

### PRODUCT DATASHEET

# ChemiScreen™ Chem-6 Wild-type Parental Control

CATALOG NUMBER: HTSCHEM-6

**CONTENTS**: Pack contains 2 vials of mycoplasma-free cells, 1 ml per vial.

**STORAGE**: Vials are to be stored in liquid N<sub>2</sub>.

#### **BACKGROUND**

Eurofins Discovery Services' Chem-6 Parental Control Frozen Cells are designed for use as a negative control alongside our GPCR-expressing Ready-to-Assay Frozen Cells in the Chem-6, Chem-7, and Chem-8 backgrounds. Chem-6 cells lack endogenous expression of most GPCRs, although they endogenously express trypsin and thrombin receptors that may be used to confirm responsiveness of the cells in calcium assays. The Chem-6 cell line is adherent to tissue culture-treated surfaces and contains high levels of the promiscuous G protein  $G\alpha15$  to couple many classes of GPCRs to the calcium signaling pathway.

#### **USE RESTRICTIONS**

Please see Limited Use Label License Agreement (Label License Agreement) for further details.

### **WARNINGS**

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

#### GMC

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.



### **APPLICATIONS**

Calcium Flux Assays; cAMP

### **CELL CULTURE**

Table 1. Recommended Cell Culture Reagents (not provided)

Description	Component	Concentration	Supplier and Product Number
<b>Basal Medium</b>	F-12 Kaighn's	-	Hyclone: SH3052601
	Fetal Bovine Serum (FBS)	10%	Hyclone: SH30070.03
Selection Medium	Basal Medium (see above)	-	
Dissociation	Sterile PBS	-	Hyclone: SH30028.03
	0.25% Trypsin-EDTA	-	Hyclone: SH30042.01
CryoMedium	Basal Medium (see above)	40%	
	Fetal Bovine Serum (FBS)	50%	Hyclone: SH30070.03
	Dimethyl Sulfoxide (DMSO)	10%	Sigma: D2650

### **Cell handling**

- 1. Upon receipt, directly place cells in liquid nitrogen storage. Consistent cryopreservation is essential for culture integrity.
- 2. Prepare Basal Medium. Prepare 37°C Water Bath. Thaw cells rapidly by removing from liquid nitrogen, and immediately immersing in a 37°C water bath, until 90% thawed. Immediately sterilize the exterior of the vial with 70% ethanol.
- 3. Add vial contents to 15 mL Basal Medium in T75 Tissue Culture Treated Flask. Gently swirl flask and place in a humidified, tissue culture incubator, 37°C, 5% CO<sub>2</sub>.
- 4. 18-24 Hours Post–Thaw, all live cells should be attached. Viability of the cells is expected to be 60-90%, At this time, exchange Basal Medium with Selection Medium.
- 5. When cells are approximately 80% confluent, passage the cells. It is suggested that user expand culture to create >20 vial Master Cell Bank at low passage number. *Cells should be maintained at less than 80% confluency for optimal assay results*.
- 6. Cell Dissociation: Aspirate Culture Medium. Gently wash with 1x Volume PBS. Add 0.1x Volume Warm Trypsin-EDTA. Incubate 4 min, 37°C, until cells dislodge. *If cells do not round up, place in 37°C incubator for additional 2 min.* Neutralize Trypsin and collect cells in 1x Volume Basal Medium.
- 7. Seed Cells for expansion of culture. It is recommended that cell lines are passaged at least once before use in assays.

Table 3. Cell Culture Seeding Suggestions: User should define based on research needs.

Flask Size (cm <sup>2</sup> )	Volume (mL)	Total Cell Number (x10 <sup>6</sup> )	Growth Period (hrs)
T75	15	5.0	24
T75	15	2.0	48
T75	15	0.45	72



### **Discovery Services**

### **ASSAY SETUP**

#### **ASSAY MATERIALS**

Description	Supplier and Product Number		
HBSS	Hyclone: SH3026802		
HEPES 1M Stock	EMD Millipore: TMS-003-C		
Probenicid	Sigma: P8761		
Quest Fluo-8 <sup>™</sup> , AM	AAT Bioquest: 21080		
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 TETRA® 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	0 μΙ
Analysis	Subtract Bias Sample 1

### **Assay Protocol – Fluorescence**

- 1. Dissociate Culture as Recommended. Collect in Basal Medium. Document Cell Count and Viability
- 2. Centrifuge the cell suspension at 190 x a for six min
- 3. Remove supernatant. Gently resuspend the cell pellet in Basal Medium. *It is suggested that end user optimize cell plating based on individual formats.* (Default: Resuspend in volume to achieve 5x10<sup>5</sup>cells/ml (i.e, if collected 5e6 TC, 5e6/5e5/ml =10 mL volume)
- 4. Seed cell suspension into black, clear bottom plate (100 μL/well for 96-well plate). When seeding is complete, place the assay plate at room temperature for 30 min.
- 5. Move assay plate to a humidified 37°C 5% CO<sub>2</sub> incubator for 18-24 h.
- 6. Next day, prepare Assay buffer (HBSS, 20mM HEPES, 2.5 mM Probenicid, pH 7.4) and Loading buffer (Assay buffer with 5 mM Fluo8 Dye). *Note: Please prepare Fluo8 stock according to Manufacturer's Recommendations*
- 7. Remove plate from incubator and quickly invert plate on an absorbent pad and blot to remove all Media Component.
- 8. Add Loading buffer to assay plate (100  $\mu$ L/well for 96-well plate). Incubate plate for 1.5 h at room temperature, protected from light.
- 9. Prepare ligands in assay buffer at 3x final concentration in non-binding plates. Use Buffer Only Control Wells for Background Subtraction.
- 10. Create protocol for ligand addition. Please refer to FLIPR<sup>TETRA</sup>® settings provided in Table 2. Set time course for 180 s. with ligand addition at 10 s.
- 11. After the run is complete, apply subtract bias on sample 1. We recommend using Negative Control Correction with Buffer Only Wells. Export data to according to research needs. For most Calcium Flux analysis using Export of Max Signal to end of run is sufficient.



### **HOST CELL**

Chem-6, an adherent Chinese Hamster Ovary cell line expressing endogenous Gα15 protein.

### **RELATED PRODUCTS**

PRODUCT NUMBER DESCRIPTION

HTSCHEM-6RTA Ready-to-Assay™ ChemiScreen™ Chem-6 Wild-Type Parental Control

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