



User Manual

DiR-Membrane EVs Labeling & Purification Kit (near-infrared red)

Cat. # EXOPDiR10-1
EXOPDiR20-1

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Storage and Application

【Storage】

The DiR-Membrane EVs Labeling & Purification Kit (near-infrared red) is shipped on ice, the components should be stored at recommended temperatures and protected from light. Properly stored kits are stable for 6 months. please read the instructions before use.

【Application】

For research use only. Not for use in diagnostic procedures.

Product Description

DiR is a near-infrared fluorescent, lipophilic carbocyanine and widely used as a lipophilic tracer. The dye uniformly labels cells via lateral diffusion in the plasma membrane. The fluorescence of free DiR is very weak, but greatly enhanced when incorporated into membranes. DiR emits red fluorescence when excited, its maximum excitation and emission wavelength is **750nm** and **780nm**, respectively.

Extracellular vesicles (EVs) are membrane-derived particles surrounded by a phospholipid bilayer that are released by cells. We use DiR to efficiently label EVs membranes. Then the excess unbound dyes are fast removed through our excellent Spin Columns, which separate molecules on the basis of the differences in size. Contaminant removal is easier and faster than traditional clean-up methods such as ultracentrifugation, spin filters.

With the better tissue permeability and low spontaneous fluorescence of cells and tissues in the near infrared region, the kit shows higher specificity and sensitivity than traditional visible dyes. It can be used to track the migration and distribution of labeled exosomes in vivo in real time.

List of Components

Specification : **20** reactions/Kit (Cat.#EXOPDiR20-1) ; **10** reactions/Kit (Cat.#EXOPDiR10-1)

| Item (Cat.#EXOPDiR20-1) | Volume | Storage Temp. |
|---|----------|---------------|
| DiR Labeling Dye* | 100µL | -20°C |
| Reaction Buffer | 1.0mL | 2-8°C |
| Spin Columns# within Collections Tubes | 20 sets | 2-8°C |
| 1.5mL Light-proof Microcentrifuge Tubes | 20 tubes | RT |
| Item (Cat.#EXOPDiR10-1) | Volume | Storage Temp. |
| DiR Labeling Dye* | 50µL | -20°C |
| Reaction Buffer | 500µL | 2-8°C |
| Spin Columns# within Collections Tubes | 10 sets | 2-8°C |

| | | |
|---|----------|----|
| 1.5mL Light-proof Microcentrifuge Tubes | 10 tubes | RT |
|---|----------|----|

* **Protect labeling dye from light.**

Keep the Spin Columns upright stand on end.

The sterilized PBS buffer is not provided in the Kit. Please prepare at least 600uL PBS buffer for each reaction.

General Information

1. This Kit can label exosomes of any resources, including cell culture supernatants and body fluids (such as serum, plasma, urine, CSF or saliva).
2. When extracting exosomes for Labeling, the minimum dosage of serum and plasma is 500 μ L, the urine is 10mL, and the cell culture supernatant is 5mL.
3. It is not recommended to use exosomes extracted by PEG precipitation method, which contains too many impurities. The exosomes extracted by ultracentrifugation, affinity method or our company's **Exosomes Extraction and Purification Kit** is preferable.
4. Heat the dye at 37 °C until dissolved completely when the dye has crystallized.
5. Fluorescent dyes have quenching problems, please protect dyes from light during operating.
6. For your safety and health, please wear a lab gown and wear a disposable glove when operating.
7. The whole procedure is non-aseptic, please filter the labeled exosomes through 0.45um filter membrane before performing the downstream experiments.

Protocols

Staining Protocol

1. Vortex the DiR Labeling Dye and then instantaneous centrifugation, add 5 μ L DiR Labeling Dye to 50 μ L Reaction Buffer, and mix well until the dye is dissolved completely.
2. Add 50 μ L exosomes into the dye mixture slowly and mix well.
3. Incubate the mixture for 30 minutes at 37 °C, pipet twice during the incubation

period.

