



BD Cytometric Bead Array (CBA) Mouse/Rat Soluble Protein Master Buffer Kit Instruction Manual

Catalog No. 558266 (100 Tests)
Catalog No. 558267 (500 Tests)



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Regulatory information

BD flow cytometers are class I (1) laser products.

For Research Use Only. Not for use in diagnostic or therapeutic procedures.

History

Revision	Date	Change made
23-12722-00 Rev. 01	2/2011	New document

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About this kit

This section covers the following topics:

- [Purpose of the kit \(page 6\)](#)
- [Limitations \(page 8\)](#)
- [Kit contents \(page 10\)](#)
- [Storage and handling \(page 11\)](#)

Purpose of the kit

Use of the kit The BD™ CBA Mouse or Rat Soluble Protein Flex Set and BD™ CBA Mouse/Rat Soluble Protein Master Buffer Kit employ particles with discrete fluorescence intensities to detect soluble analytes at very low concentrations. The working assay range for most analytes in this system is 10 to 2,500 pg/mL for mouse assays and 40 to 10,000 pg/mL for rat assays.

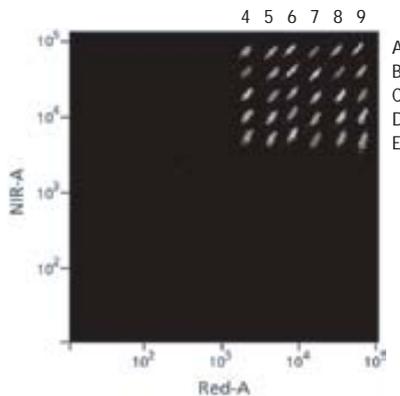
The BD CBA Mouse/Rat Soluble Protein Master Buffer Kit contains all of the supporting reagents necessary to perform an assay using a BD CBA Mouse or Rat Soluble Protein Flex Set. The buffers and instrument setup reagents provided in this kit have been optimized for analysis of analytes in tissue culture supernatants, plasma, and serum samples. The BD CBA Mouse/Rat Soluble Protein Master Buffer Kit provides sufficient reagents for the quantitative analysis of 100 samples and 10 instrument setup procedures (Catalog No. 558266) or 500 samples and 10 instrument setup procedures (Catalog No. 558267).

Principle of CBA assays BD CBA assays provide a method of capturing a soluble analyte or set of analytes with beads of known size and fluorescence, making it possible to detect analytes using flow cytometry.

Each capture bead in the BD CBA Mouse or Rat Soluble Protein Flex Set System has a distinct fluorescence and is coated with a capture antibody specific for a soluble protein. The detection reagent used in the BD CBA Mouse or Rat Soluble Protein Flex Set System provides a fluorescent signal in proportion to the amount of bound analyte.

When the capture beads and detection reagents are incubated with standards or unknown samples containing recognized analytes, sandwich complexes (capture bead + analyte + detection reagent) are formed. These complexes can be measured using flow cytometry to identify particles with fluorescence characteristics of both the bead and the detector.

A BD CBA Mouse or Rat Soluble Protein Flex Set Capture Bead is a single bead population with distinct fluorescence intensity and is coated with a capture antibody specific for a soluble protein. The bead population is resolved in two fluorescence channels of a flow cytometer. For specific instruments and the channels used to resolve the beads, see the flow cytometers listed in [Materials required but not provided \(page 15\)](#).



Each bead population is given an alphanumeric position designation indicating its position relative to other beads in the BD CBA Flex Set System. Beads with different positions can be combined in assays to create a multiplex assay. The intensity of PE fluorescence of each sandwich complex reveals the concentration of that particular analyte. After acquiring samples on a flow cytometer, use

FCAP Array™ software to generate results in graphical and tabular format.

Advantages over ELISA

The broad dynamic range of fluorescence detection via flow cytometry and the efficient capturing of multiple analytes via suspended particles enable the BD CBA Flex Set System to obtain the concentration of an unknown in substantially less time and using fewer sample dilutions compared to conventional ELISA methodology.

- The BD CBA Mouse or Rat Soluble Protein Flex Set assays allow for multiplexed analysis of multiple proteins from a single sample.
 - A single set of diluted standards is used to generate a standard curve for each analyte.
 - The BD CBA Mouse or Rat Soluble Protein Flex Set assays are more sensitive than most comparable ELISAs.
-

Limitations

Limitations

The BD CBA Mouse or Rat Soluble Protein Flex Set System is not recommended for use on stream-in-air instruments in which signal intensities might be reduced, adversely affecting assay sensitivity. Stream-in-air instruments include the BD FACStar™ Plus, BD FACSVantage™, and BD Influx™ flow cytometers (BD Biosciences).

