

Magnetic Luminex® Performance Assay Human Osteopontin (OPN) Kit

Catalog Number: LHK1433
Pack Size: 100 Tests

SPECIFICATIONS AND USE

Recommended Sample Types Microparticle Region Components

- Serum, EDTA plasma, heparin plasma, and urine.
- Region-28
- Microparticle Concentrate (Part 894305) is supplied as a 100X concentrated stock (0.075 mL) with preservatives.
- Biotin-Antibody Concentrate (Part 894316) is supplied as a 100X concentrated stock solution (0.075 mL) with preservatives.

Other Supplies Required

 Magnetic Luminex Performance Assay Human Kidney Biomarker Base Kit (Catalog Number LHK000).

Storage

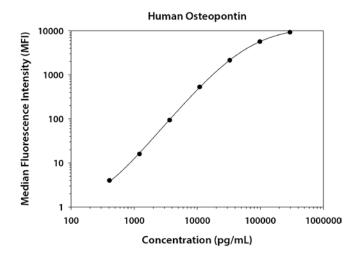
- Store the unopened kit at 2-8 °C. Do not use past the expiration date on the label.
- Avoid freezing microparticles.
- Protect microparticles from light.

Instructions for Use

Refer to the Base Kit insert for the Magnetic Luminex Performance Assay procedure.

TYPICAL DATA

This human Osteopontin standard curve is provided only for demonstration. A standard curve must be generated each time an assay is run, utilizing values from the Standard Value Card included in the Base Kit.



Standard	pg/mL	MFI	Average	Corrected
Blank	0	11 11	11	
1	295,500	9202 9324	9263	9252
2	98,500	5657 5741	5699	5688
3	32,833	2128 2179	2154	2143
4	10,944	533 548	541	530
5	3648	105 105	105	94
6	1216	27 27	27	16
7	405	15 15	15	4

PERFORMANCE CHARACTERISTICS

All data were collected with assays run as a multiplex.

Sensitivity - The Minimum Detectable Dose (MDD) was determined by adding two standard deviations to the MFI of twenty zero standard replicates and calculating the corresponding concentration.

Thirty-two assays were evaluated, and the MDD of human Osteopontin ranged from 8.1-247 pg/mL. The mean MDD was 133 pg/mL.

CORRELATION

This assay has been correlated to the respective Quantikine® ELISA kit with a slope of 0.9-1.1 and an R² value greater than 0.9.

FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES.

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Intra-assay Precision (precision within an assay) - Three samples of known concentration were tested twenty times on one plate to assess precision within an assay.

Inter-assay Precision (precision between assays) - Three samples of known concentration were tested in separate assays to assess precision between assays. Assays were performed by at least three technicians using two lots of components.

	Intra-assay Precision			Inter-assay Precision		
Sample	1	2	3	1	2	3
n	20	20	20	72	70	72
Mean (pg/mL)	2702	17,630	121,750	2717	17,696	118,682
Standard Deviation	59.6	232	3919	277	1031	11,549
% CV	2.2	1.3	3.2	10.2	5.8	9.7

Linearity - Samples containing and/or spiked with high concentrations of Osteopontin were serially diluted to evaluate assay linearity.

		Serum (n=4)	EDTA plasma (n=4)	Heparin plasma (n=4)	Urine (n=4)
1.0	Average % of Expected	106	106	96	97
1:2	Range (%)	101-111	101-110	94-97	90-102
1:4	Average % of Expected	108	102	96	97
	Range (%)	97-120	101-102	91-102	93-100
1:8	Average % of Expected	110	103	73	102
	Range (%)	96-133	101-104	63-83	100-104

Specificity - This assay recognizes natural and recombinant human Osteopontin. The assay was tested for cross-reactivity and interference with the following factors. Less than 0.5% cross-reactivity and interference was observed.

Recombinant human:				Recombinant mouse:	Other recombinants:	Recombinant human multiplex partners:
ApoA1	Cathepsin O	CXCL2/GROB	HPRG	Clusterin	bovine Osteopontin	Clusterin
ApoA2	Cathepsin S	CXCL3/GROγ	IFN-γ	Cystatin C	·	Cystatin C
ApoB	Cathepsin V	CXCL5/ENA-78	Lipocalin-1	CXCL10/IP-10/CRG-2	Natural human	Lipocalin-2/NGAL
ApoB100	Cathepsin Z	CXCL6/GCP-2	MMP-3	HGF	proteins:	CXCL10/IP-10
ApoC1	CCL2/MCP-1	CXCL7/NAP-2	MMP-7	Lipocalin-2/NGAL	lpha1-Acid Glycoprotein	HGF
ApoC2	CCL5/RANTES	CXCL8/IL-8	MMP-9	Osteopontin (OPN)	Kininogen	Fetuin A
ApoD	Cystatin A	CXCL9/MIG	MSP	TIM-1/KIM-1/HAVCR		RBP4
ApoE	Cystatin B	CXCL11/I-TAC	Plasminogen			TFF3
АроН	Cystatin E/M	CXCL12/SDF-1	Serpin A1	Recombinant		TIM-1/KIM-1/HAVCR
ApoM	Cystatin F	CXCL13/BLC/BCA-1	TIM-3	rat:		
Cathepsin A	Cystatin S	Enterokinase	TIM-4	Clusterin		
Cathepsin B	Cystatin SA	Fetuin B	TFF-1	Fetuin A		
Cathepsin C	Cystatin SN	Fibronectin	TFF-2	TIM-1/KIM-1/HAVCR		
Cathepsin D	Clusterin-like 1	HAI-1	Thrombin			
Cathepsin E	COX-2	HAI-2				
Cathepsin F	CRP	HGF R/c-MET				
Cathepsin L	CXCL1/GR0 $lpha$	HGF Activator				

TECHNICAL HINTS

- Protect the microparticles and streptavidin-PE from light at all times.
- Refer to the Base Kit Standard Value Card for reconstitution volume and values of the reconstituted standard.
- Diluted microparticles cannot be stored. Make a fresh dilution of microparticles each time the assay is run.
- The use of a magnetic device made to accommodate a microplate is necessary for washing.
- Discrepancies may exist in values obtained for the same analyte utilizing different technologies.

Luminex Performance Assays afford the user the benefit of multianalyte analysis of biomarkers in a complex sample. A single, multipurpose diluent is used to optimize recovery, linearity, and reproducibility. Such a multipurpose diluent may not optimize any single analyte to the same degree that a unique diluent selected for analysis of that analyte can optimize conditions. Therefore, some performance characteristics may be more variable than those for assays designed specifically for single analyte analysis.