<sup>2+</sup> dye by dissolving 1mg of Fluo-8 NW in 200 μL of DMSO. Once dissolved place 10 μL of Fluo-8 NW Ca<sup>2+</sup> dye solution into 10 mL of HBSS 20mM HEPES, 2.5mM Probenecid pH 7.4 buffer and apply to assay microplate (Ca<sup>2+</sup> dye at 10 μL /10 mL is sufficient for loading one (1) microplate).
- 11. Set-up FLIPR to dispense 3x ligand to appropriate wells in the assay plate. Set excitation wavelength at 470-495 nm (FLIPR<sup>TETRA</sup>) or 485 nm (FLIPR1, FLIPR2, FLIPR3) and emission wavelength at 515-565 nm (FLIPR<sup>TETRA</sup>) or emission filter for Ca<sup>2+</sup> dyes (FLIPR1, FLIPR2, FLIPR3). Set pipet tip height to 5 μL below liquid level and dispense rate to 75 μL/sec (96-well format) or 50 μL/sec (384-well format). Set up plate layout and tip layout for each individual experiment. Set time course for 180 seconds, with ligand addition at 10 seconds.
- 12. Ligands are prepared in non-binding surface Corning plates (Corning 3605 96-well or Corning 3574 384well).
- 13. After the run is complete, negative control correction is applied and data analyzed utilizing the maximum statistic.

#### **ASSAY MATERIALS**

| Description  | Supplier and Product Number           |
|--|---------------------------------------|
| HBSS   | Hyclone: SH30268.02                   |
| HEPES 1M Stock                                     | EMD Millipore.: TMS-003-C             |
| Probenicid   | Sigma: P8761                          |
| Quest Fluo-8™, AM                                  | AAT Bioquest: 21080                   |
| MIP-1a ligand                                      | Peprotech: 300-08                     |
| Non-binding white plates (for ligand prep)         | Corning: 3605(96-well)/3574(384-well) |
| Black (clear bottom) tissue-culture treated plates | Corning: 3904(96-well)/3712(384-well) |

#### **FLIPR SETTINGS**

Settings for FLIPR<sup>TETRA</sup>® with ICCD camera option

| Option          | Setting                          |
|-----------------|----------------------------------|
| Read Mode       | Fluorescence                     |
| Ex/Em           | Ex470_495 / Em515_575            |
| Camera Gain     | 2000                             |
| Gate Open       | 6 %                              |
| Exposure Time   | 0.53                             |
| Read Interval   | 1s                               |
| Dispense Volume | 50 µl (25 µl for 384-well)       |
| Dispense Height | 25 μl (50 μl for 384-well)       |
| Dispense Speed  | 75 μl L/sec (50 μl for 384-well) |
| Expel Volume    | Ο μΙ                             |
| Analysis        | Subtract Bias Sample 1           |

# **HOST CELL**

Chem-1, an adherent rat hematopoietic cell line expressing endogenous  $G\alpha 15$  protein.

# **EXONGENOUS GENE EXPRESSION**

CCR1 cDNA (Accession Number: L09230) expressed from a proprietary plasmid.



## **RELATED PRODUCTS**

| PRODUCT NUMBER | DESCRIPTION  |
|----------------|--|
| HTSCHEM-1RTA   | Ready-to-Assay™ Chem-1 host frozen cells (control cells)       |
| HTS005M        | ChemiScreen <sup>™</sup> CCR1 Chemokine receptor membrane prep |

## REFERENCES

- 1. Anders, H.J., *et al.* (2002) A chemokine receptor CCR-1 antagonist reduces renal fibrosis after unilateral ureter ligation. *J. Clin. Invest.* 109: 251-9.
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- 3. Gao, W., *et al.* (2000) Targeting of the chemokine recpetor CCR1 suppresses development of acute and chronic cardiac allograft rejection. *J. Clin. Invest.* 105: 35-44.
- 4. Gladue, R.P., *et al.* (2003) CP-481,715, a potent and selective CCR1 antagonist with potential therapeutic implications for inflammatory diseases. *J. Biol. Chem.* 278: 40473-80.
- 5. Horuk, R., et al. (2001a) CCR1-specific non-peptide antagonist: efficacy in a rabbit allograft rejection model. *Immunol. Lett.* 76: 193-201.
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- 7. Kaufmann, A., *et al.* (2001) Increase of CCR1 and CCR5 expression and enhanced functional response to MIP-1 alpha during differentiation of human moncytes to macrophages. *J. Leukoc. Biol.* 69: 248-52.
- 8. Liang, M., *et al.* (2000) Identification and characterization of a potent, selective, and orally active antagonist of the CC chemokine receptor-1. *J. Biol. Chem.* 275: 19000-8.
- 9. Olson, T.S. and Ley, K. (2002) Chemokines and chemokine receptors in leukocyte trafficking. *Am. J. Physiol. Regul. Integr. Comp. Physiol.* 283: R7-R28.

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