Quantitative results or protein levels for the same sample or recombinant protein run in ELISA and BD CBA assays may differ. A spike recovery assay can be performed using an ELISA standard followed by BD CBA analysis to assess possible differences in quantitation.

When several BD CBA Mouse or Rat Soluble Protein Flex Set assays are multiplexed, it is possible that the background (MFI of the 0 pg/mL standard point) might increase and the overall assay signals of other standard points may be reduced. This can result in lower dynamic range or loss in sensitivity in some assays. This effect might be greater as more assays are added to the multiplex.

For assays that will be acquired on a BD FACSCalibur™ flow cytometer, we recommend that additional dilutions of the standard be prepared (ie, 1:512 and 1:1024), since it is possible that in multiplex experiments containing a large number of assays the Top Standard, 1:2, and 1:4 standard dilutions cannot be analyzed by FCAP Array™ software. In those cases, the Top Standard, 1:2, and 1:4 standard dilutions can be run in the experiment but might need to be excluded from the final analysis in FCAP Array software.

The BD CBA Mouse/Rat Soluble Protein Master Buffer Kit should not be used with any non-Mouse or Rat Soluble Protein Flex Sets. BD CBA Mouse and Rat Soluble Protein Flex Set assays cannot be performed in the same experiment.

Kit contents

Contents

The kit contains the following components sufficient for 100 tests (Catalog No. 558266) and 500 tests (Catalog No. 558267).

Reagent	Quantity (558266)	Quantity (558267)
Assay Diluent	1 bottle, 30 mL	1 bottle, 150 mL
Capture Bead Diluent	1 bottle, 5 mL	1 bottle, 30 mL
Detection Reagent Diluent	1 bottle, 5 mL	1 bottle, 30 mL
Wash Buffer	1 bottle, 130 mL	1 bottle, 650 mL
Instrument Setup Bead A1	1 vial, 0.25 mL	1 vial, 0.25 mL
Instrument Setup Bead A9	1 vial, 0.25 mL	1 vial, 0.25 mL
Instrument Setup Bead F1	1 vial, 1.0 mL	1 vial, 1.0 mL
Instrument Setup Bead F9	1 vial, 0.25 mL	1 vial, 0.25 mL
PE Instrument Setup Bead F1	1 vial, 0.25 mL	1 vial, 0.25 mL
PE Positive Control Detector	1 vial, 0.5 mL	1 vial, 0.5 mL

Note: Source of all serum proteins is from USDA-inspected abattoirs located in the United States.

Storage and handling

Storage Store all kit components at 2 to 8°C. Do not freeze.

Warning All components of this kit contain sodium azide. Sodium azide yields highly toxic hydrazoic acid under acidic conditions. Dilute azide compounds in running water before discarding to avoid accumulation of potentially explosive deposits in plumbing.

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Before you begin

This section covers the following topics:

- [Workflow overview \(page 14\)](#)
- [Required materials \(page 15\)](#)

Workflow overview

Workflow The overall workflow consists of the following steps.

Step	Description
1	Preparing Mouse/Rat Soluble Protein Flex Set Standards (page 18)
2	Mixing Mouse/Rat Soluble Protein Flex Set Capture Beads (page 21)
3	Diluting test samples (page 23)
4	Preparing Mouse/Rat Soluble Protein Flex Set Detection Reagents (page 24) Note: Can be prepared during the first incubation in step 6 below.
5	Performing instrument setup with Instrument Setup Beads, if necessary (instructions can be found at bdbiosciences.com/cbasetup) Note: Can be performed during one of the incubations in step 6 .
6	Performing the Mouse or Rat Soluble Protein Flex Set Assay (page 28)
7	Acquiring samples (instructions can be found at bdbiosciences.com/cbasetup)
8	Data analysis (page 33)

Incubation times To help you plan your work, the incubation times are listed in the following table.

