

# **CERTIFICATE OF ANALYSIS**

### **Product Information**

Product Name HEK293/CRE-luciferase/GLP1R

 Cat. No.
 M00562

 Lot No.
 R10031809

 Host Cell:
 HEK293

 Target Gene:
 GLP1R

Quantity: 2 vials of frozen cells, > 1x10<sup>6</sup> cells/vial

Shipping Condition: Dry Ice

Recommended Liquid Nitrogen

Storage Condition:

### **Stable Cell Line Information**

Recommended Cell Culture Medium: DMEM, 10% FBS, 400 µg/ml G418, 100 µg/ml Hygromycin B

Freeze Medium: 90% FBS, 10% (V/V) DMSO

Application: Functional assay for HEK293/CRE-luciferase/GLP1R

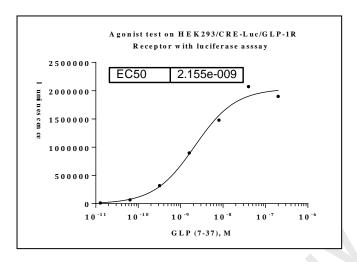
| Test Item      | Specification    | Result                     |
|----------------|------------------|----------------------------|
| Mycoplasma     | Not detected*    | Not detected*              |
| Cell viability | >90%             | 95%                        |
| Function assay | cAMP assay       | EC <sub>50</sub> =13.42 nM |
| Function assay | Luciferase assay | EC <sub>50</sub> =2.16 nM  |

<sup>\*</sup> The mycoplasma test was performed with MycoAlert™ PLUS Mycoplasma Detection Kit of Lonza.



## **Appendix**

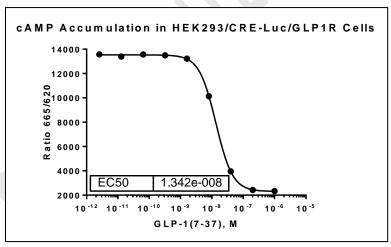
### 1. Luciferase assay



GLP-1(7-37)-induced luciferase expression in HEK293/CRE-Luc/GLP1R cells. After stimulation by agonist GLP-1(7-37), the cells are determined with One-Glo™ Luciferase Assay System and the relative luminescence units (RLU) were recorded by PheraStar.

The RLU were plotted against the log of the cumulative doses of GLP-1(7-37) (Mean  $\pm$  SD, n = 3). The EC<sub>50</sub> of GLP-1(7-37) on HEK293/CRE-Luc/GLP-1R cells was 2.16 nM. The S/B of the assay was 74.

### 2. cAMP assay



GLP-1(7-37) induced concentration-dependent stimulation of intracellular cAMP accumulation in HEK293/CRE-luciferase/GLP1R cells.  $EC_{50}=1.342\times10^{-8}$  M.

#### Caution

For research use only. Not intended for household use. If you have any questions about the Certificate of Analysis, please contact our customer service representative at 1-877-436-7274 (Toll-Free), or 1-732-885-9188.

Certified by: Vary Tue

Date: 11/11/2019

Department of New Technology