**HLA-B27 SOP 1: BD FACSCanto™ II Cytometer Startup Procedure**

**Purpose**
To prepare the BD FACSCanto™ II flow cytometer for acquisition of stained samples.

**Scope**
This procedure applies to the clinical laboratory environment with the BD FACSCanto II flow cytometer for the purpose of detecting the HLA-B27 antigen using whole blood specimens. We recommend that all personnel who operate the instrument be sufficiently trained to fully perform and implement this guideline.

**Equipment Required**
BD FACSCanto II flow cytometer and workstation

**Materials Required**
- Biohazard safety manual
- Biohazard sharps waste container
- Personal protective equipment (PPE)
  - Protective gloves
  - Protective eyewear
  - Closed-toe shoes
  - Lab coat
- BD FACSFLOW™ sheath fluid (Catalog No. 342003)
- BD FACS™ Clean solution (Catalog No. 340345)
- BD FACS™ Shutdown solution (Catalog No. 334224)
- 12 x 75-mm Falcon® tubes

**Procedure**

**Starting the system**
1. Turn on the computer main power.
2. Log in to Windows.
3. Turn on the cytometer main power.
4. Double-click the BD FACSCanto™ clinical software icon on the desktop to start the software.
5. Log in to the software with the appropriate user and password information.
6. Wait for the cytometer to connect. The icon in the bottom right-hand corner of the software window will turn green when the cytometer is connected.
7. Allow the cytometer to warm up for 30 minutes (20-minute minimum) before running any beads. The elapsed time will be displayed in the bottom right-hand corner of the window.

**During the warm-up time for the cytometer:**
1. Check the BD FACSFlow level in the Status window. If necessary, replace the cubitainer with a full one.
2. Empty the waste container, if necessary.
3. Check the BD FACSFlow filter on the fluidics cart for air.
   a. If bubbles are present, loosen the cap on the top of filter and allow the air to escape.
   b. Once fluid starts dripping from the filter, tighten the cap on the filter.
4. Select Cytometer > Fluidics Startup.
5. Click OK in the dialog to confirm.
   Fluidics startup will take about seven minutes to complete.
6. After fluidics startup has completed, select Cytometer > Cleaning Modes > Bubble Filter Purge. When the process is complete, click OK.
7. Select Cytometer > Cleaning Modes > De-gas Flow Cell. When the process is complete, click OK.
8. Repeat Step 7 to de-gas the flow cell a second time.

**Performing instrument QC with BD FACS 7-color setup beads**
1. Open a foil pouch containing one tube of BD FACS 7-color setup beads.
2. Add the BD FACS 7-color setup bead diluent to the line on the tube.
3. Vortex the tube for two seconds to completely mix the beads and diluent.
5. Select the current bead lot ID, as listed on the sticker included in the box.
6. To enter a new bead lot using a 2D barcode reader:
   a. Click the Scan Barcodes button.
   b. Scan the 2D barcode on the lot ID sticker and the information will be automatically populated into the appropriate fields.
   c. Check all affected software fields for accuracy against the setup beads label.
7. To enter a new bead lot manually:
   a. Click the New Lot ID button.
   b. Enter the lot ID and expiration date and click OK.
   c. In the Targets tab, enter the appropriate target values printed on the sticker included in the box.
**HLA-B27 SOP 1:**
**BD FACSCanto™ II Cytometer Startup Procedure**

- d. In the **Spectral Overlap Factors** tab, enter the appropriate values printed on the sticker included in the box.
- e. Click **Finish**.

8. Click **Next**. If you changed lot ID values, click **Yes** when prompted.

9. Select **Run setup in Manual mode**, and click **Next**.

10. When prompted, load the tube of beads onto the SIT.

11. Click **OK** and wait for setup to finish.

12. Unload the beads when prompted. When the run is complete, a dialog is displayed with the message **Setup Completed Successfully**.

13. Click **View Setup Report** to print the report and review the values obtained.

Performing application setup for the HLA-B27 assay

- 1. Add 1 mL of 0.22-micron filtered sheath to a 12 x 75-mm Falcon polystyrene tube.
- 2. Mix the HLA-B27 bead vial by inversion and dispense two drops into the filtered sheath.
- 3. From the **Cytometer** menu, select **Setup > HLA B27 setup**.
- 4. Enter the bead lot number and suffix and reagent lot number and suffix information in the window, using the following pictures for guidance.

5. Install the well mixed HLA-B27 bead tube on the cytometer and click **Start**.

Once the setup is complete, a report will be generated.
6. Click **View Report** and confirm that the Overall Result is **PASS**.

7. To close the report and return to the wizard, click the **Close Preview** button.

8. Click **Save**.

9. Unload the tube when prompted.

10. Click **Finish** to close the Cytometer Setup Wizard.
HLA-B27 SOP 1:
BD FACSCanto™ II Cytometer Startup Procedure

References

BD FACSCanto™ II Instructions for Use, document 23-12882-01.


BD HLA-B27 Application Guide for BD FACSCanto Flow Cytometers, document 343366 Rev. A.