Procedure	Incubation time
Preparing standards	15 minutes
Assay Procedure	
<ul style="list-style-type: none"> • First incubation–Capture Beads 	1 hour
<ul style="list-style-type: none"> • Second incubation–PE Detection Reagent 	1 hour (mouse assays) 2 hours (rat assays)

Required materials

Materials required but not provided

In addition to the reagents provided in the BD CBA Mouse/Rat Soluble Protein Master Buffer Kit and the BD CBA Mouse or Rat Soluble Protein Flex Set, the following items are also required.

- A dual-laser flow cytometer equipped with a 488-nm or 532-nm and a 633-nm or 635-nm laser capable of distinguishing 576-nm, 660-nm, and >680-nm fluorescence. The following table lists examples of compatible instrument platforms.

Flow cytometer	Reporter channel	Bead channels
BD FACSAria™	Yellow	Red and NIR
BD FACSCanto™ platform BD™ LSR platform BD FACSAria™ platform	PE	APC and APC-Cy™7
BD FACSCalibur™	FL2	FL4 and FL3
Note: Visit bdbiosciences.com/cbasetup for setup protocols.		

- BD Falcon™ 12 × 75-mm sample acquisition tubes for a flow cytometer (Catalog No. 352008)
 - 15-mL conical polypropylene tubes (BD Falcon, Catalog No. 352097), or equivalent
 - FCAP Array software (Catalog No. 641488 [PC] or 645447 [Mac®])
 - Microcentrifuge
-

**Materials
required for
plate loader-
equipped flow
cytometers**

- Millipore MultiScreen_{HTS}-BV 1.2-µm Clear non-sterile filter plates [Catalog No. MSBVN1210 (10 pack) or MSBVN1250 (50 pack)]
 - Millipore MultiScreen_{HTS} Vacuum Manifold (Catalog No. MSVMHTS00)
 - MTS 2/4 Digital Stirrer, IKA Works, VWR (Catalog No. 82006-096)
 - Vacuum source
 - Vacuum gauge and regulator (if not using the recommended manifold)
-

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Assay preparation

This section covers the following topics:

- [Preparing Mouse/Rat Soluble Protein Flex Set Standards \(page 18\)](#)
- [Mixing Mouse/Rat Soluble Protein Flex Set Capture Beads \(page 21\)](#)
- [Diluting test samples \(page 23\)](#)
- [Preparing Mouse/Rat Soluble Protein Flex Set Detection Reagents \(page 24\)](#)

Preparing Mouse/Rat Soluble Protein Flex Set Standards

Purpose of this procedure

The BD CBA Mouse or Rat Soluble Protein Flex Set Standards are lyophilized and must be reconstituted and serially diluted immediately before mixing with the Capture Beads and the Detection Reagent.

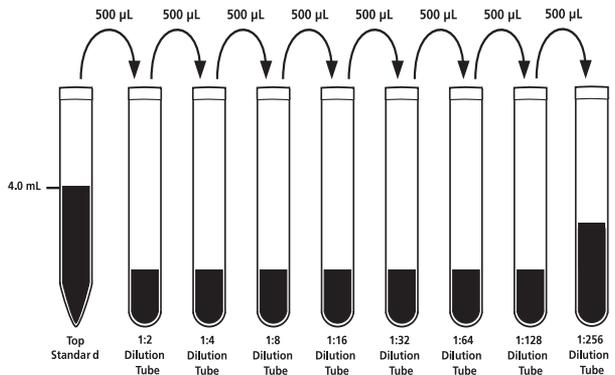
Note: You must prepare fresh standards to run with each single bead or multiplex experiment. Do not store or reuse reconstituted or diluted standards.

Procedure

To reconstitute and serially dilute the standards:

1. Open one vial of lyophilized standard from each BD CBA Mouse or Rat Soluble Protein Flex Set that will be tested.
2. Pool all lyophilized standard spheres into one 15-mL polypropylene tube. Label the tube “Top Standard.”
3. Reconstitute the standards with 4 mL of Assay Diluent.
 - a. Allow the reconstituted standard to equilibrate for at least 15 minutes at room temperature.
 - b. Gently mix the reconstituted standard by pipet only. Do not vortex or mix vigorously.
4. Label eight 12 × 75-mm tubes and arrange them in the following order: 1:2, 1:4, 1:8, 1:16, 1:32, 1:64, 1:128, and 1:256.
5. Pipette 500 µL of Assay Diluent into each of the 12 × 75-mm tubes.
6. Perform a serial dilution.

- a. Transfer 500 μL from the Top Standard to the 1:2 dilution tube and mix thoroughly by pipet only. Do not vortex.
- b. Continue making serial dilutions by transferring 500 μL from the 1:2 tube to the 1:4 tube and so on to the 1:256 tube.



