G-Series Human Cytokine Antibody Array 640

A combination of 16 non-overlapping arrays to measure the relative expression levels of 640 human cytokines

Catalog #: GSH-CAA-640

User Manual
Last revised October 1, 2021

Caution:
Extraordinarily useful information enclosed

RayBiotech
Empowering your proteomics

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Please read the entire manual carefully before starting your experiment

I. Overview

Cytokines Detected (640)	Arrays Included: GSH-INF-3 (40); GSH-GF-1 (40); GSH-CHE-1 (40); GSH-REC-1 (40); GSH-CYT-4 (40); GSH-CYT-5 (40); GSH-CYT-6 (40); GSH-CYT-7 (40); GSH-CYT-8 (40); GSH-CYT-9 (40); GSH-CYT-10 (40); GSH-CYT-11 (40); GSH-CYT-12 (40); GSH-CYT-13 (40); GSH-CYT-14 (40); GSH-CYT-15 (40) *See Section IX for Array Map*
Format	One standard glass slide is spotted with 16 wells of identical cytokine antibody arrays. Each antibody is arrayed in quadruplicate.
Detection Method	Fluorescence. Go to www.RayBiotech.com/Scanners for a list of compatible laser scanners.
Sample Volume	50 - 100 µl per array
Reproducibility	CV <20%
Assay Duration	6 hours

