

# VitroView<sup>TM</sup> RTU Golgi Staining Solution SKU#: VB-1005

**Description:** The Golgi apparatus is a folded flattened membrane-enclosed organelle that consists of multiple compartments within the cytoplasm of most eukaryotic cells, whose function is involved in protein secretion and intracellular transport of lipids and carbohydrates released from the endoplasmic reticulum (ER) to export outside of the cell. It also resides at the intersection of the secretory, lysosomal, and endocytic pathways. Here Vitrovivo Biotech offers a cell-permeant probe that can be used to distinguish the Golgi morphology in both live and fixed cells. Additionally, this product has application in lipid metabolism, trafficking studies.

NBD C6-ceramide is a sensitive fluorescent dye (Ex:466nm/Em:536nm) working in aprotic solvents and other nonpolar environments. It can be used as a selective stain for the Golgi apparatus in live and fixed cells. It is also used to study sphingolipid transport and metabolism mechanisms.

#### **Contents:**

VB-1005 RTU Golgi Staining Solution-----30 ml

#### **Storage**

Store at 2-8 °C and protect from light.

### Material Needed But NOT Supplied with the Kit:

- 1. Staining jar
- 2. Glass slides
- 3. PBS (Wash Buffer)
- 4. Mount media

#### **General Protocol:**

## Adherent cells for fluorescence microscopy

- 1. Grow cultured cells on sterile glass cover slips or slides overnight at 37 °C.
- 2. Follow appropriate protocol to fix cultured cells.
- 3. Completely wash the cells with PBS as needed.
- 4. Add adequate RTU Golgi Staining Solution to cover the whole sample.
- 5. Incubate under dark at room temperature for 15-30 minutes.
- 6. Rinse the sample several times with PBS and remove excess dye.
- 7. Add antifade aqueous mounting medium and mount.
- 8. Use appropriate filters and detect under fluorescence microscope according to standard protocol.

Suspension cells for fluorescence microscopy

Page 1/2

- 1. The cells are harvested into a 15 mL polypropylene centrifuge tube and spin down for 8 min at 600 RPM.
- 2. The supernatant is discarded and the cells are resuspended in 0.5 ml of culture medium
- 3. 1-2 drops of the cell suspension were placed on a slide in the central area and moved around to form a thin and even film with a glass spreader.
- 4. (Option) You also can use cytocentrifuge to prepare cell slides.
- 5. Air dry and follow appropriate protocol to fix cultured cells.
- 6. Drop adequate RTU Golgi Staining Solution to cover the whole sample on slide.
- 7. Incubate under dark at room temperature for 15-30 minutes.
- 8. Cover with coverslip and view under fluorescence microscope according to standard protocol.

**Note:** This product is intended for research purposes only. This product is <u>not</u> intended to be used for therapeutic or diagnostic purposes in humans or animals.

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