7. Prepare one 12 x 75-mm tube containing Assay Diluent to serve as the 0-pg/mL negative control.

Note: We recommend that the first 10 wells or tubes in the experiment be the standards. Standards should be run in order from least concentrated (0 pg/mL) to most concentrated (Top Standard) to facilitate analysis in FCAP Array software.

Concentration of standards The approximate concentration (pg/mL) of each BD CBA Mouse or Rat Soluble Protein Flex Set Standard in each dilution tube is shown in the following table.

Note: See the technical data sheet for each individual assay to verify the concentration of the Top Standard.

	Dilution tube								
Mouse Soluble Protein Flex Set	Top Standard	1:2	1:4	1:8	1:16	1:32	1:64	1:128	1:256
Protein (pg/mL)	2,500	1,250	625	312.5	156	80	40	20	10

	Dilution tube								
Rat Soluble Protein Flex Set	Top Standard	1:2	1:4	1:8	1:16	1:32	1:64	1:128	1:256
Protein (pg/mL)	10,000	5,000	2,500	1,250	625	312.5	156	80	40

Next step Proceed to [Mixing Mouse/Rat Soluble Protein Flex Set Capture Beads \(page 21\)](#).

Mixing Mouse/Rat Soluble Protein Flex Set Capture Beads

Purpose of this procedure The Capture Beads provided in each BD CBA Mouse or Rat Soluble Protein Flex Set are at a 50X concentration and must be diluted to their optimal concentration before use.

Mixing the beads To mix the Capture Beads:

1. Determine the number of BD CBA Mouse or Rat Soluble Protein Flex Sets to be used in the experiment (size of the multiplex).
2. Determine the number of tests in the experiment.

Note: Extra tests of Capture Beads should be mixed to ensure that the necessary number of tests will be recovered from the mixed Capture Beads tube. Add an additional 2 to 3 assay tubes to the number determined.

3. Vortex each Capture Bead stock vial for at least 15 seconds to resuspend the beads thoroughly.
4. Determine the total volume of diluted beads needed for the experiment. Each tube/well requires 50 μL of the diluted beads. Calculate the total volume of diluted beads by multiplying the number of tests (determined in [step 2](#)) by 50 μL .

Example: 35 tests \times 50 μL = 1,750 μL total bead volume

5. Determine the volume needed for each capture bead. Beads are supplied so that 1.0 μL = 1 test. Therefore, the required volume (μL) of beads is equal to the number of tests.

Example: 35 tests requires 35 μL of each Capture Bead included in the assay.

- Determine the volume of Capture Bead Diluent needed to dilute the beads. Calculate the Diluent volume by subtracting the volume for each bead tested from the total volume of diluted beads needed to perform the assay. See [Capture Bead and Detection Reagent Diluent](#) (page 38).

Example: 1,750 μL total volume of beads – 35 μL for each bead = volume of Capture Bead Diluent

- if testing one analyte: $1,750 \mu\text{L} - (35 \mu\text{L} \times 1) = 1,715 \mu\text{L}$ diluent
 - if testing five analytes: $1,750 \mu\text{L} - (35 \mu\text{L} \times 5) = 1,575 \mu\text{L}$ diluent
- Pipette the Capture Beads and Capture Bead Diluent into a tube labeled “Mixed Capture Beads.”

Next step

The Capture Beads are now ready to be transferred to the assay tubes. Discard excess prepared Capture Beads. Do not store after mixing.

If you need to dilute samples having high-protein concentrations (for example, serum or plasma samples), proceed to [Diluting test samples](#) (page 23). Otherwise, proceed to [Preparing Mouse/Rat Soluble Protein Flex Set Detection Reagents](#) (page 24).

If sample dilution is not required, you can save time by proceeding directly to [Performing the Mouse or Rat Soluble Protein Flex Set Assay](#) (page 28). Please note that you will need to prepare the Detection Reagents during the first assay incubation step. You will also need to perform the cytometer setup procedure during one of the incubation steps.

Diluting test samples

Purpose of this procedure The standard curve for each BD CBA Mouse or Rat Soluble Protein Flex Set covers a defined set of concentrations (10 to 2,500 pg/mL for mouse assays and 40 to 10,000 pg/mL for rat assays). It might be necessary to dilute test samples to ensure that their median fluorescence values fall within the range of the generated standard curve. For best results, dilute samples that are known or assumed to contain high levels of a given protein. This procedure is not required for all samples.