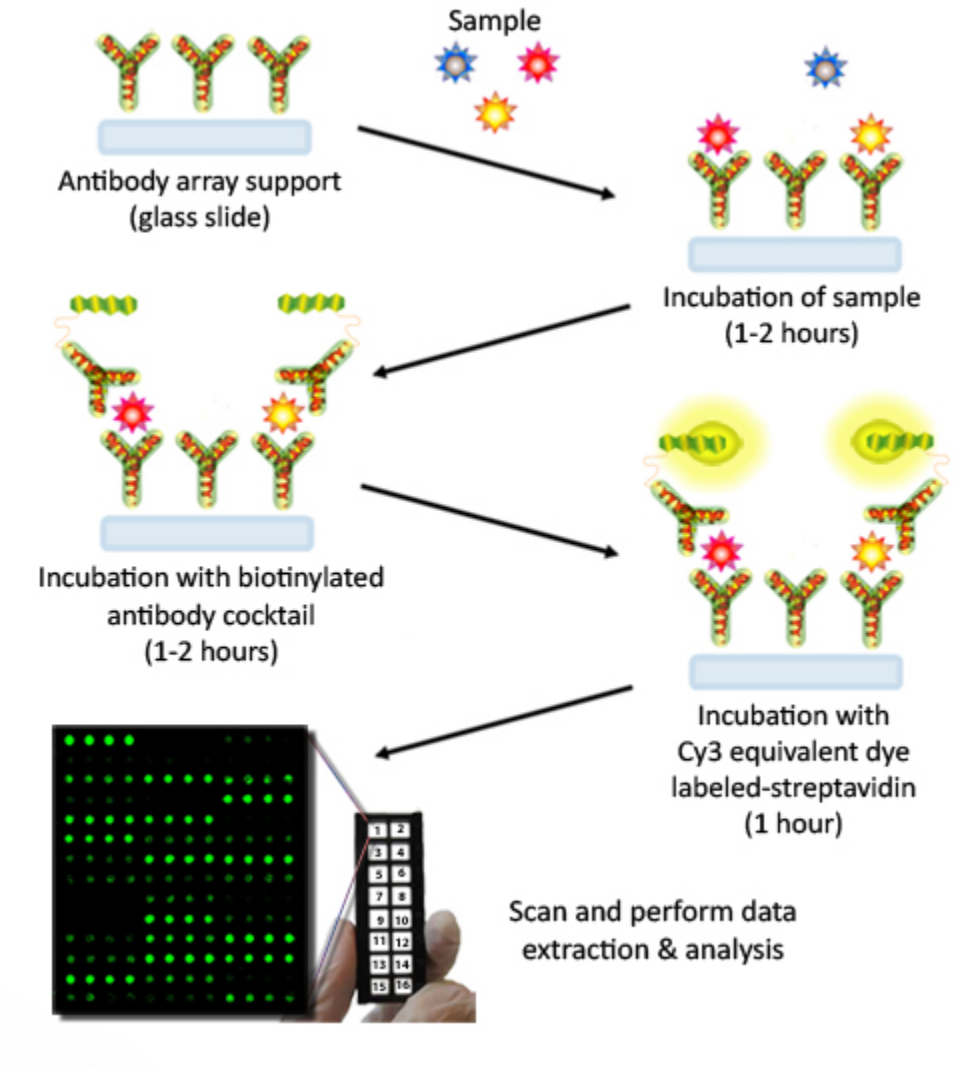
II. Introduction

Cytokines play an important role in innate immunity, apoptosis, angiogenesis, cell growth and differentiation. They are involved in interactions between different cell types, cellular responses to environmental conditions, and maintenance of homeostasis. In addition, cytokines are also involved in most disease processes, including cancer and cardiac diseases. RayBio® G-Series Arrays are glass slide-based antibody arrays which allow researchers to conduct rapid, accurate expression profiling of hundreds of cytokines, chemokines, growth factors, proteases, soluble receptors and other proteins from any biological fluid. Like a traditional sandwich-based ELISA, this array uses a matched pair of cytokine-specific antibodies for detection. After incubation with the sample, the target cytokines are captured by the antibodies printed on the solid surface. A second biotin-labeled detection antibody is then added, which recognizes a different epitope of the target cytokine. The cytokine-antibody-biotin complex can then be visualized through the addition of the streptavidin-conjugated Cy3 equivalent dye. Like the Quantibody® arrays, G-Series utilizes a highly sensitive and stable fluorescent readout which can be detected by most laser fluorescent scanner systems. After capturing the spot densities with a laser scanner,

normalization of the raw data can be easily calculated by the researcher, or by a quick copy-paste into our excel-based Analysis Tool software.

This array as well as all catalog numbers beginning with 'GS' differ from the classic G-Series Arrays in a few important ways. First, each capture antibody is printed in quadruplicate instead of duplicate, delivering higher precision. Secondly, this array features the same antibody panels used in our Quantibody Arrays, allowing a seamless transition to our quantitative multiplex assay platform. Lastly, all 16 wells are spotted as sub-arrays, delivering easy handling of 16 samples simultaneously while consuming low sample volumes (10 - 100 µl per array).

III. How It Works



IV. Materials Provided

This product is a combination of multiple arrays. Items 1, 5, & 6 are array-specific.

	Catalog #	Component Name	1 Slide Box	2 Slide Box*
1	*[Array-Cat-#]*S	*Array-specific* Glass Slide	1	2
2	QA-SDB	Sample Diluent	15 ml	
3	AA-WB1-30ML	20X Wash Buffer I	2 x 30 ml	3 x 30 ml
4	AA-WB2-30ML	20X Wash Buffer II	30 ml	
5	*[Array-Cat-#]*B	*Array-specific* Biotinylated Antibody Cocktail	1-25 µl	2 x 1-25 µl
6	QA-CY3E	Cy3 equivalent dye-conjugated Streptavidin	5 µl	2 x 5 µl
7	QA-SWD	Slide Washer/Dryer	1 x 30 ml Tube	
8	QA-ADH	Adhesive Film	1	2

* 4 slide kits are comprised of 2 separate 2 slide kits.

V. Storage

Upon receipt, all components should be stored at -20°C. The kit will retain activity for up to 6 months. Once thawed, the glass slide, antibody cocktail and dye-conjugated Streptavidin should be kept at -20°C. All other components may be stored at 4°C. The entire kit should be used within 6 months of purchase.

VI. Additional Materials Required

- Benchtop rocker or orbital rocker
- Laser scanner for fluorescence detection
- Aluminum foil
- Distilled water
- 1.5 ml Polypropylene microcentrifuge tubes

VII. General Considerations

A. Preparation of Samples

- Use serum-free conditioned media if possible.
- If serum-containing conditioned media is required, it is highly recommended that complete medium be used as a control since many types of sera contains cytokines.
- Each array needs 100 µl of total sample volume. To avoid matrix effects, we recommend using a minimum of 2x dilution for serum, plasma, cell culture media, or other body fluids, or 500 µg/ml-1 mg/ml (after a 5-fold to 10-fold dilution to minimize the effects of any detergent(s)) total protein for cell and tissue lysates. Please be aware, more sample volume is required for combination arrays. For example, the minimum sample volume for a 10-array kit is 500 µl, or 500 µg cell lysate.

