

Cytokine Report – Luminex 34-plex Panel

ORDER DETAILS

Patient Name	HELIX, Onco	Physician	Dr. XXXXX XXXXX
Medical Record#	XXX-XXXXX-XXX	Organization	University Hospital
Date of Birth	XX-Feb-1955	Address	
Gender	M		
Diagnosis	COVID-19		

SPECIMEN AND TEST

Specimen Type	Frozen Serum	Date Received	1-Apr-2020	Report Status	Final
Specimen ID	XX-XXX-XXXX	Analysis Date	2-Apr-2020	Report Date	5-Apr-2020

REPORT SUMMARY

No.	Cytokine	Results (pg/mL)		
1	IFN-gamma	33.32	Moderate	
2	IL-12p70	4.04	Very Low	
3	IL-13	-	ND	
4	IL-1-beta	4.16	Low	
5	IL-2	101.87	High	
6	IL-4	74.34	High	
7	IL-5	105.61	Moderate	
8	IL-6	150	High	
9	TNF-alpha	56.02	Moderate	
10	GM-CSF	-	ND	
11	IL-18	13.45	Very Low	
12	IL-10	6.25	Moderate	
13	IL-17A	65.43	High	
14	IL-21	-	ND	
15	IL-22	-	ND	
16	IL-23	-	ND	
17	IL-27	375.41	High	

No.	Cytokine	Results (pg/mL)		
18	IL-9	-	ND	
19	IFN-alpha	-	ND	
20	IL-31	10.84	High	
21	IL-15	72.01	Moderate	
22	IL-1-alpha	2.48	Very Low	
23	IL-1RA	2498.3	Moderate	
24	IL-7	3.06	Moderate	
25	TNF-beta	-	ND	
26	Eotaxin	128.36	High	
27	GRO-alpha	-	ND	
28	IL-8	400	High	
29	IP-10	218.91	High	
30	MCP-1	14.06	Low	
31	MIP-1-alpha	-	ND	
32	MIP-1-beta	70.94	Moderate	
33	SDF-1-alpha	890.44	High	
34	RANTES	153.1	High	

Code	Label	Definition	Description
	ND	Not detected	Value was not detected in the assay
	Very Low	< Lower limit of detection (LLOD)	Value smaller than the lower limit of detection
	Low	0-30th percentile	30% of the reference samples have values less than the cutoff
	Moderate	30th-70th percentile	40% of the reference samples fall in this range
	High	> 70th percentile	Value higher than 70% of the reference samples.
	Very High	> Upper limit of detection (ULOD)	Higher than upper limit of detection

COMPARISON WITH PREVIOUS RESULTS

No.	Cytokine	01-Apr-20		25-Mar-20		Date	
		Results (pg/mL)		Results (pg/mL)		Results (pg/mL)	
1	IFN-gamma	33.32	Moderate	24.25	Low		
2	IL-12p70	4.04	Very Low	3.3	Very Low		
3	IL-13	-	ND	-	ND		
4	IL-1-beta	4.16	Low	3.2	Low		
5	IL-2	101.87	High	71.84	Moderate		
6	IL-4	74.34	High	43.67	Moderate		
7	IL-5	105.61	Moderate	77.5	Low		
8	IL-6	150	High	120.68	Moderate		
9	TNF-alpha	56.02	Moderate	42.8	Moderate		
10	GM-CSF	-	ND	-	ND		
11	IL-18	13.45	Very Low	2.2	Very Low		
12	IL-10	6.25	Moderate	20.4	High		
13	IL-17A	65.43	High	52.4	Moderate		
14	IL-21	-	ND	-	ND		
15	IL-22	-	ND	-	ND		
16	IL-23	-	ND	-	ND		
17	IL-27	375.41	High	249.7	High		
18	IL-9	-	ND	-	ND		
19	IFN-alpha	-	ND	-	ND		
20	IL-31	10.84	High	5.2	High		
21	IL-15	72.01	Moderate	51.5	Moderate		
22	IL-1-alpha	2.48	Very Low	1.1	Very Low		
23	IL-1RA	2498.3	Moderate	1854.6	Moderate		
24	IL-7	3.06	Moderate	1.7	Low		
25	TNF-beta	-	ND	-	ND		
26	Eotaxin	128.36	High	60	Moderate		
27	GRO-alpha	-	ND	-	ND		
28	IL-8	400	High	292.3	Moderate		
29	IP-10	218.91	High	111.8	High		
30	MCP-1	14.06	Low	12.5	Low		
31	MIP-1-alpha	-	ND	-	ND		
32	MIP-1-beta	70.94	Moderate	55.68	Low		
33	SDF-1-alpha	890.44	High	550	High		
34	RANTES	153.1	Very High	135	High		

COMMENT

There is a significant increase in the level of many cytokines in the current specimens when compared to the previous specimen of the patient.

METHOD

Multiplex immunoassay using Luminex xMAP (multi-analyte profiling) technology was performed on the submitted serum sample to quantitate 34 cytokine and chemokine protein targets simultaneously using Luminex instrument. The assay is based on the principle of sandwich ELISA (Enzyme Linked Immunosorbent Assay). A 5-parameter (5-P) logistic regression model was generated from the mean florescence intensity(MFI) detected by the Luminex instrument and the known standards for each analyte. Standards with a serial dilution of 1:4 were used for generating the model. Criteria of acceptance of the model are R² value >95%, percent recovery of each point of the standard curve < 30%, and percent CV of duplicates <30%. MFI values of the unknown samples were compared to the 5-P logistic regression curve of each cytokine to calculate the levels. Interpretation of the results (low, moderate or high) was based on the comparison of the cytokine levels generated from 136 healthy individuals using the same cytokine panel in Hematology Translational Lab(HTL). Details of lower and upper limit of detection along with the healthy range can be provided upon request. Test performance characteristics for this laboratory validated test has been determined by the College of Physicians and Surgeons of Alberta (CPSA) accredited laboratory.

DISCLAIMER

This report is based on the assumption that the sample received from the individual noted by the unique identifiers and has not been contaminated with that of another individual prior to receipt at the Hematology Translational Lab(HTL). It is assumed that the serum sample has been processed by following the strict guidelines recommended by HTL. Absolute value of the cytokine levels can vary based on kit used and the lab in which the assay is performed. Variability in the absolute value of the cytokine levels between different runs was less than 25%. Cytokine levels of IL23, TNF-beta and IL31 were not detected in healthy individuals and therefore the interpretations for these cytokines in the analyzed samples should be considered with caution. Results from this panel should be interpreted in the context of all clinical, biochemical, genetic and other molecular data.

REVIEW AND APPROVAL

Report Reviewed by

Clinical Scientist

Verified and Approved by

Laboratory Director

Medical Director

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