Procedure To dilute samples with known high-protein concentrations:

1. Dilute the sample by the desired dilution factor (eg, 1:10 or 1:100) using the appropriate volume of Assay Diluent.

Serum or plasma samples must be diluted at least 1:4 before transferring the samples to the assay tubes or wells.
2. Mix sample dilutions thoroughly before transferring samples to the appropriate assay tubes containing Capture Beads.
3. To facilitate analysis in FCAP Array software, load serial diluted samples in sequential wells from most concentrated to least concentrated (eg, Sample 1 – 1:4, 1:8, 1:16; Sample 2 – 1:4, 1:8, 1:16, etc).

Next step

Proceed to [Preparing Mouse/Rat Soluble Protein Flex Set Detection Reagents](#) (page 24).

Or, you can save time by proceeding directly to [Performing the Mouse or Rat Soluble Protein Flex Set Assay](#) (page 28). Please note that you will need to prepare the Detection Reagents during the first assay incubation step. If cytometer setup is required, you will also need to perform this procedure during an assay incubation step.

Preparing Mouse/Rat Soluble Protein Flex Set Detection Reagents

Purpose of this procedure

The PE Detection Reagent provided with each BD CBA Mouse or Rat Soluble Protein Flex Set is a 50X bulk concentration (1 μL per test). It should be mixed with other BD CBA Mouse or Rat Soluble Protein Flex Set PE Detection Reagents and diluted to the optimal volume per test (50 μL per test) before adding the PE Detection Reagents to a given tube or assay well.

Note: Protect the PE Detection Reagents from exposure to direct light. They can become photobleached and will lose fluorescence intensity.

Procedure

To prepare the PE Detection Reagent:

Note: You can use the same calculations for the number of tests and volume that you used for the Capture Beads in [Mixing Mouse/Rat Soluble Protein Flex Set Capture Beads \(page 21\)](#).

1. Determine the number of BD CBA Mouse or Rat Soluble Protein Flex Sets to be used in each assay tube or well in the experiment (size of the multiplex).
2. Determine the number of tests to be run in the experiment. Prepare a few additional tests than necessary to ensure that there is enough material prepared for the experiment.
3. Determine the total volume of diluted PE Detection Reagent needed for the experiment. Each tube/well requires 50 μL of the diluted PE Detection Reagent. Calculate the total volume by multiplying the number of tests (determined in [step 2](#)) by 50.

Example: 35 tests \times 50 μL = 1,750 μL total volume

4. Determine the volume needed for each PE Detection Reagent. The PE Detection Reagent is supplied so that 1.0 μL = 1 test. Therefore, the required volume (μL) of Detection Reagent is equal to the number of tests.

Example: 35 tests requires 35 μL of each Detection Reagent included in the assay

5. Determine the volume of Detection Reagent Diluent needed to dilute the PE Detection Reagents. Calculate the Diluent volume by subtracting the volume for each bead tested from the total volume of diluted beads needed to perform the assay. See [Capture Bead and Detection Reagent Diluent \(page 38\)](#).

Example: 1,750 μL total volume Detection Reagent – 35 μL for each Detection Reagent = volume of Detection Reagent Diluent

- if testing one analyte: $1,750 \mu\text{L} - (35 \mu\text{L} \times 1) = 1,715 \mu\text{L}$ diluent
 - if testing five analytes: $1,750 \mu\text{L} - (35 \mu\text{L} \times 5) = 1,575 \mu\text{L}$ diluent
6. Pipette the Detection Reagents and Detection Reagent Diluent into a tube labeled “Mixed PE Detection Reagents.” Store at 4°C , protected from light until ready to use.

Next step

Perform cytometer setup, if necessary, using the instrument setup beads. For details on setup, go to bdbiosciences.com/cbasetup and select the appropriate flow cytometer under CBA Flex Sets: Instrument Setup.

Or, if you wish to begin staining your samples for the assay, proceed to [Performing the Mouse or Rat Soluble Protein Flex Set Assay \(page 28\)](#), and you can perform the cytometer setup procedure during one of the incubation steps.

Note: It is not necessary to set up most digital cytometers before every experiment. Templates can be created by performing the setup and saving a template with the appropriate settings for subsequent experiments. When using a template, be sure to confirm the settings by running either setup beads or an extra well/tube from the assay to ensure that the template settings are acceptable.