  If you experience high background or if the fluorescent signal intensities exceed the detection range, further dilution of your sample is recommended.

B. Handling Glass Slides

- Do not touch the surface of the slides, as the microarray slides are very sensitive. Hold the slides by the edges only.
- Handle all buffers and slides with powder free gloves.
- Handle glass slide/s in clean environment.
- Permanent marker ink can significantly interfere with fluorescent signal detection. To help distinguish one slide from another, you may make a small marking (such as a number or a star) along the top or bottom edge, using a green or blue ultra-fine point Sharpie® brand marker. This can also serve to orient the slide. For best results during scanning, please **DO NOT**:
  - Write anywhere on the front (arrayed) side of the slide
  - Write on the slide while it is wet
  - Use red or black colored ink anywhere on the slide
  - Write over the arrayed well areas of the slide, as this interferes with scanning.

C. Incubation

- Completely cover array area with sample or buffer during incubation.
- Avoid foaming during incubation steps.

- Perform all incubation and wash steps under gentle rocking or rotation.
- Cover the incubation chamber with adhesive film during incubation, particularly when incubation is more than 2 hours or <70 µl of sample or reagent is used.
- Several incubation steps such as step 6 (blocking), step 7 (sample incubation), step 10 (detection antibody incubation), or step 13 (Cy3 equivalent dye-streptavidin incubation) may be done overnight at 4°C. Please make sure to cover the incubation chamber tightly to prevent evaporation.

VIII. Protocol

Note: *This product contains sets of reagents for different arrays. Always ensure you are using the proper glass slide and biotinylated antibody cocktail for the correct corresponding array.*
The following procedure is for processing any one of the arrays in the kit.

A. Completely Air Dry The Glass Slide

1. Take out the glass slide from the box, and let it equilibrate to room temperature inside the sealed plastic bag for 20-30 minutes. Remove slide from the plastic bag, peel off the cover film, and let it air dry for another 1-2 hours.

Incomplete drying of slides before use may cause the formation of "comet tails," thin directional smearing of antibody spots.

B. Blocking & Incubation

2. Add 100 µl Sample Diluent into each well and incubate at room temperature for 30 minutes to block slides.

3. Decant buffer from each well. Add 100 µl of sample to each well. Incubate arrays at room temperature for 1-2 hour.

Longer incubation time is preferable for higher signals. This step may be done overnight at 4°C.

We recommend using 50 to 100 µl of original or diluted serum, plasma, conditioned media, or other body fluid, or 50-500 µg/ml of protein for cell and tissue lysates. Cover the incubation chamber with adhesive film during incubation, especially if less than 70 ul of sample or reagent is used.

4. Wash:

- Decant the samples from each well, and wash 5 times (5 min each) with 150 µl of 1X Wash Buffer I at room temperature with gentle shaking. Completely remove wash buffer in each wash step. Dilute 20x Wash Buffer I with H2O.

- *(Optional for Cell and Tissue Lysates)* Put the glass slide with frame into a box with 1X Wash Buffer I (cover the whole glass slide and frame with Wash Buffer I), and wash at room temperature with gentle shaking for 20 min.

- Decant the 1x Wash Buffer I from each well, wash 2 times (5 min each) with 150 µl of 1X Wash Buffer II at room temperature with gentle shaking. Completely remove wash buffer in each wash step. Dilute 20X Wash Buffer II with H2O.

Incomplete removal of the wash buffer in each wash step may cause "dark spots," the background signals higher than the spots.

C. Incubation with Biotinylated Antibody Cocktail & Wash

5. Reconstitute the detection antibody by adding 1.4 ml of Sample Diluent to the tube. Spin briefly.

6. Add 80 µl of the detection antibody cocktail to each well. Incubate at room temperature for 1-2 hour.

Longer incubation time is preferable for higher signals

7. Decant the samples from each well, and wash 5 times (5 mins each) with 150 µl of 1X Wash Buffer I and then 2 times with 150 µl of 1x Wash Buffer II at room temperature with gentle shaking. Completely remove wash buffer in each wash step.

