Simoa® IL-10 Advantage PLUS Kit HD-X Data Sheet Item 105259

Description

This data sheet summarizes data from the analytical validation performed at Quanterix to characterize performance of the IL-10 Advantage PLUS kit on the HD-X platform.

IL-10

Calibration Curve: Representative calibrator concentrations and Lower Limit of Quantification (LLOQ) depicted in Figure 1. The assigned concentrations of calibrator levels and reconstitution volume may vary between different kit lots.

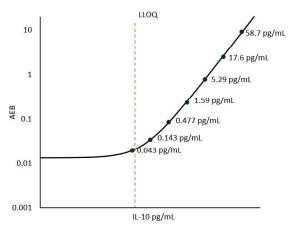


Figure 1. Example calibrator curve.

Minimum Required Dilution (MRD)

Diluted Sample Volume	100 μL per measurement	
Serum/EDTA Plasma Dilution	1:4	
Tests per Kit	96	

See Kit Instruction for details.

Lower Limit of Quantification (LLOQ): The analytical LLOQ was set at the lowest concentration that read back within 80-120% of the expected value with a CV $\leq 20\%$. The functional LLOQ (fLLOQ) values below are for serum and EDTA plasma and represent the analytical LLOQ multiplied by the dilution factor used for the samples.

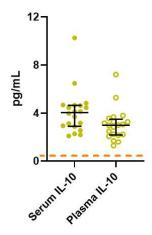
Limit of Detection (LOD): The LOD is calculated as 2.5 standard deviations from the mean of background signal read back on each calibration curve.

Assay Range: The upper end of the dynamic range is equal to the top calibrator concentration multiplied by MRD. The representative ranges below are for serum and EDTA plasma.

IL-10		
Analytical	0.048 pg/mL Pooled CV: 20%	
LLOQ	Mean Recovery: 92%	
Functional LLOQ	Serum/EDTA Plasma (4x): 0.192 pg/mL	
LOD	0.015 pg/mL Range: 0.01 - 0.021 pg/mL	
Dynamic Range	Serum/EDTA Plasma (4x): 0 - 300 pg/mL	

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Endogenous Sample Reading: Healthy donor matched EDTA plasma (n=20), and serum (n=20) concentrations (pg/mL) were measured using the IL-10 Advantage PLUS kit on HD-X. Bars depict median with interquartile range. The orange line is a representative value for functional LLOQ, which may vary from lot to lot.



IL-10				
Sample Type	Mean (pg/mL)	Median (pg/mL)	% Above LOD	% Above fLLOQ
Serum	4.09	4.05	100%	100%
EDTA Plasma	3.06	3.01	100%	100%

Precision: Measurements of 2 calibrator-based controls, 3 commercial pooled serum and 3 commercial pooled plasma. Triplicate measurements were made for 2 runs each for 1 reagent lot across 2 instruments (4 runs total, 12 measurements). All samples were diluted at the appropriate MRD for the sample matrix.

IL-10				
Sample	Mean (pg/mL)	Within Run CV	Between Run CV	Between Instr CV
Control 1	1.36	3.2%	5.9%	5.0%
Control 2	74.3	1.9%	6.1%	0.7%
Plasma 1	1.85	4.7%	12%	9.4%
Plasma 2	216	2.7%	7.3%	4.6%
Plasma 3	211	3.2%	7.8%	6.4%
Serum 1	0.754	6.8%	2.6%	0.9%
Serum 2	140	6.0%	9.4%	11%
Serum 3	156	3.9%	7.4%	3.0%

Spike and Recovery: 2 serum and 2 EDTA plasma samples were spiked at high and low concentrations of IL-10 within the range of each assay and analyzed on HD-X. Percent recovery is defined as the difference between the measured concentration of the analytes in the spiked sample and the measured concentration in unspiked sample relative to the concentration of the analytes in spiked calibrator diluent.

Dilution Linearity: 2 serum and 2 EDTA plasma samples were spiked with endogenous antigen and then diluted 2x serially with sample diluent. Total dilution of each sample ranged from 4x to 64x.

IL-10 †			
Spike and Recovery	Mean: 93%		
Serum	Range: 85 – 100%		
Spike and Recovery	Mean: 79%		
EDTA Plasma	Range: 71 – 88%		
Dilution Linearity	Mean: 104%		
Serum (4X-64X)	Range: 88 - 141%		
Dilution Linearity EDTA	Mean: 96%		
Plasma (4X-64X)	Range: 88 - 107%		

[†] The assay design is conserved between C4PA Advantage PLUS and IL-10 Advantage PLUS. Values are sourced from the C4PA Advantage PLUS Validation report.

The Simoa® IL-10 Advantage PLUS assay kit is formulated for use on the HD-X platform. Verification and validation results for the fully automated HD-X instrument are summarized in this report.