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Assay procedure

This section covers the following topics:

- [Performing the Mouse or Rat Soluble Protein Flex Set Assay \(page 28\)](#)
- [Data analysis \(page 33\)](#)

Performing the Mouse or Rat Soluble Protein Flex Set Assay

- Before you begin**
1. Prepare the standards as described in [Preparing Mouse/Rat Soluble Protein Flex Set Standards \(page 18\)](#).
 2. Mix the Capture Beads as described in [Mixing Mouse/Rat Soluble Protein Flex Set Capture Beads \(page 21\)](#).
 3. If necessary, dilute the unknown samples. See [Diluting test samples \(page 23\)](#).
 4. Prepare the Detection Reagents as described in [Preparing Mouse/Rat Soluble Protein Flex Set Detection Reagents \(page 24\)](#). You can also prepare these reagents during the first assay incubation.

Overview

Following the preparation and dilution of the individual assay components, transfer the standards or samples, mixed Capture Beads, and mixed PE Detection Reagents to the appropriate assay wells or tubes for incubation and analysis.

Note: Protect Capture Beads and PE Detection Reagents from direct exposure to light.

Assay procedure for plates

To prepare the standards and samples for analysis using plates:

1. Wet the filter plate by adding 100 μL of Wash Buffer to each well. To remove the excess volume, apply to a vacuum manifold. Do not exceed 10" Hg of vacuum pressure. Aspirate for 2 to 10 seconds until wells are drained.
2. Add 50 μL of Mouse/Rat Soluble Protein Flex Set Standard dilutions to the first 10 wells as listed in the following table.

Well label	Standard dilution	Mouse conc. (pg/mL)	Rat conc. (pg/mL)
1	no standard dilution (Assay Diluent only)	0 (negative control)	0 (negative control)
2	1:256	10	40
3	1:128	20	80
4	1:64	40	156
5	1:32	80	312.5
6	1:16	156	625
7	1:8	312.5	1,250
8	1:4	625	2,500
9	1:2	1,250	5,000
10	Top Standard	2,500	10,000

3. Add 50 μL of each unknown sample to the appropriate wells.

To facilitate analysis in FCAP Array software, load serial diluted samples in sequential wells from most concentrated to least concentrated (eg, Sample 1 – 1:4, 1:8, 1:16; Sample 2 – 1:4, 1:8, 1:16, etc).

4. Vortex the mixed Capture Beads for at least 5 seconds and add 50 μL of the Capture Beads to each assay well. Mix the plate for 5 minutes at 500 rpm using a digital shaker (do not exceed 600 rpm).
5. Incubate the plate for 1 hour at room temperature.
Note: If you have not yet performed cytometer setup, you may wish to do so during this incubation or the incubation in [step 7](#).
6. Add 50 μL of the mixed PE Detection Reagent to each well. Mix the plate for 5 minutes at 500 rpm using a digital shaker.
7. Incubate the plate at room temperature for 1 hour for mouse assays and 2 hours for rat assays.
8. Apply the plate to the vacuum manifold and vacuum aspirate (do not exceed 10" Hg of vacuum pressure) for 2 to 10 seconds until the wells are drained.
9. Add 150 μL of Wash Buffer to each assay well. Mix the plate for 5 minutes at 500 rpm using a digital shaker to resuspend the beads.
10. Proceed to sample acquisition. See [Next step \(page 32\)](#) for helpful information on acquisition.

Assay procedure for tubes

To prepare the standards and samples for analysis using tubes:

1. Add 50 μL of Mouse/Rat Soluble Protein Flex Set Standard dilutions to the first 10 tubes as listed in the following table.

Well label	Standard dilution	Mouse Conc (pg/mL)	Rat Conc (pg/mL)
1	no standard dilution (Assay Diluent only)	0 (negative control)	0 (negative control)
2	1:256	10	40
3	1:128	20	80
4	1:64	40	156
5	1:32	80	312.5
6	1:16	156	625
7	1:8	312.5	1,250
8	1:4	625	2,500
9	1:2	1,250	5,000
10	Top Standard	2,500	10,000

2. Add 50 μL of each unknown sample to the appropriate assay tubes.
3. Vortex the mixed Capture Beads for at least 5 seconds and add 50 μL of the Capture Beads to each assay tube. Gently mix the tubes.
4. Incubate the tubes for 1 hour at room temperature.