D. Incubation with Cy3 Equivalent Dye-Streptavidin & Wash

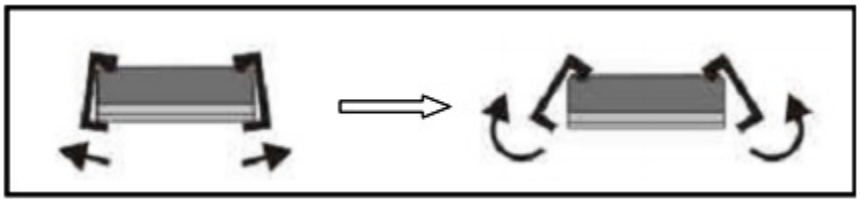
8. After briefly spinning down, add 1.4 ml of Sample Diluent to Cy3 equivalent dye-conjugated streptavidin tube. Mix gently.

9. Add 80 µl of Cy3 equivalent dye-conjugated streptavidin to each well. Cover the device with aluminum foil to avoid exposure to light or incubate in dark room. Incubate at room temperature for 1 hour.

10. Decant the samples from each well, and wash 5 times (5 mins each) with 150 µl of 1X Wash Buffer I at room temperature with gentle shaking. Completely remove wash buffer in each wash step.

E. Fluorescence Detection

11. Disassemble the device by pushing clips outward from the slide side. Carefully remove the slide from the gasket.



Be careful not to touch the surface of the array side.

12. Place the slide in the Slide Washer/Dryer (a 4-slide holder/centrifuge tube), add enough 1x Wash Buffer I (about 30 ml) to cover the whole slide, and then gently shake at room temperature for 15 minutes. Decant Wash Buffer I. Wash with 1x Wash Buffer II (about 30 ml) and gently shake at room temperature for 5 minutes.

13. Remove water droplets completely by gently applying suction with a pipette to remove water droplets. Do not touch the array, only the sides.

    You may also dry the glass slide by a compressed N_2 stream.

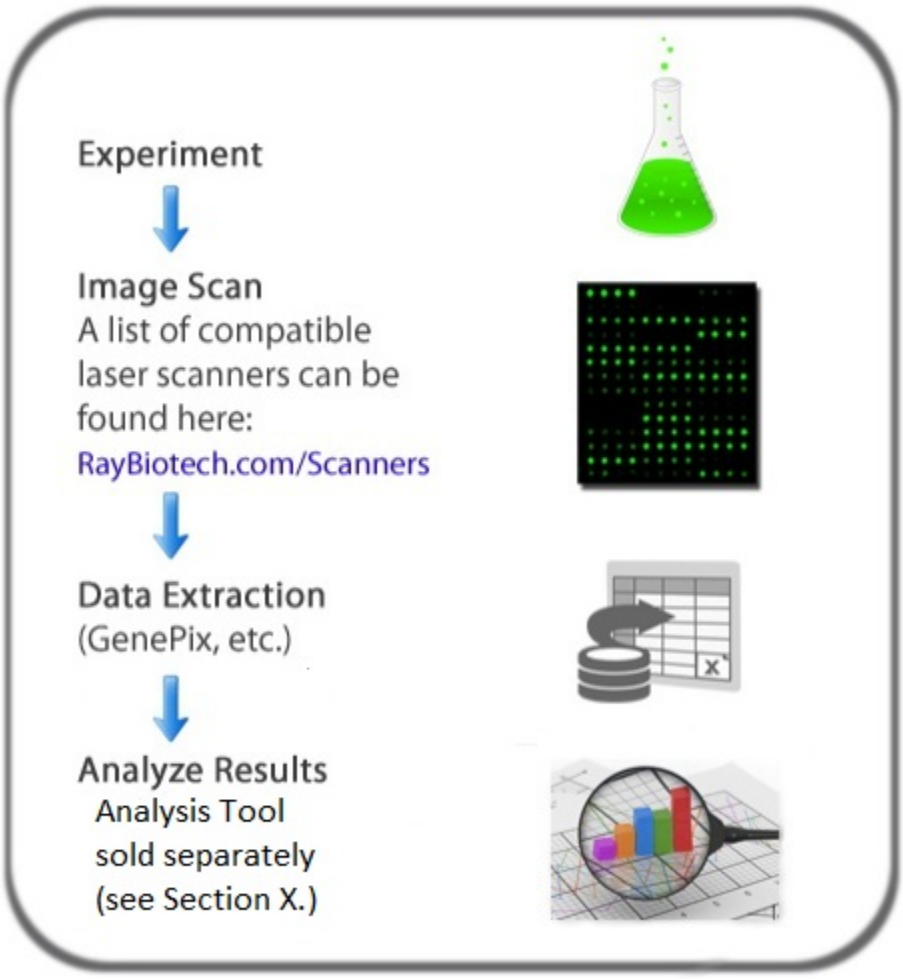
14. Imaging: The signals can be visualized through use of a laser scanner equipped with a Cy3 wavelength (green channel) such as Axon GenePix or Innopsys Innoscan.

