

Product Description

D-Luciferin, Potassium Salt is a substrate used for reporting gene expression of luciferase in bioluminescence imaging assays for live organisms. The specific mechanism of action is as follows: In the presence of ATP, magnesium ions, and oxygen, luciferase catalyzes the oxidation of D-Luciferin, Potassium Salt as a substrate, resulting in the production of a strong bioluminescent signal. In vivo imaging experiments involve the introduction of cells or mRNA expression vectors capable of expressing luciferase into experimental animals. Subsequently, D-Luciferin, Potassium Salt is injected, and the changes in luminescence intensity are detected using a small animal optical imaging system, thereby reflecting the level of luciferase expression. It can be used for tumor research, immunology and stem cell research, drug studies, cell/protein/mRNA labeling, protein-protein interactions, and more.

Components

Components	DD1210-01	DD1210-02	DD1210-03
D-Luciferin, Potassium Salt	10 mg	100 mg	1 g

Storage

Store at -30 ~ -15°C and protect from light, ship at ≤0°C.

Applications

It is applicable for bioluminescence imaging detection in plants and animals.

Self-provided Materials

Self-provided Reagents

DPBS (without Ca^{2+} , Mg^{2+}) ddH_2O

Other Materials and Equipments

Sprayer

1 ml Syringe

0.22 µm Filter Membrane

Optical Imaging System for Live Plants and Animals

Notes

- 1. Please wear laboratory coats, masks, and gloves in accordance with biosafety regulations, and dispose of experimental waste in accordance with medical waste disposal requirements.
- 2. This product should be stored sealed and protected from light. After opening and dissolving, it is prone to oxidation. Aliquot and store at -20°C or -80°C to minimize freeze-thaw cycles.
- 3. After thawing, the product should be dissolved on ice or at 4°C. Thawed D-Luciferin can be temporarily stored at 4°C or on ice.
- 4. For in vivo injection in small animals, filter sterilization is required. If the product accidentally splashes into the eyes, skin, or other body parts, rinse immediately with plenty of water.
- 5. During ATP detection, wear disposable gloves to avoid interference from exogenous ATP.
- 6. Use DPBS without calcium and magnesium ions when dissolving the product, as these ions can interfere with the luminescence reaction
- 7. Before in vivo imaging detection, it is recommended to conduct a pilot experiment to establish an in vivo luciferase kinetics curve to determine the signal plateau and detection time.

Experiment Process

Plant In Vivo Imaging Detection

- 1. Dissolve D-Luciferin, Potassium Salt in sterile ddH₂O to prepare a 30 mg/ml stock solution (100 200 ×), mix well. Use immediately or aliquot and store at -20°C in the dark, avoiding repeated freeze-thaw cycles.
- 2. Dilute the stock solution to a working concentration of 0.3 0.5 mg/ml with sterile ddH₂O.
- 3. Wet the underside of the leaves with the working solution, let stand in the dark at room temperature for 5 10 min before detection.

Animal In Vivo Imaging Detection

- 1. Prepare a 15 mg/ml luciferin stock solution in sterile DPBS (without Ca2+, Mg2+), mix well.
- 2. Filter sterilize using a 0.22 µm filter membrane, use immediately or aliquot and store at -20°C in the dark, avoiding repeated freeze-thaw cycles.
- 3. Administer intraperitoneally (i.p.) at a concentration of 150 mg/kg luciferin/body weight.
- 4. Perform imaging detection 10 15 min after injection (when the light signal reaches its strongest and most stable plateau).