Note: If you have not yet performed cytometer setup, you may wish to do so during this incubation or the incubation in [step 6](#).
5. Add 50 μL of the mixed PE Detection Reagent to each assay tube. Gently mix the tubes.

6. Incubate the tubes at room temperature for 1 hour for mouse assays and 2 hours for rat assays.
7. Add 1 mL of Wash Buffer to each assay tube and centrifuge at 200g for 5 minutes.
8. Carefully aspirate and discard the supernatant from each assay tube.
9. Add 300 μ L of Wash Buffer to each assay tube. Vortex the assay tubes briefly to resuspend the beads.

Next step

Acquire the samples on the flow cytometer. For details, go to bdbiosciences.com/cbasetup and select the appropriate flow cytometer under CBA Flex Sets: Instrument Setup.

Acquire samples on the same day they are prepared. Prolonged storage of samples, once the assay is complete, can lead to increased background and reduced sensitivity.

To facilitate the analysis of samples in FCAP Array software, we recommend the following guidelines:

- Acquire standards from lowest (0 pg/mL) to highest (Top Standard) concentration, followed by the test samples.
 - If running sample dilutions, acquire sequentially starting with the most concentrated sample (eg, Sample 1 – 1:4, 1:8, 1:16; Sample 2 – 1:4, 1:8, 1:16, etc).
 - Store all FCS files (standards and samples) in a single folder.
-

Data analysis

How to analyze data Analyze data using FCAP Array software. For instructions on analysis, go to bdbiosciences.com/cbasetup and see the *FCAP Array Software User's Guide*.

When analyzing the BD CBA Mouse/Rat Soluble Protein assay data with FCAP Array software and choosing a curve fitting option, try both 4 Parameter and 5 Parameter Logistic, and select the one that results in the best curve fit (highest R^2 value).

5

Reference

This section covers the following topics:

- [Troubleshooting \(page 36\)](#)
- [Capture Bead and Detection Reagent Diluent \(page 38\)](#)
- [References \(page 41\)](#)

Troubleshooting

Recommended actions These are the actions we recommend taking if you encounter the following problems.

Note: For best performance, vortex samples immediately before acquiring on a flow cytometer.

Problem	Recommended action
Poor standard curves	If there is no change in signal above background across the entire standard curve range, ensure that all of the components (Capture Beads, Detection Reagent, and standard) were added to each tube.
	If the curve is relatively flat and then increases at higher concentrations but not to the expected levels, make sure standards are not being vortexed or vigorously mixed while being reconstituted. The best approach is to allow the standards to equilibrate for 15 minutes in Assay Diluent prior to mixing. Mix by gently pipetting several times.
	Check that all components have been properly prepared and stored. Use freshly reconstituted standards. Ensure that incubation times are of proper length and that the assay did not sit for a prolonged period of time after the wash step.
Low event count	The beads can aggregate. Thoroughly vortex individual Capture Bead bulk vials prior to preparation of the master bead mix and vortex the master bead mix prior to dispensing into the individual assay wells. Thoroughly shake plates or vortex sample tubes prior to acquisition.
	Ensure that the stopping rule, singlet gate, and thresholds are set correctly.
	Ensure that the vacuum is not too strong and that filter membranes are not compromised (filter plates). Avoid aspiration of beads during the wash step (tubes).
Variation between duplicate samples	Vortex Capture Beads before pipetting. The beads can aggregate.

Problem	Recommended action
Little or no detection of protein in samples	Samples might be too dilute. Try various sample dilutions.
All samples are positive or above the high standard mean fluorescence value	Samples might be too concentrated. Try various sample dilutions.
High background	Remove excess detection reagent by increasing the number of wash steps, since the background might be due to non-specific binding.
	Background can be produced by precipitated buffers. Check for visible precipitate and filter through a 0.2- μ m filter, if necessary.
Sample dilution	We recommend diluting serum and plasma samples at least 1:4 because spike recoveries are generally better, suggesting that factors might be present at lower dilutions that inhibit the binding kinetics of the assay. If using the filter plate protocol, diluting the samples also prevents clogging of the filter membrane, which can lead to insufficient washing and high background. If using a BD FACSCalibur system, diluting the samples along with adding additional standard dilutions ensures that sample MFIs fall on the linear portion of the curve and prevents spillover of excessive PE signal into the FL3 channel, which can cause gating issues during analysis.
Sample storage	Cytokines in general are quite labile and will degrade over time even when stored frozen at -70°C . Samples can usually be stored in single-use aliquots for 3 to 6 months. Sample storage strategies should be determined empirically prior to making them standard practice.