In case the signal intensity for different cytokine varies greatly in the same array, we recommend using multiple scans, with a higher PMT for low signal cytokines, and a low PMT for high signal cytokines.

F. Data Analysis

15. >Data extraction can be done using the GAL file that is specific for this array (QAH-CAA-640) along with the microarray analysis software (GenePix, ScanArray Express, ArrayVision, MicroVigene, etc.). The GAL file can be found on the product web page under the 'Files' tab.

    Need help analyzing all that data? All RayBiotech array analysis tools are now free to download! Just like the GAL file, you can find this analysis tool on the product web page under the 'Files' tab. More information can be found in Section X.



IX. Array Map

QAH-INF-3

(hINF-3 Map) Each antibody is printed in quadruplicate horizontally

	1	2	3	4	1	2	3	4	1	2	3	4
A	POS1				POS2				BLC (CXCL13)			
B	Eotaxin-1 (CCL11)				Eotaxin-2 (MPIF-2)				G-CSF			
C	GM-CSF				I-309				ICAM-1 (CD54)			
D	IFN gamma				IL-1 alpha				IL-1 beta			
E	IL-1ra (IL-1 F3)				IL-2				IL-4			
F	IL-5				IL-6				IL-6sR			
G	IL-7				IL-8				IL-10			
H	IL-11				IL-12p40				IL-12p70			
I	IL-13				IL-15				IL-16			
J	IL-17				MCP-1 (CCL2)				M-CSF			
K	MIG (CXCL9)				MIP-1 alpha (CCL3)				MIP-1 beta (CCL4)			
L	MIP-1 delta (CCL15)				PDGF-BB				RANTES (CCL5)			
M	TIMP-1				TIMP-2				TNF alpha			
N	TNF beta				TNF RI				TNF RII			

QAH-GF-1

(hGF-1 Map) Each antibody is printed in quadruplicate horizontally

	1	2	3	4	1	2	3	4	1	2	3	4
A	POS1				POS2				AR			
B	BDNF				bFGF				BMP-4			
C	BMP-5				BMP-7				beta-NGF			
D	EGF				EGF R				EG-VEGF			
E	FGF-4				FGF-7 (KGF)				GDF-15			
F	GDNF				Growth Hormone (GH)				HB-EGF			
G	HGF				IGFBP-1				IGFBP-2			
H	IGFBP-3				IGFBP-4				IGFBP-6			
I	IGF-I				Insulin				MCF R			
J	NGFR (TNFSR16)				NT-3				NT-4			
K	Osteoprotegerin (OPG)				PDGF-AA				PlGF (PLGF)			
L	SCF				SCF R (CD117)				TGF alpha			
M	TGF beta 1				TGF beta 3				VEGF-A (VEGF)			
N	VEGF R2				VEGF R3				VEGF-D			

QAH-CHE-1

(hCHE-1 Map) Each antibody is printed in quadruplicate horizontally

	1	2	3	4	1	2	3	4	1	2	3	4
A	POS1				POS2				6Ckine (CCL21)			
B	Axl				Betacellulin (BTC)				CCL28 (MEC)			
C	CTACK (CCL27)				CXCL16				ENA-78 (CXCL5)			
D	Eotaxin-3 (CCL26)				GCP-2 (CXCL6)				GRO			
E	HCC-1 (CCL14)				HCC-4 (CCL16)				IL-9			
F	IL-17F				IL-18 BP alpha				IL-28A			
G	IL-29				IL-31				IP-10 (CXCL10)			
H	I-TAC (CXCL11)				LIF				LIGHT (TNFSF14)			
I	Lymphotactin				MCP-2 (CCL8)				MCP-3 (CCL7)			
J	MCP-4 (CCL13)				MDC (CCL22)				MIF			
K	MIP-3 alpha				MIP-3 beta				MPIF-1 (CCL23)			
L	MSP				NAP-2 (CXCL7)				Osteopontin (OPN)			
M	PARC (CCL18)				Platelet Factor 4 (PF4)				SDF-1 alpha			
N	TARC (CCL17)				TECK (CCL25)				TSLP			

