Exosome Isolation Kit (from cell culture media)

Catalog No.: EZB-exo2

Description

The **EZ**Bioscience® Exosome Isolation Kit (from cell culture media) provides a simple, reliable, and rapid method for isolating high-quality intact total exosome from cell culture media samples, without the need for ultracentrifugation.

Exosomes are small vesicles (30 ~ 120 nm) containing RNA and protein that are secreted by various types of cells in culture, and found in abundance in body fluids, such as blood, saliva, urine, and breast milk. Exosomes were reported to function as intercellular messengers, delivering their cargo of effector or signaling macromolecules between specific cells; however, their formation, the makeup of the cargo and biological pathways in which they are involved remain unclear.

The biological and medical study of exosome function and trafficking requires the isolation of intact exosomes, but the current methods used are tedious and difficult. The EZBioscience® Exosome Isolation Kit (from cell culture media) provides a simple and reliable method of concentrating intact exosomes from cell culture media samples. By tying up water molecules, the Exosome Precipitation Reagent forces less-soluble components (i.e. exosomes) out of solution, allowing them to be collected with a brief and relatively low-speed centrifugation. After that, the exosomes could be used for RNA purification, protein extraction, etc.

Components

| Component | EZB-exo2 |
|-------------------------------------|----------|
| Exosome Precipitation Reagent (EPR) | 50 ml |

The Exosome Precipitation Reagent contains enough reagent for processing up to 200 ml of cell culture media.

Storage

Store the Exosome Precipitation Reagent at 2°C to 8°C.

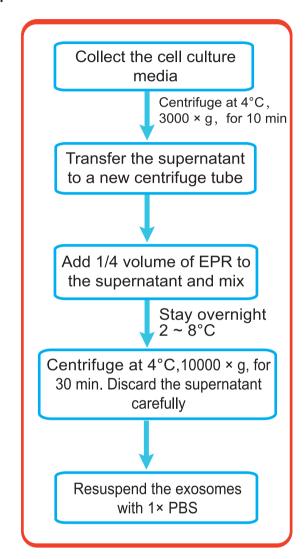
Protocol

General Guidelines

- a) To ensure that isolated exosomes originate from your cells of interest, culture the cells with exosome depleted fetal bovine serum (FBS) or serum-free culture medium, because normal FBS contains extremely high levels of exosomes that will contaminate the cell derived exosomes. If you cannot obtain exosome depleted FBS, certain cell lines can be grown for up to 12 hours in media without FBS.
- b) The Exosome Precipitation Reagent (EPR) is not recommended for isolation of exosomes from other kinds

of samples. If you are isolating intact exosomes from plasma, use the Total Exosome Isolation (from plasma) reagent.

Experimental Procedure Overview:



Prepare Sample

- When the cell confluence reach to about 60%, change the media to DMEM, or to media with exosome-depleted- FBS.
- Culture the cells for 12 ~ 24 hours, until the cell confluence reach to about 90%. Harvest cell culture media.
- Centrifuge the cell media at 3000 × g for 10 minutes to remove cells and debris.
- Transfer the supernatant containing the cell-free culture media to a new tube without disturbing the pellet.

Isolate Exosomes

Transfer the required volume of cell-free culture media to a new tube and add 1/4 volume of the **Exosome Precipitation Reagent (EPR).**

| Culture Media | Reagent |
|---------------|---------|
| 5 ml | 1.25 ml |
| 20 ml | 5 ml |

- Mix the culture media/reagent mixture well either by vortexing or inversion until the solution is homogenous. Incubate the sample at 2 ~ 8°C overnight.
- After incubation, centrifuge the sample at 10000 × g 3. for 30 minutes at 4°C.
- Carefully aspirate the supernatant by pipetting and discard the supernatant. Exosomes are contained in the pellet at the bottom of the tube (not visible in most cases).

Resuspend Exosomes

Add 1× PBS or similar buffer to the pellet and vortex or pipette up and down to resuspend the exosomes.

| Starting Cell Culture Media Volume | Resuspension Volume |
|------------------------------------|------------------------|
| 10 ml | 30 µl |
| 20 ml | 60 µl |

The purified exosomes are ready for downstream analysis or further purification through affinity methods.

Note: The purified exosome sample may be stored at 2 ~ 8°C for up to 7 days or can be stored at -80°C for long term storage. To minimize the risk of RNA degradation, we recommend proceeding directly with further downstream sample processing.