

# TBNK Panel

Multi-Color Flow Cytometry Panel

Catalog Number: FMC-P-001

### PRODUCT DESCRIPTION

T cells, B cells, and NK cells are the major lymphocyte subsets in peripheral blood. This panel contains 5 conjugated antibodies that can be used for the single step staining of T cells, B cells, and NK cells.

#### **INTENDED USE**

This multicolor flow cytometry panel allows for easy identification of each cell type.

### **MATERIALS PROVIDED & STORAGE**

Store the unopened kit at 2-8 °C. Do not use past expiration date.

MARKER	CLONE	FLUOROCHROME	SIZE	RECOMMENDED CONCENTRATION
CD3	UCHT1	Alexa Fluor® 405	100 tests	5 μL/10 <sup>6</sup> cells
CD4	11830	FITC	100 tests	10 μL/10 <sup>6</sup> cells
CD8	37006	APC	100 tests	10 μL/10 <sup>6</sup> cells
CD19	4G7-2E3	PE	100 tests	10 μL/10 <sup>6</sup> cells
CD56	2524C	Alexa Fluor® 700	100 μg	0.25-1 μg/10 <sup>6</sup> cells

Note: Recommended concentrations are given as reference point for antibody titration. Optimal concentrations should be determined by each laboratory for their experimental conditions.

### **PRECAUTIONS**

Wear protective gloves, clothing, eye, and face protection. Wash hands thoroughly after handling. Refer to the SDS on our website prior to use.

#### **OTHER SUPPLIES REQUIRED**

- PBS
- Flow Cytometry Staining Buffer (R&D Systems®, Catalog # FC001)
- Fc-block (blocking IgG)
- Isotype Control Antibodies (optional)
- 5 mL Flow cytometry tubes

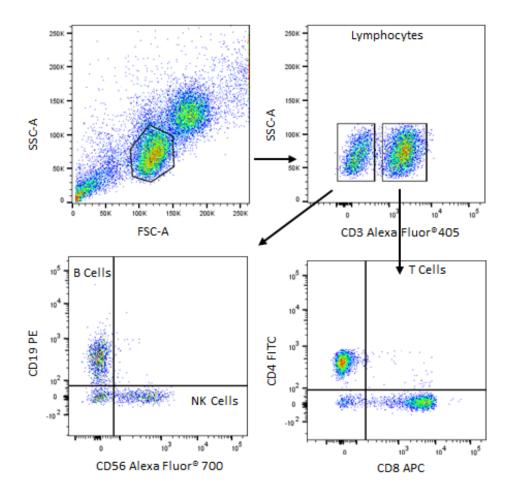
#### **PROTOCOL**

- 1. Wash human PBMCs (1 x 10<sup>6</sup> cells per sample) with 2 mL of <u>Staining Buffer (1X) (Catalog # FC001)</u> or other BSA-containing buffer, by spinning at 300 x g for 5 minutes, using 5 mL flow cytometry tubes. Decant/aspirate supernatant.
- 2. Fc-block cells with blocking IgG (1  $\mu$ g IgG/10 $^{\circ}$  cells) for 10 minutes at room temperature.
- 3. Add 5 µL (or previously titrated amount) of each primary conjugated antibody. Vortex tubes.
- 4. (optional) Separate tube, add 5 µL of each of the isotype control antibodies. Vortex tubes.
- 5. Incubate the mixtures for 30-45 minutes at room temperature in the dark.
- 6. Wash with 2 mL of Staining Buffer (1X), by spinning at 300 x g for 5 minutes. Decant/aspirate supernatant.
- 7. Resuspend the cells in 0.2 0.5 mL Staining Buffer (1X) and acquire on a Flow Cytometer.

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## **DATA EXAMPLES**



**Human TBNK Cell Panel Data**. Multicolor flow cytometry panel to identify human T Cells, B Cells, and NK cells. PBMCs were stained with anti-human CD3 Alexa Fluor® 405, CD4 FITC, CD8 APC, CD56 Alexa Fluor 700, and CD19 PE. All antibodies are validated for flow cytometry.

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