Problem	Recommended action
Biohazardous samples	It is possible to treat samples with 1% paraformaldehyde before analyzing on the flow cytometer. This might affect assay performance and should be validated. The antibody pairs are optimized for detection of native protein, so fixation should be attempted only after the samples have been incubated with the Capture Beads and Detection Reagents.
Anticoagulant for plasma samples	Only EDTA plasma samples have been verified by BD Biosciences.
Clogged filter plate	Serum and plasma proteins can settle and clog the membrane during incubation. Dilute samples further or perform assay incubations in a standard polystyrene U-bottom plate (Catalog No. 353910) and transfer to the filter plate immediately prior to aspiration. Resuspend the beads well prior to transfer.

Capture Bead and Detection Reagent Diluent

How to calculate Calculate the Diluent volume by subtracting the volume for each bead tested from the total volume of diluted

beads needed to perform the assay. The following table lists the appropriate volumes.

No. of Flex Sets to be used	Volume per test			
	Each Capture Bead or Detection Reagent	Total Capture Bead or Detection Reagent	Capture Bead or Detection Reagent Diluent	Mixed Capture Beads or Detection Reagent
1	1 µL	1 µL	49 µL	50 µL
2	1 µL	2 µL	48 µL	50 µL
3	1 µL	3 µL	47 µL	50 µL
4	1 µL	4 µL	46 µL	50 µL
5	1 µL	5 µL	45 µL	50 µL
6	1 µL	6 µL	44 µL	50 µL
7	1 µL	7 µL	43 µL	50 µL
8	1 µL	8 µL	42 µL	50 µL
9	1 µL	9 µL	41 µL	50 µL
10	1 µL	10 µL	40 µL	50 µL
11	1 µL	11 µL	39 µL	50 µL
12	1 µL	12 µL	38 µL	50 µL
13	1 µL	13 µL	37 µL	50 µL
14	1 µL	14 µL	36 µL	50 µL
15	1 µL	15 µL	35 µL	50 µL
16	1 µL	16 µL	34 µL	50 µL
17	1 µL	17 µL	33 µL	50 µL
18	1 µL	18 µL	32 µL	50 µL
19	1 µL	19 µL	31 µL	50 µL
20	1 µL	20 µL	30 µL	50 µL

No. of Flex Sets to be used	Volume per test			
	Each Capture Bead or Detection Reagent	Total Capture Bead or Detection Reagent	Capture Bead or Detection Reagent Diluent	Mixed Capture Beads or Detection Reagent
21	1 µL	21 µL	29 µL	50 µL
22	1 µL	22 µL	28 µL	50 µL
23	1 µL	23 µL	27 µL	50 µL
24	1 µL	24 µL	26 µL	50 µL
25	1 µL	25 µL	25 µL	50 µL
26	1 µL	26 µL	24 µL	50 µL
27	1 µL	27 µL	23 µL	50 µL
28	1 µL	28 µL	22 µL	50 µL
29	1 µL	29 µL	21 µL	50 µL
30	1 µL	30 µL	20 µL	50 µL
31	1 µL	31 µL	19 µL	50 µL
32	1 µL	32 µL	18 µL	50 µL
33	1 µL	33 µL	17 µL	50 µL
34	1 µL	34 µL	16 µL	50 µL
35	1 µL	35 µL	15 µL	50 µL
36	1 µL	36 µL	14 µL	50 µL
37	1 µL	37 µL	13 µL	50 µL
38	1 µL	38 µL	12 µL	50 µL
39	1 µL	39 µL	11 µL	50 µL
40	1 µL	40 µL	10 µL	50 µL
41	1 µL	41 µL	9 µL	50 µL
42	1 µL	42 µL	8 µL	50 µL
43	1 µL	43 µL	7 µL	50 µL
44	1 µL	44 µL	6 µL	50 µL
45	1 µL	45 µL	5 µL	50 µL
46	1 µL	46 µL	4 µL	50 µL
47	1 µL	47 µL	3 µL	50 µL
48	1 µL	48 µL	2 µL	50 µL
49	1 µL	49 µL	1 µL	50 µL
50	1 µL	50 µL	0 µL	50 µL

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