



Firefly Luciferase Assay Kit

Catalog Number: 30003-1 (150 assays)
30003-2 (1000 assays)

Description

Firefly luciferase is widely used as a reporter for studying gene regulation and function, and for pharmaceutical screening^{1,2}. It is a very sensitive genetic reporter due to the lack of any endogenous activity in mammalian cells or tissues^{3,4}. The *Firefly* luciferase is a 62,000 Dalton protein, which is active as a monomer and does not require subsequent processing for its activity. The enzyme catalyzes ATP-dependent D-luciferin oxidation by oxygen into oxyluciferin with emission of light centered on 560nm (Figure 1). As with many enzymes, *Firefly* luciferase follows Michaelis-Menten kinetics and, as a result, maximum light output is not achieved until the substrate and co-factors are present in large excess. When assayed under these conditions, light emitted from the reaction is directly proportional to the number of luciferase enzyme molecules. This *Firefly* luciferase assay kit is designed for detection and quantification of the *Firefly* luciferase reporter enzyme from cultured cells in a simple, efficient, and linear fashion (Figure 2).

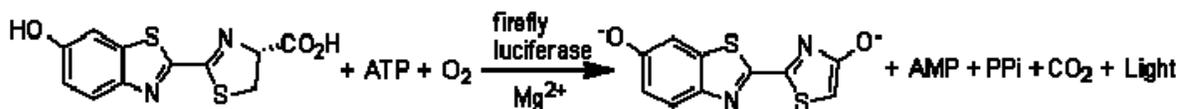


Figure 1. Bioluminescent reaction catalyzed by *Firefly* luciferase.

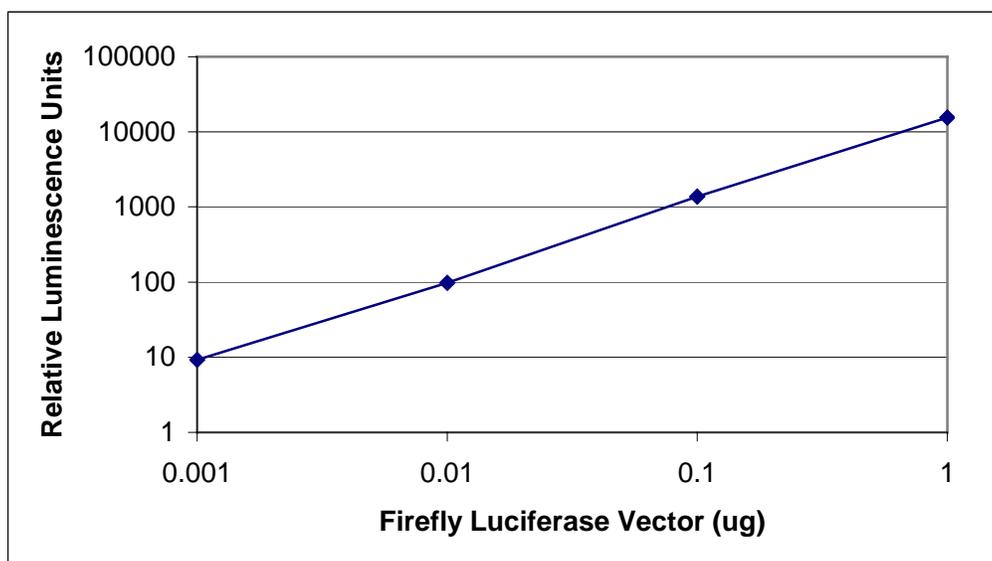


Figure 2. Dose response curve of transfected *Firefly* Luciferase gene. PC3 cells were transfected with 0.001µg, 0.01µg, 0.1µg, and 1µg pFL-CMV vector encoding *Firefly* luciferase gene by Fugene 6 (Roche) in 6-well cell culture dishes. pGL2 Basic vector (Promega) was used as a control and for normalizing total DNA vector level to 1µg per transfection. Twenty-four hours after transfection, cells were harvested using 500µL lysis buffer contained in Biotium's *Firefly* Luciferase Assay Kit. To assay luciferase activity, 20µL of lysate from each sample was then added to 100µL of assay buffer also in Biotium's *Firefly* Luciferase Assay Kit. Luminescence was measured on a luminometer (Turner Designs). Light emission was integrated over 10 seconds without initial pre-read delay.

Product Components

Firefly Luciferase Assay Kit, 30003-1 (150 assays)

| Catalog Number | Component |
|----------------|--|
| 99907 | 3 vials (1mg each) of D-Luciferin |
| 99923 | 15mL 5X <i>Firefly Luciferase Assay Lysis Buffer</i> |
| 99932 | 15mL <i>Firefly Luciferase Assay Buffer</i> |

Firefly Luciferase Assay Kit, 30003-2 (1000 assays)

| Catalog Number | Component |
|----------------|--|
| 99908 | 2 vials (10mg each) of D-Luciferin |
| 99923 | 30mL 5X <i>Firefly Luciferase Assay Lysis Buffer</i> |
| 99909 | 100mL <i>Firefly Luciferase Assay Buffer</i> |

Storage Conditions

Store *Firefly Luciferase Assay Kit* at -70°C . *Firefly luciferase assay solution* (Assay Buffer + Substrate) should be prepared fresh for each use. Avoid repeated freeze-thaw cycles. Aliquot *Firefly Luciferase Assay Buffer* for storage if necessary.

