BD™ Cytometric Bead Array (CBA)



Technical Data Sheet Human IL-1α Flex Set

Product Information

- Material Number: Size: Bead Position: Assay Range: Reactivity:
- 560153 100 Tests D6 10-2,500 pg/mL QC Testing: Human

Component Description: Component Mat. No: Component Storage Buffer:

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51-9005594 Aqueous buffered solution containing BSA and $\leq 0.09\%$ sodium azide.

Human IL-1a PE Detection Reagent

Human IL-1 α Capture Bead D6 51-9005596 Aqueous buffered solution containing fetal bovine serum and $\leq 0.09\%$ sodium azide.

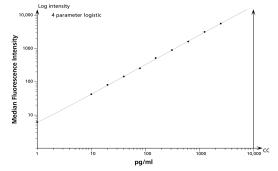


Figure 1. Example BD CBA Human IL-1α Flex Set standard curve. Data acquired on a BD FACSArray bioanalyzer and analyzed using FCAP Array Software.

Description

The BDTM CBA Human IL-1 α Flex Set is a bead-based immunoassay capable of measuring human interleukin-1 α (IL-1 α) in serum, plasma, and cell culture supernatant samples. Human reactivity was determined by testing samples with the BD CBA Human IL-1 α Flex Set. The biology and function of IL-1 α has been extensively reviewed in the literature. For more information on bead-based immunoassays, refer to the product insert for the BD CBA Human Soluble Protein Master Buffer Kit (Cat. No. 558264 or 558265).

Preparation and Storage

This BDTM CBA Flex Set contains one vial each of Capture Bead and PE Detection Reagent and two vials of Standard. The Capture Bead and PE Detection Reagent components of this flex set have been formulated to a 50x concentration to ensure product performance when multiplexed. The Standard component is lyophilized and should be transferred to a 15 mL polypropylene tube for reconstitution. When reconstituted in 4.0 mL Assay Diluent, the standard has a protein concentration of 2,500 pg/mL. Discard unused reconstituted standard, do not store or reuse. Store lyophilized standard and other components at 4°C. Protect Capture Beads and the PE Detection Reagent from prolonged exposure to light.

Application Notes

Recommended Assay Procedure: The BD CBA Human IL-1 α 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 ArrayTM 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-1 α Flex Set will be 2,500 pg/mL. An example standard curve is shown in Figure 1.

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The BD CBA Human IL-1 α Flex Set should not be used in the same assay well with any non-BD CBA Human Soluble Protein Flex Set reagents (such as BD CBA Mouse Soluble Protein or Cell Signaling Flex Sets). For an updated assay compatibility chart for the BD CBA Human Soluble Protein Flex Sets, please refer to the BD CBA Flex Set System homepage at http://www.bdbiosciences.com/cbasetup.

Performance

Limit of Detection: The theoretical limit of detection is 1.0 pg/mL and was determined by evaluating the estimated result of the average MFI of the negative control (0 pg/mL, n=30) + 2 standard deviations.

Specificity		Inter-Assay Reproducibility			Intra-Assay Reproducibility			
		Mean (pg/mL)	Standard Deviation	%CV	Mean (pg/mL)	Standard Deviation	%CV	
	Sample 1	36.2	3.2	9%	31.8	0.4	1%	
Human IL-1 α	Sample 2	140.9	12.0	9%	123.1	2.6	2%	
	Sample 3	577.5	32.9	6%	546.4	14.2	3%	

Reproducibility: The intra-assay and inter-assay reproducibility were determined for the BD CBA Human IL-1α Flex Set by evaluating ten replicates of three different sample levels (intra-assay) and three replicates of three different sample levels from four separate experiments (inter-assay).

	Cell Culture Supernatant		Serum		Plasma	
Specificity	Average % Recovery	Range	Average % Recovery	Range	Average % Recovery	Range
Human IL-1 α	99%	95 - 102%	66%	62 - 71%	73%	67 - 80%

Recovery: Cell culture supernatant, serum, or EDTA-treated plasma were spiked with three different levels of protein. The spiked samples were assayed and the results were compared with expected values. Serum and plasma samples were diluted 1:4 before the protein was spiked into each. Serum is a pool of 800 - 1000 donors and the plasma was pooled from at least 20 donors.

Comula Dilution	Cell Culture Supernatant		Ser	um	Plasma	
Sample Dilution	Detected (pg/mL)	% of Expected	Detected (pg/mL)	% of Expected	Detected (pg/mL)	% of Expected
Spiked sample	658.0	100%	412.5	100%	463.3	100%
1:2	305.7	93%	224.3	109%	226.1	102%
1:4	147.0	89%	116.7	113%	117.1	101%

Linearity: Cell culture supernatant, 1:4 diluted serum, or 1:4 diluted EDTA-treated plasma were spiked with protein and serially diluted. The diluted samples were assayed and the results were compared with the original spiked sample.

Product Notices

- 1. ProClin is a trademark of Rohm and Haas Company.
- This product is manufactured and sold under license from Pestka Biomedical Laboratories, Inc. (d/b/a PBL InterferonSource) and may be used solely as indicated. This product may not be resold or incorporated in any manner into another product for resale. Any use for therapeutics is strictly prohibited. This product is covered by U.S. Patent No. 5,831,022.
- 3. Source of all serum proteins is from USDA inspected abattoirs located in the United States.
- 4. Caution: 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.
- 5. Please refer to www.bdbiosciences.com/pharmingen/protocols for technical protocols.
- Warning: CBA lyophilized standard contains 0.02% (w/w) of a CMIT/MIT mixture (3:1), which is a mixture of: 5-chloro-2-methyl-4-isothiazolin-3-one [EC No 247-500-7] and 2-methyl-4-isothiazolin-3-one [EC No 220-239-6] (3:1). Hazard statement: May cause an allergic skin reaction.

Precautionary statements: Wear protective gloves/eye protection. Wear protective clothing. Avoid breathing mist/vapours/spray. If skin irritation or rash occurs: Get medical advice/attention. IF ON SKIN: Wash with plenty of water. Dispose of contents/container in accordance with local/regional/national/international regulations.

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