



## DATA SHEET

### Dil-LABELED LOW DENSITY LIPOPROTEIN, HUMAN

**Catalog No:** YB-0011

**Lot No:** lot specific

**Quantity:** 500ug(micrograms)Protein/vial

**Concentration:** lot specific (mg/ml Protein)

#### **Product Introduction:**

Purified Yiyuan Biotechnologies LDL is labeled with the fluorescent probe, Dil, and reisolated by ultracentrifugation (1.019-1.063). The resultant product is exhaustively dialyzed against phosphate buffered saline, (pH 7.4), sterilized by membrane filtration and then aseptically packaged in a solution containing phosphate-buffered saline at pH 7.4 and 0.2 mM EDTA. Each lot is evaluated on a murine macrophage cell line for fluorescence uptake.

#### **Storage & Stability:**

Dil-LDL should be kept sterile at 2-8°C. **NEVER FREEZE**. The stability of this product is in the 6 week range after receipt. Clarify by centrifugation if needed. Before using, spin the vial in a microfuge for 10 minutes over 5000g.

#### **Typical Lipoprotein Labeling Protocol**

1. Aseptically dilute the Dil-LDL to 10-40µg/ml in your culture media.
2. Add to live cells and incubate for 2-4 hours at 37°C.
3. Remove media containing Dil-LDL from your culture.
4. Wash cells several times with probe-free media.
5. A. Fluorescence Microscopy:

Visualize using standard rhodamine excitation: emission filters (or suggested wavelengths excitation:emission at 554nm:571nm or near). If fixation is desired use 3% formaldehyde in PBS. (Never use methanol or acetone fixation - Dil is soluble in organic solvents). Note: A positive culture must be stained for comparison purposes.

B. Cell Sorting:

Label as in steps 1-5. Trypsinize or treat cultures with EDTA to produce a single cell suspension. Use labeled pure cultures of positive and negative cell types to set gates on the cell sorter.

Suggested Wavelengths for Cell Sorting: Excitation: 554nm  
Emission: 571nm

**Special Note:** All our LDL products have a natural tendency to aggregate. Aggregates of this product can interfere with its use. To clarify these aggregates out, simply spin in a microfuge for 2 minutes.

**FOR RESEARCH USE ONLY**

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