QAH-REC-1

(hREC-1 Map) Each antibody is printed in quadruplicate horizontally

	1	2	3	4	1	2	3	4	1	2	3	4
A	POS1				POS2				4-1BB (CD137)			
B	ALCAM (CD166)				B7-1 (CD80)				BCMA (TNFRSF17)			
C	CD14				CD30 (TNFRSF8)				CD40 Ligand			
D	CEACAM-1				DR6 (TNFRSF21)				Dtk			
E	Endoglin (CD105)				ErbB3				E-Selectin			
F	Fas				Flt-3 Ligand				GITR (TNFRSF18)			
G	HVEM (TNFRSF14)				ICAM-3 (CD50)				Contactin-2			
H	IL-1 RI				IL-2 R gamma				IL-10 R beta			
I	IL-17R				IL-21 R				LIMPII			
J	Lipocalin-2 (NGAL)				L-Selectin (CD62L)				LYVE-1			
K	MICA				MICB				NRG1-beta 1			
L	PDGF R beta				PECAM-1 (CD31)				RAGE			
M	TIM-1 (KIM-1)				TRAIL R3				Trappin-2			
N	uPAR				VCAM-1				XEDAR			

QAH-CYT-4

(hCYT-4 Map) Each antibody is printed in quadruplicate horizontally

	1	2	3	4	1	2	3	4	1	2	3	4
A	POS1				POS2				Activin A			
B	AgRP				Angiogenin (ANG)				Angiopoietin-1 (ANG-1)			
C	Angiostatin				Catheprin S				CD40			
D	Cripto-1				DAN				DKK-1			
E	E-Cadherin				EpCAM (TROP1)				Fas Ligand (TNFSF6)			
F	Fc gamma RIIB/C				Follistatin				Galectin-7			
G	ICAM-2 (CD102)				IL-13 R1				IL-13 R alpha 2			
H	IL-17B				IL-2 R alpha				IL-2 R beta			
I	IL-23				LAP/TGF beta 1				NrCAM			
J	PAI-I				PDGF-AB				Resistin			
K	SDF-1 beta				sgp130				Shh N			
L	Siglec-5 (CD170)				ST2 (IL-1 R4)				TGF-beta 2			
M	Tie-2				Thrombopoietin (TPO)				TRAIL-R4			
N	TREM-1				VEGF-C				VEGF-R1			

QAH-CYT-5

(hCYT-5 Map) Each antibody is printed in quadruplicate horizontally

	1	2	3	4	1	2	3	4	1	2	3	4
A	POS1				POS2				Adiponectin (ACRP30)			
B	Adipsin				Alpha-fetoprotein (AFP)				ANGPTL4			
C	Beta-2 Microglobuliin (B2M				BCAM				CA125			
D	CA15-3				CEA				CRP			
E	ErbB2				Ferritin				FSH			
F	GRO alpha (CXCL1)				HCG beta (HCGb)				IGF-I R			
G	IL-1 RII				IL-3				IL-18 R beta			
H	IL-21				Leptin				MMP-1			
I	MMP-2				MMP-3				MMP-8			
J	MMP-9				MMP-10				MMP-13			
K	NCAM-1 (CD56)				Nidogen-1				NSE			
L	Oncostatin M (OSM)				Procalcitonin (PCT)				Prolactin			
M	PSA-free				Siglec-9				TACE			
N	Thyroglobulin				TIMP-4				TSH			

