(FOR RESEARCH USE ONLY, DO NOT USE IT IN CLINICAL DIAGNOSIS!)

Catalog No: E-BC-F074

Specification: 48T(46 samples)/96T(94 samples)

Measuring instrument: Chemiluminescence immunoassay analyzer

# Elabscience® Dual Luciferase Reporter Gene Luminescence Assay Kit (Glow Type)

This manual must be read attentively and completely before using this product.

If you have any problem, please contact our Technical Service Center for help:

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Tell: 1-832-243-6086

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Email: techsupport@elabscience.com

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Please kindly provide us the lot number (on the outside of the box) of the kit for more efficient service.

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#### Intended use

This kit can be used to measure firefly luciferase reporter gene and renilla luciferase reporter gene expressed level in cell samples.

### **Detection principle**

Dual luciferase reporter gene is used to detect the luminescence reaction produced by luciferase catalyzed substrates. The detection system includes firefly luciferase reporter gene and renilla luciferase reporter gene. As an internal parameter of transfection, the renilla luciferase can normalize the experimental data. Firefly luciferase reporter gene refers to a reporting system that uses fluorescein as substrate to detect fluorescein luciferase activity. The reaction can occur in the presence of ATP, magnesium ions and oxygen, resulting in yellow-green fluorescence signals. The advantage of the reporter gene is that the expression product of the reporter gene has detectable enzyme activity as soon as the translation is completed. Renilla luciferase reporter gene refers to a reporting system that uses coelenterazine as substrate to detect the activity of renilla luciferase. This reaction does not depend on ATP to participate in the enzymatic reaction. During the reaction, a blue fluorescence signal is generated.

The reaction principle of this kit is that in the presence of oxygen, ATP and magnesium ions at the same time, firefly luciferin and coelenterazine are catalyzed and oxidized successively by firefly luciferase and renilla luciferase in the samples to produce different fluorescence signals, and the expression levels of firefly luciferase and renilla luciferase in the samples can be detected by chemiluminescence immunoassay analyzer.

#### Kit components & storage

Item	Component	Size 1 (48 T)	Size 2 (96 T)	Storage
Reagent 1	Firefly Luciferase Buffer Solution	7 mL × 1 vial	14 mL × 1 vial	-20°C, 12 months
Reagent 2	Firefly Luciferase Substrate	Powder × 2 vials	Powder × 4 vials	-20°C, 12 months shading light
Reagent 3	Renilla Luciferase Buffer Solution	6 mL × 1 vial	12 mL × 1 vial	-20°C, 12 months
Reagent 4	Renilla Luciferase Substrate	0.12 mL × 1 vial	0.24 mL × 1 vial	-20°C, 12 months shading light
	Black Clear-bottom Culture Plate	96 wells		No requirement
	Plate Sealer	2 pieces		
	Sample Layout Sheet	1 piece		

Note: The reagents must be stored strictly according to the preservation conditions in the above table. The reagents in different kits cannot be mixed with each other. For a small volume of reagents, please centrifuge before use, so as not to obtain sufficient amount of reagents.

## Materials prepared by users

#### Instruments:

Chemiluminescence immunoassay analyzer or multifunctional microplate reader (with the function of detecting luminescence)

## **Reagent preparation**

- ① Keep renilla luciferase substrate on ice to thawing. Equilibrate other reagents to 25℃ before use.
- ② The preparation of firefly luciferase substrate working solution: Dissolve one vial of firefly luciferase substrate with 3 mL of firefly luciferase buffer solution, mix well to dissolve. Aliquoted storage at -20°C for 1 month protected from light.

## The key points of the assay

The firefly luciferase substrate working solution can be aliquoted storage at -20°C, and avoid repeated freeze/thaw cycles is advised.

### Sample preparation

#### Sample preparation

#### Cell samples:

The cells were inoculated and designed according to the following groups:

Blank group: The cells without transfection treatment;

Control group: The cells were transfected with plasmids without drug stimulation.

Experiment group: The cells were transfected with plasmids and stimulated with drugs according to experimental design.

### **Operating steps**

- ① Add 100  $\mu$ L of firefly luciferase substrate working solution into black clear-bottom culture plate, and mix fully with chemiluminescence immunoassay analyzer. Measure the luminescence values of each well, as  $F_1$ .
- ② Sample well: add 100  $\mu$ L of renilla luciferase substrate working solution into the sample wells, and mix fully with chemiluminescence immunoassay analyzer. Measure the luminescence values of each well, as  $F_2$ .

**Note:**  $F_1$  is the expression level of luciferase in firefly and  $F_2$  is the expression level of luciferase in renilla.

#### Calculation

#### Relative Light Units (RLUs) formula:

RLUs = 
$$\frac{\Delta F_1}{\Delta F_2}$$

#### [Note]

 $\Delta F_1$ : The  $F_1$  values of experiment group - The  $F_1$  values of blank group.

 $\Delta F_2$ : The  $F_2$  values of experiment group - The  $F_2$  values of blank group.

#### Statement

- 1. This assay kit is for Research Use Only. We will not response for any arising problems or legal responsibilities causing by using the kit for clinical diagnosis or other purpose.
- Please read the instructions carefully and adjust the instruments before the experiments. Please follow the instructions strictly during the experiments.
- 3. Protection methods must be taken by wearing lab coat and latex gloves.
- 4. If the concentration of substance is not within the detection range exactly, an extra dilution or concentration should be taken for the sample.
- 5. It is recommended to take a pre-test if your sample is not listed in the instruction book.
- 6. The experimental results are closely related to the situation of reagents, operations, environment and so on. Elabscience will guarantee the quality of the kits only, and NOT be responsible for the sample consumption caused by using the assay kits. It is better to calculate the possible usage of sample and reserve sufficient samples before use.