

Technical Data Sheet

Human IL-12p70 Flex Set

Product Information

Material Number:558283Size:100 TestsBead Position:E5

 Assay Range:
 10-2,500 pg/mL

 Reactivity:
 QC Testing: Human

Component Description: Human IL-12p70 Standard

Component Mat. No: 51-9003505

Component Storage Buffer: Lyophilized in an aqueous buffered solution containing BSA

and ProClin™ 150.

Component Description: Human IL-12p70 PE Detection Reagent

Component Mat. No: 51-9004063

Component Storage Buffer: Aqueous buffered solution containing BSA and ≤0.09%

sodium azide.

Component Description: Human IL-12p70 Capture Bead E5

Component Mat. No: 51-9005304

Component Storage Buffer: Aqueous buffered solution containing fetal bovine serum

and ≤0.09% sodium azide.

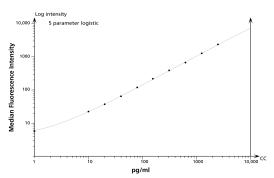


Figure 1. Example BD CBA Human IL-12p70 Flex Set standard curve. Data acquired on a BD FACSArray bioanalyzer and analyzed using FCAP Array Software.

Description

The BDTM CBA Human IL-12 p70 Flex Set is a bead based immunoassay capable of measuring human interleukin-12p70 (IL-12p70) in serum, plasma, and cell culture supernatant samples. Human reactivity was determined by testing samples with the BD CBA Human IL-12p70 Flex Set. The biology and function of IL-12p70 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-12p70 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-12p70 Flex Set will be 2,500 pg/mL. An example standard curve is shown in Figure 1.

The BD CBA Human IL-12p70 Flex Set cannot be used in the same assay well with the following BD CBA Human Soluble Protein Flex Set

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Flex Set Bead Position Catalog Number BD CBA Human IL-12/IL-23p40 Flex Set E5 560154

The BD CBA Human IL-12p70 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 0.6 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.

		Inter-Assay Reproducibility			Intra-Assay Reproducibility			
Specificit	У	Mean (pg/ml)	Standard Deviation	%CV	Mean (pg/ml)	Standard Deviation	%CV	
Human IL-12p70	Sample 1	39.4	1.8	4%	39.7	0.9	2%	
	Sample 2	151.3	3.1	2%	155.6	7.1	5%	
	Sample 3	623.7	20.3	3%	640.8	20.0	3%	

Reproducibility: The inter-assay and intra-assay reproducibility were determined for the BD CBA Human IL-12p70 Flex Set by evaluating ten replicates of three different sample levels (intra-assay) and two replicates of three different sample level from four separate experiments (inter-assay).

Specificity	Cell Culture Supernatant		Ser	um	Plasma		
	Average % Recovery	Range	Average % Recovery	Range	Average % Recovery	Range	
Human IL-12p70	83%	80 - 85%	76%	64 - 86%	67%	56 - 74%	

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.

Sample Dilution	Cell Culture Supernatant		Ser	um	Plasma		
	Detected (pg/ml)	% of Expected	Detected (pg/ml)	% of Expected	Detected (pg/ml)	% of Expected	
Spiked sample	583.1	100%	580.6	100%	543.3	100%	
1:2	286.7	98%	296.9	102%	279.7	103%	
1:4	134.1	92%	145.7	100%	144.1	106%	

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.
- 2. Source of all serum proteins is from USDA inspected abattoirs located in the United States.
- 3. 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.
- 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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