)-PLEX[®] NHP IFN-γ

MSD Customer Service Phone: 1-240-314-2795 Fax: 1-301-990-2776 Email: CustomerService@ mesoscale.com

Scientific Support

Phone: 1-240-314-2798 Email: ScientificSupport@

Company Address MESO SCALE DISCOVERY®

Meso Scale Diagnostics, LLC.

1601 Research Boulevard Rockville, MD 20850-3173 USA

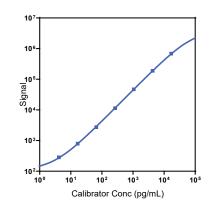
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www.mesoscale.com [®] Ordering Information MSD Customer Service Phone: 1-240-314-2795 Fax: 1-301-990-2776 Email: CustomerService@ magacable.com	Product Options	Catalog Number	Description		
	Multiplex	K15068M, K25068M	U-PLEX Biomarker Group 1 (NHP)		
		K151TTK-1/-2/-4	U-PLEX NHP IFN-γ Assay with SECTOR [™] plates		
	Singleplex	K156TTK-21	U-PLEX NHP IFN- γ Assay with QuickPlex® APT plates		
		K256TTK-2/-4	U-PLEX NHP IFN- γ Assay with 384-well plates		
	Antibody Set	B21TT-2/-3	U-PLEX Human IFN-y Antibody Set		
	Assay Protocol	U-PLEX Product Inserts are available at www.mesoscale.com			

The U-PLEX® platform was designed to provide ultimate flexibility for detection of biomarkers in a wide variety of sample types. This datasheet provides the representative performance of the U-PLEX NHP IFN-y Assay tested on U-PLEX 96-well SECTOR plates run as a multiplex. The data do not represent the product specifications. Under your experimental conditions, the assay may perform differently from the representative data. U-PLEX assays are offered in either singleplex or multiplex; both are available in 96- or 384well plates. See a U-PLEX product insert for instrument compatibility.

Representative Calibration Curve and Sensitivity



Assay	Median LLOD (pg/mL)	LLOD Range (pg/mL)		
IFN-γ	1.07	0.68-1.35		

The Calibrator curve was fitted with a 4-parameter logistic model with a $1/Y^2$ weighting. The lower limit of detection (LLOD) is a calculated concentration corresponding to 2.5X the standard deviations above the background (zero Calibrator).

Precision

	Control	Average Conc. (pg/mL)	Average Intra-run Conc. %CV	Inter-run Conc. %CV	
	High	11,300	5.9	8.4	
IFN-γ	Mid	1,120	3.6	9.5	
	Low	108	5.3	12.6	

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Controls were made by spiking Calibrator into assay diluent at 3 levels within the quantitative range of the assay. Average intra-run concentration %CV is the average %CV of the control replicates within an individual run. Inter-run concentration %CV is the variability of controls across multiple runs.

Spike Recovery

		Serum (N=5)		Plasma (N=5)		Cell Culture Media (N=5)	
Spike Level		Average % % Recovery Recovery Range		Average % % Recovery Recovery Range		Average % % Recovery Range	
Cynomolgus Monkey	High	72.7	37-93	90.8	80-99	121	116-128
	Mid	74.3	47-93	91.9	80-99	113	106-120
	Low	72.8	49-87	88.3	76-95	114	109-119
Rhesus Monkey	High	101.3	89-117	107	92-122	121	116-128
	Mid	99.8	81-114	106.2	95-114	113	106-120
	Low	99.1	70-116	104.6	90-113	114	109-119

Normal serum, EDTA plasma, and cell culture media were spiked with Calibrator at 3 levels. Undiluted samples were tested to determine the expected concentration of the analyte. Samples may benefit from additional dilution with assay diluent to reduce matrix effects.

% Recovery = (measured concentration / expected concentration) x 100

Tested Samples

	Sample Type	Serum (N=10)	Plasma (N=10)	Spiked Serum (N=5)
Ormonialaria	Median (pg/mL)	ND	NA	5.64
Cynomolgus Monkey	Range (pg/mL)	ND-5.79	NA	ND-645
	% Detected	10	0	60
Dhassa	Median (pg/mL)	NA	ND	105
Rhesus Monkey	Range (pg/mL)	NA	ND-1.70	1.80-208
	% Detected	0	20	100

Normal serum and plasma samples were diluted 2-fold prior to the assay. ND = not detected (<LLOD); NA = not applicable due to 0% detected

Dilution Linearity

	Serum (N=5)			Plasma (N=5)			Cell Culture Media (N=5)		
	Fold Dilution	Average % Recovery	% Recovery Range	Fold Dilution	Average % Recovery	% Recovery Range	Fold Dilution	Average % Recovery	% Recovery Range
0	2	99	87-114	2	111	103-115	2	91	89-93
Cynomolgus Monkey	4	104	87-133	4	112	103-123	4	85	82-89
	8	112	90-155	8	112	100-128	8	86	78-100
Dharra	2	99	70-117	2	103	99-112	2	91	89-93
Rhesus Monkey	4	109	97-121	4	107	99-120	4	85	82-89
	8	112	98-130	8	110	100-126	8	86	78-100

Normal serum, EDTA plasma, and cell culture media were spiked with Calibrator and tested at different dilutions. Undiluted samples were tested to determine the expected concentration of the analyte. Samples may benefit from additional dilution with assay diluent to reduce matrix effects.

% Recovery = (measured concentration / expected concentration) x 100





Specificity

To assess specificity, the IFN- γ Antibody Set was tested individually against a larger panel of recombinant human analytes for nonspecific binding (CTACK, Eotaxin, Eotaxin-2, Eotaxin-3, ENA-78, FLT3L, Fractalkine, G-CSF, GM-CSF, GRO- α , I-309, IFN- α 2a, IFN- γ , IL-1 α , IL-1 β , IL-1RA, IL-2, IL-4, IL-5, IL-6, IL-7, IL-8, IL-9, IL-10, IL-12/IL-23p40, IL-12p70, IL-13, IL-15, IL-16, IL-17A, IL-17A/F, IL-17B, IL-17C, IL-17D, IL-17F, IL-18, IL-22, IL-23, IP-10, IL-TAC, MCP-1, MCP-2, MCP-3, MCP-4, M-CSF, MDC, MIF, MIP-1 α , MIP-1 β , MIP-3 α , MIP-3 β , MIP-5, SDF-1 α , TARC, TNF- α , TNF- β , TPO, TRAIL, VEGF-A, and YKL-40). Nonspecific binding was less than 0.5%.

% Nonspecificity = (nonspecific signal / specific signal) x 100

Diluent Compatibility

Diluents 57 and 3 are provided with this assay. MSD offers a range of assay and antibody diluents for separate purchase. Depending on your assay needs, other diluents may be tested.

Assay Components

Calibrator: IFN- γ is included in Calibrator 1. The full-length recombinant protein is expressed in *E. coli*. **Antibodies:** The U-PLEX NHP IFN- γ Assay uses a mouse monoclonal antibody for capture and a mouse monoclonal antibody for detection. **Assay generation:** C

Note: This datasheet contains representative assay performance data. In custom multiplex formats, the assay may perform differently than the representative data shown.

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