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## Fcap Array Software V3.0 Download Firefox [BEST]

the bd cba mouse il-5 flex set is a multi-analyte flow cytometry detection set used to measure levels of interleukin-5 (il-5) in the blood of mice. this assay is used to evaluate allergic reactions to a variety of antigens, including dust mite, ragweed, and aspergillus. to run this assay, the following general steps must be performed: (a) prepare the sample. if the sample is a plasma or serum, the sample must be centrifuged for 5 min at 10,000 x g to remove cells and large particles. the supernatant is then removed and stored at 4° c. if the sample is a whole blood sample, the sample is prepared in a similar manner except that the sample must be diluted 1:2 or 1:4 with facs flow and the sample is then incubated with bd fix/perm solution for 20 min at 4° c. (b) prepare the reagents. dilute the bd cba mouse il-5 flex set (1:10 dilution of the reagent stock) with perm/wash buffer to the appropriate concentration (2,000 - 10,000 pg/ml). (c) prepare a standard curve and a negative control sample. prepare a standard curve by diluting standards from the bd cba mouse il-5 flex set (100 pg/ml, 1,000 pg/ml, 5,000 pg/ml, 10,000 pg/ml, and 20,000 pg/ml) in perm/wash buffer to the appropriate concentration. prepare a negative control by diluting an aliquot of the perm/wash buffer 1:10 with perm/wash buffer. (d) load the test sample and standards. load the test sample into a flow cytometer and the bd cba mouse il-5 flex set into the bd facsarray bioanalyzer. add the test sample and standards to the assay plate and mix by shaking. after the plate is mixed, place the plate on the instrument. (e) analyze the data.

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when using bd cba human il-8 flex set (in conjunction with a bd cba human soluble protein master buffer kit (cat. no. 558264, 100 tests, or 558265, 500 tests), a flow cytometer, and fcap array software, analyze the results of the standard curve. set up the software to receive the normal distribution of all the data. in a perfect situation, you should have 3 as the first cut point; 8,500 pg/ml for most of the samples (and as low as 1,800 pg/ml for the most highly expressing samples); 15,300 pg/ml for the least expressing samples; and 21,300 pg/ml for the top two expression samples. normally, a large number of samples are tested on a day. because of that, the total time needed to test 100,000 beads per sample is about 9 minutes (average). there is a slight reduction of the time needed to test a sample when using an influx or fluorospot, compared to using a becton and dickinson facscalibur. the software deals with up to 12 samples per run. each sample can be tested for as many as 500,000 beads. the bd cba human il-8 flex set must be used in conjunction with a bd cba human soluble protein master buffer kit (cat. no. 558264, 100 tests, or 558265, 500 tests), a flow cytometer, and fcap array software. detailed instructions on the use of this product can be found in the manual for the bd cba human soluble protein master buffer kit. when following the directions in the master buffer kit, the top standard point for the bd cba human il-8 flex set will be 2,500 pg/ml. an example standard curve is shown in figure 3. when using bd cba human granzyme b flex set (in conjunction with a bd cba human soluble protein master buffer kit (cat. no. 558264, 100 tests, or 558265, 500 tests), a flow cytometer, and fcap array software, analyze the results of the standard curve. set up the software to receive the normal distribution of all the data. in a perfect situation, you should have 3 as the first cut point; 8,500 pg/ml for most of the samples (and as low as 1,800 pg/ml for the most highly expressing samples); 15,300 pg/ml for the least expressing samples; and 21,300 pg/ml for the top two expression samples. 5ec8ef588b

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