QAH-CYT-6

(hCYT-6 Map) Each antibody is printed in quadruplicate horizontally

	1	2	3	4	1	2	3	4	1	2	3	4
A	POS1				POS2				2B4 (CD244)			
B	ADAM-9				Angiopoietin-2 (ANG-2)				APRIL			
C	BMP-2				BMP-9				C5a			
D	Cathepsin L				CD200				CD97			
E	Chemerin				DcR3				FABP2			
F	FAP				FGF-19				Galectin-3			
G	HGF R				IFN alpha/beta R2				IGF-II			
H	IGF-II R				IL-1 R6 (IL-1 Rrp2)				IL-24			
I	IL-33 (IL-1 F11)				Kallikrein 14				Legumain			
J	LOX-1				MBL				Neprilysin			
K	Notch-1				NOV (CCN3)				Osteoactivin			
L	PD-1				PGRPs				Serpin A4			
M	sFRP-3				Thrombomodulin				TLR2			
N	TRAIL R1				Transferrin				WIF-1			

QAH-CYT-7

(hCYT-7 Map) Each antibody is printed in quadruplicate horizontally

	1	2	3	4	1	2	3	4	1	2	3	4
A	POS1				POS2				ACE-2			
B	Albumin				AMICA				Angiopoietin-4 (ANG-4)			
C	BAFF				CA19-9				CD163			
D	Clusterin				CRTAM				CXCL14 (BRAK)			
E	Cystatin C				Decorin				Dkk-3			
F	DLL1				Fetuin A				aFGF (FGF-1)			
G	FOLR1				Furin				GASP-1			
H	GASP-2				G-CSF R (CD114)				HAI-2			
I	IL-17B R (IL-17 RB)				IL-27				LAG-3			
J	LDL R				Pepsinogen I (PG1)				RANK			
K	RBP4				SOST				Syndecan-1			
L	TACI				TFPI				Thrombospondin 1			
M	TRAIL R2				TRANCE				Troponin I			
N	uPA				VE-Cadherin (CDH5)				WISP-1 (CCN4)			

QAH-CYT-8

(hCYT-8 Map) Each antibody is printed in quadruplicate horizontally

	1	2	3	4	1	2	3	4	1	2	3	4
A	POS1				POS2				ANGPTL3			
B	beta IG-H3				CA9				Cathepsin B			
C	CD23				CHI3L1				CTLA4			
D	Dkk-4				DPPIV				EDA-A2			
E	Epo R				FGF-6				FGF-9			
F	Gas1				IGFBP-5				IL-1F5			
G	IL-1F6				IL-1F7				IL-1F8			
H	IL-1F9				IL-1F10				IL-1R5			
I	IL-17C				IL-18				IL-20			
J	IL-34				IL-5 R alpha				IL-10 R alpha			
K	Layilin				Leptin R				Marapsin			
L	Mer				MMP-7				P-Cadherin			
M	Prostasin				PSMA				SIGIRR			
N	TGF beta RIII				Tissue Factor (TF)				TWEAK			

QAH-CYT-9

(hCYT-9 Map) Each antibody is printed in quadruplicate horizontally

	1	2	3	4	1	2	3	4	1	2	3	4
A	POS1				POS2				ADAMTS13			
B	Aggrecan				Angiotensinogen (AGT)				B7-H1 (CD274)			
C	BMPR-IA (ALK-3)				BMPR-II				Cadherin-11			
D	CD27 (TNFRSF7)				CD6				Ck beta 8-1 (CCL23)			
E	CNTF				DNAM-1 (CD226)				EMMPRIN (CD147)			
F	FLRG				Follistatin-like 1 (FSL1)				Fractalkine (CX3CL1)			
G	Galectin-1				GITR Ligand				Granulysin (LAG-2)			
H	IL-1 R3 (IL-1 R Acp)				IL-15 R alpha				IL-17E (IL-25)			
I	IL-32 alpha				L1CAM-2 (CHL-1)				LRIG3			
J	LRP-6				MEPE (OF45)				Nectin-4			
K	Periostin				Persephin				Renin			
L	RGM-B				ROBO3				S100A8			
M	Siglec-7 (CD328)				Syndecan-3				Thrombospondin 2			
N	Thrombospondin 5				Tie-1				ULBP-2			

