Cell-ID™ Intercalator-Ir 500 µM

Catalog #201192B (500 µL)

WARNING! CHEMICAL HAZARD. Before handling any chemicals, refer to the Safety Data Sheet (SDS) provided by the manufacturer, and observe all relevant precautions.

NOTICE: HIGH CONCENTRATION. Cell-ID Intercalator-Ir 500 µM is a highly concentrated metal intercalator solution and must be diluted in accordance with this protocol to avoid early failure of the detector.

Description

Cell-ID Intercalator-Ir is a cationic nucleic acid intercalator that contains natural abundance Iridium (^{191}Ir and ^{193}Ir) and is used for identifying nucleated cells in CyTOF® system analysis. When cells are stained with Intercalator-Ir, it will bind to cellular nucleic acid, and detection of both stable isotopes will enable identification of nucleated cells. It is a live cell membrane-impermeable dye and therefore requires cells to be fixed and/or permeabilized before staining.

Note: While dilutions of the 500 µM stock solution are suggested in the protocols below, the concentration can be titrated for individual cell types and experiments for optimal Cell-ID Intercalator staining. It is suggested not to exceed 1 µM intercalator concentration in the staining solution.

Storage

Upon arrival, aliquot and store at -20 °C. Aliquots stored at 4 °C are stable for up to three months.

Staining Protocol

1. After cell staining is complete, prepare 1 mL of cell intercalation solution for each sample by diluting Cell-ID Intercalator-Ir 1:4000 into Maxpar® fix and perm buffer (Fluidigm Cat. 201067) and mix by vortexing.
2. Add 1 mL of the intercalation solution prepared in step 1 to each tube and gently vortex. Incubate for one hour at room temperature or leave overnight at 4 °C.

   Note: Cells can be left at 4 oC in the intercalation solution up to 48 hours.

3. Wash cells by adding 2 mL of Maxpar® cell staining buffer (Fluidigm Cat. 201068), centrifuge and discard supernatant by aspiration.
4. Repeat for a total of two washes with Maxpar cell staining buffer.
5. Wash cells with 2 mL of Maxpar® water (Fluidigm Cat. 201069), centrifuge and discard supernatant by aspiration.
6. Leave cells pelleted until ready to run on the CyTOF system. Immediately prior to data acquisition, adjust cell concentration to 2.5–5 x 10^5/mL with Maxpar water and filter cells into cell strainer cap tubes.
7. Acquire data on the CyTOF system.

For technical support visit fluidigm.com/support
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