Preparation of Cell Lysates

A. Preparation of *Firefly Luciferase Lysis Buffer*

Firefly Luciferase Lysis Buffer 1X working solution is prepared by adding 1 volume of 5X *Firefly Luciferase Lysis Buffer* to 4 volumes of de-ionized water and mixing well. The 1X Lysis Buffer may be stored at 4°C for up to one month. Store the 5X *Firefly Luciferase Assay Lysis Buffer* at -20°C .

B. Lysis of Cells Cultured in Multiwell Plates

1. Remove growth medium from cultured cells and gently add a sufficient volume of phosphate buffered saline (PBS) to wash the surface of the culture vessel. Add *Firefly Luciferase Assay Lysis Buffer* to each well using the volume recommended below for each type of culture plate:

| Wells/plate | Lysis buffer/well |
|-------------|-------------------|
| 6 well | 500 μL |
| 12 well | 250 μL |
| 24 well | 100 μL |
| 48 well | 65 μL |
| 96 well | 20 μL |

Note for Cat# 30003-2: Sufficient *Firefly Lysis Buffer* is provided to perform 1000 assays with cells grown in 96 – 24 well plates. For applications requiring more lysis buffer (e.g. $>100\ \mu\text{L/well}$), additional 5X lysis buffer (Cat. # 99923, 2 X 15 mL) may be purchased separately.

2. Place the culture plates on a rocking platform or orbital shaker with gentle rocking/shaking to ensure complete and even coverage of the cell monolayer with 1X *Firefly Luciferase Lysis Buffer*. Rock the culture plates at room temperature for 15 minutes.

Note: Cultures that are overgrown are often more resistant to complete lysis and typically require an increased volume of *Firefly Luciferase Lysis Buffer* and/or an extended treatment period to ensure complete lysis. Lifting cells from the plate will facilitate the process of cell lysis.

3. Transfer the lysate to a tube or vial and place in 4°C for further assay. Although it is not necessary, the lysate can be cleared by centrifugation for 30 seconds at top speed in a refrigerated microcentrifuge and transferred into a new tube.

Firefly Luciferase Assay

A. Preparation of *Firefly Luciferase Assay Solution*

1. Thaw a bottle of *Firefly Luciferase Assay Buffer* and pipette a desired volume (5mL or 50mL) from the bottle into a new container.

2. Dissolve the supplied D-luciferin with the above assay buffer at 0.2mg/mL concentration in a clean container. 5 mL assay solution is prepared from each 1 mg D-luciferin vial. Similarly, 50 mL assay solution is prepared from each 10 mg D-luciferin vial. *Firefly Luciferase Assay Solution* should be prepared fresh and used within a day.

Note: *D-luciferin in Assay Buffer has limited stability. If you need less than 5mL or 50mL Luciferase Assay Solution as described in step 2, you may dissolve D-Luciferin in DI water as 10X or 50X stock solution and store it at -20°C or below for repeated use. The D-luciferin stock solution should be stable for at least one month, depending on the frequency of free-thaw cycle. A desired volume of the final Assay Solution can be prepared by diluting the stock solution with the supplied Assay Buffer to 0.2mg/mL D-luciferin.*

B. Standard Protocol

For manual luminometer:

1. Set up luminometer with appropriate parameters (delay time, integration time, sensitivity, etc.).
2. Add 100µL of *Firefly Luciferase Assay Solution* to the luminometer tube.
3. Add 20µL of cell lysate. Mix quickly by flicking the tube with a finger for thorough mixing.
3. Place tube in luminometer and initiate measurement. Luminescence is normally integrated over 10 seconds without delay. Other integration times may also be used.
4. If the luminometer is not connected to a printer or computer, record the *Firefly luciferase* activity measurement.
5. Discard the reaction tube, and proceed to the next *Firefly Luciferase Assay*.

For luminometer with injector:

1. Format the luminometer so that the injector dispenses 100µL. Prime the injector with *Firefly Luciferase Assay Solution*.
2. For each reaction, carefully add 20µL of cell lysate to an individual luminometer tube or to the wells of a multiwell plate.
3. Place the samples in a luminometer.
4. Initiate measurement. This action will cause *Firefly Luciferase Assay Solution* to be injected into the reaction vessel and the measurement to be subsequently taken. Luminescence is normally integrated over 10 seconds without delay. Other integration times may also be used.
5. Record the *Firefly luciferase* activity measurement.
6. If using a single tube luminometer, discard the reaction tube, and proceed to the next *Firefly Luciferase Assay* reaction. If using a plate luminometer, the luminometer will automatically begin injecting *Firefly Luciferase Assay Solution* into the next well indicated on the luminometer plate.

References

1. Alam, J. and J.L. Cook. 1990. Reporter genes: Application to the study of mammalian gene transcription. *Anal. Biochem.* 188:245-254.
2. Bronstein, I., et al. 1994. Chemiluminescent and bioluminescent reporter gene assays. *Anal. Biochem.* 219:169-181.
3. Gould, S.J. and S. Subramani. 1988. *Firefly luciferase* as a tool in molecular and cell biology. *Anal. Biochem.* 175:5-13.

4. Brasier, A.R., et al. 1989. Optimized use of the *Firefly* luciferase assay as a reporter gene in mammalian cell lines. *BioTechniques*. 7:1116-1122.

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