! Protect the tubes from light.

Purification Protocol (remove excess unbound dye)

4. Prepare the Spin Column prior to application of your sample.
 - a) Open the cap of the Spin Column, aspirate preservative buffer from the top of the column with a micropipette and discard it, then remove the outlet plug of the column. In order to prevent drying of the column bed, proceed to the next step promptly.
 - b) equilibrate the column by adding 200 μ L sterilized PBS buffer (not provided) and spin down at 100 x g for 90 seconds. If any PBS remains above the top frit, repeat spin at the same speed with 10 seconds increments. Discard the eluate.
 - c) Repeat the procedure b) again.

! Do not spin at too high speed or for too long as this may desiccate or compress the resin and decrease the function of spin column.

5. Carefully apply 100 μ L exosome labeling preparation (from step 3 above) to the top of the column.

! The maximum capacity of the spin column is 100 μ L. Do not load samples more than 100 μ L.

6. Centrifuge at 100 x g for 90 seconds. Discard the eluate.
7. Place the column into a fresh 1.5 mL Light-proof Microcentrifuge Tubes (provided). Apply 200 μ L PBS buffer (not provided) to the top of the column.
8. Centrifuge at 100 x g for 90 seconds. The 200 μ L eluate contains the labeled exosomes.
9. According to your needs, dilute the labeled exosomes with corresponding medium or not. Then filter the labeled exosomes through 0.45 μ m filter membrane before performing the downstream experiments.

Example Data and Applications

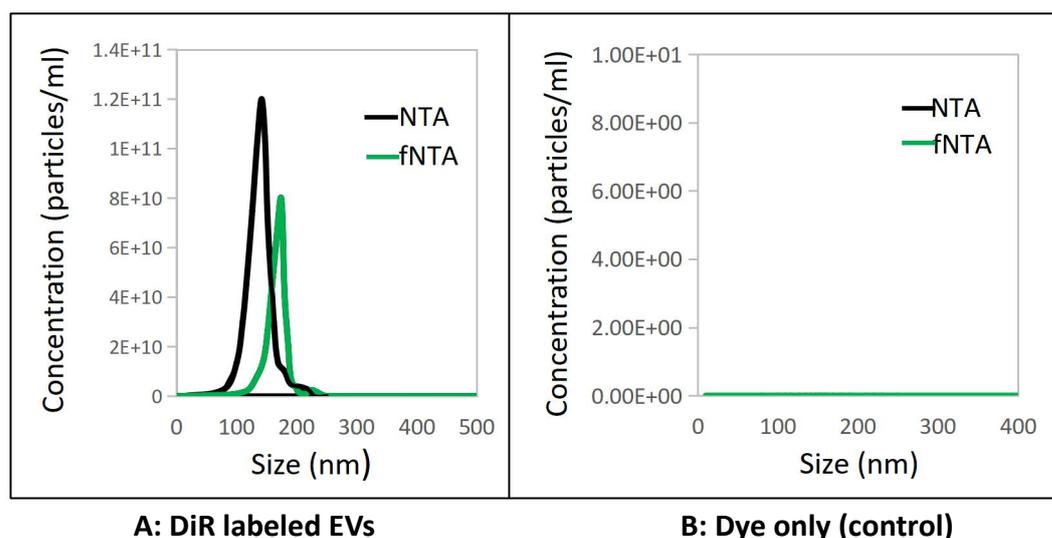


Figure 1. DiR labeled EVs and free dye control were analysed by NTA and fNTA.

The exosomes from LnCAP cell culture supernatant were concentrated by Exosome Concentration Kit (Cat# EXOCCon5-10), then were labeled and purified with DiR Membrane EVs Labeling & Purification Kit. A control involving dye but no exosomes was performed in parallel to confirm dye retention by the column. The results of NTA and fNTA indicated that a fairly considerable portion of exosomes were successfully labeled, the free dyes did not aggregate and the unbound dye can be removed successfully through the column.

Related Products

| Exosome labeling & Purification | |
|--|---------------------------|
| DiO-Membrane EVs Labeling & Purification Kit (green) | EXOPDiO10-1/EXOPDiO20-1 |
| Dii-Membrane EVs Labeling & Purification Kit (red) | EXOPDii10-1/EXOPDii20-1 |
| EVs labeling (DiO) Kit (green) | EXODiO10-1 |
| EVs labeling (Dii) Kit (red) | EXODii10-1 |
| EVs labeling (DiR) Kit (near-infrared red) | EXODiR10-1 |
| Exosome Spin Columns (MW 4000) | RGESC12-1/ RGESC24-1 |
| Exosome Isolation & Purification | |
| Exosome Extraction & Purification Kits (for blood serum/plasma) | EXORG10SP-1/ EXORG30SP-1/ |
| Exosome Extraction & Purification Kits (for cell culture media/urine) | EXORG10CU-1/ EXORG30CU-1/ |
| Exosome Concentration Kits (for cell culture media/urine) | EXOCCon5-10/ EXOUCon5-10 |
| Total Exosome Capture & Isolation Kits (for cell culture media/urine) | EXOMBoCU-10/EXOMBoCU-20 |
| Exosome Nucleic Acid Extraction | |
| Exosome Extraction & DNA Isolation Kits (for blood serum/plasma) | EXODNA30A-1/ EXODNA50A-1 |
| Exosome Extraction & DNA Isolation Kits (for cell culture media/urine) | EXODNA10B-1/EXODNA24B-1 |
| Exosome Extraction & RNA Isolation Kits (for blood serum/plasma) | EXORNA30A-1/EXORNA50A-1 |
| Exosome Extraction & RNA Isolation Kits (for cell culture media/urine) | EXORNA10B-1/EXORNA24B-1 |
| Exo-Antibody | |
| Purified Anti-human Alix Antibody | RGAB100-50/RGAB100-100 |
| Purified Anti-human CD9 Antibody | RGAB101-50/RGAB101-100 |
| Anti-human CD9 Ab Biotin Conjugated | RGAB102-50/RGAB102-100 |
| Purified Anti-human CD63 Antibody | RGAB103-50/RGAB103-100 |
| Anti-human CD63 Ab Biotin Conjugated | RGAB104-50/RGAB104-100 |
| Purified Anti-human CD81 Antibody | RGAB105-50/RGAB105-100 |
| Anti-human CD81 Ab Biotin Conjugated | RGAB106-50/RGAB106-100 |

Troubleshooting

Q1: The fluorescence signal is low than expected?

A1: There may be too low amount exosome for labeling, it is recommended to take a larger sample to extract exosome. In addition, fluorescent dye will be quenched, please keep dye from light when operating. Ensure that the columns do not dry out during the procedure. Spinning the column for too long or at too high speed may cause the column to work inefficiently and loss of the labeled exosomes.

Q2: How to store the labeled exosomes when not carry out the downstream

experiments immediately?

A2: The labeled exosomes can be stored at 2-8°C for 1-2 days aware of light, If kept for a long time, it is recommended to store at -80°C avoid repeated freezing and thawing.

Q3: Can I increase the elution volume?

A3: This is not recommended as it will result in co-elution of excess unbound dye with exosomes.

Q4: How I can know the counts of labeled exosomes?

A4: The labeled exosomes usually can be detected and counted by fluorescent Nanoparticle Tracking Analysis (fNTA) or flow cytometry. DiR labeling did not change the size of exosomes actually, but the calculation of particle size is related to many factors, which leads to the measurement value of size of the same labeled exosome sample under the fluorescence mode (fNTA) is larger than that under the conventional mode (NTA).

Technical Support

For more information about our products and to download manuals, please visit our web site: <http://www.rengenbio.com>

For additional information or technical assistance, please call or email us.

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