QAH-CYT-10

(hCYT-10 Map) Each antibody is printed in quadruplicate horizontally

	1	2	3	4	1	2	3	4	1	2	3	4
A	POS1				POS2				ADAM8			
B	ADAM12				B7-H3 (CD276)				BMPR-IB			
C	Cadherin-4				Cadherin-13				CD48 (SLAMF2)			
D	CD58 (LFA-3)				CD84 (SLAMF5)				CD99			
E	CD155 (PVR)				CD229 (SLAM3)				CEACAM-5			
F	CF XIV				Cystatin A				Cystatin B			
G	Cystatin E/M				Desmoglein 2				DR3 (TNFRSF25)			
H	ErbB4 (HER4)				ESAM				FGF-21			
I	Galectin-2				Galectin-9				ICOS			
J	JAM-A (CD321)				JAM-B (CD322)				Kallikrein 5			
K	Midkine				Pentraxin 3				Pref-1 (DLK-1)			
L	Siglec-10				SLAM (CD150)				SP-D			
M	Syndecan-4				Testican 2 (SPOCK2)				TIM-3 (KIM-3)			
N	TLR4				TRAIL (TNFSF10)				ULBP-1			

QAH-CYT-11

(hCYT-11 Map) Each antibody is printed in quadruplicate horizontally

	1	2	3	4	1	2	3	4	1	2	3	4
A	POS1				POS2				ALK-1			
B	B7-H2				BLAME				BMP-8			
C	CD28				Common beta Chain				Contactin-1			
D	Desmoglein-1				Desmoglein-3				EDAR			
E	EphA1				EphB6				Ephrin-B3			
F	Epiregulin				FGF-12				FGF-17			
G	FOLR2				Galectin-8				GHR			
H	Glypican 1				Glypican 5				IFN-gamma R1			
I	IL-22 R alpha 1				IL-22BP				IL-23 R			
J	IL-31 RA				IL-7 R alpha				Integrin alpha 5			
K	MDM2				Nectin-1				NKp30			
L	Nogo Receptor				Notch-3				OSM R beta			
M	Prolactin R				RELT				Ryk			
N	Semaphorin 6D				Semaphorin 7A				Siglec-11			

QAH-CYT-12

(hCYT-12 Map) Each antibody is printed in quadruplicate horizontally

	1	2	3	4	1	2	3	4	1	2	3	4
A	POS1				POS2				B7-2			
B	BAFF R				Calcitonin				Calsyntenin-1			
C	Cathepsin E				cIAP-2				Coagulation Factor VII			
D	Complement MASP3				Endocan				EphA2			
E	EphB4				Ephrin-A4				FGF-23			
F	FGF-5				Flt-3				GLP-1			
G	Glypican 2				GM-CSF R alpha				GP73			
H	HTRA2				IL-20 R alpha				IL-4 R alpha			
I	JAM-C				Luteinizing hormone (LH)				Matrilin-3			
J	Meprin alpha				MSP R				N-Cadherin			
K	Neprilysin-2				NKp44				PAPP-A			
L	Pepsinogen II				Presenilin 1				PTH			
M	PYY				SOX2				TFF3			
N	TFPI-2				TRACP				Ubiquitin+1			

QAH-CYT-13

(hCYT-13 Map) Each antibody is printed in quadruplicate horizontally

	1	2	3	4	1	2	3	4	1	2	3	4
A	POS1				POS2				ACE			
B	Activin RIB				ADAM23				Artemin			
C	Cardiotrophin-1				Cathepsin V				FABP1			
D	FGF-20				GDF-8				HAI-1			
E	IL-27 R alpha				Insulin R				Kallikrein 7			
F	LIF R alpha				Lipocalin-1				LTbR			
G	Mesothelin				MFRP				Neuropilin-2			
H	Neurturin				Nidogen-2				Olfactomedin-2			
I	p53				PD-ECGF				PDGF-CC			
J	Progranulin				Ret				ROBO4			
K	Semaphorin 6B				Serpin F1				SREC-I			
L	SREC-II				TLR1				TLR3			
M	TPP1				TREM-2				TrkC			
N	TROY				Uromodulin				XIAP			

QAH-CYT-14

(hCYT-14 Map) Each antibody is printed in quadruplicate horizontally												
	1	2	3	4	1	2	3	4	1	2	3	4
A	POS1				POS2				4-1BB Ligand			
B	Activin RIIB				Aminopeptidase P2				BAMBI			
C	BOC				Brevican				Carbonic Anhydrase XII			
D	Carboxypeptidase A2				CD300c				CD320			
E	CDNF				CDO				CHST1			
F	CHST4				CILP-1				CNTF R alpha			
G	CRIM1				CRTAC1				CXADR			
H	Dopa Decarboxylase				DPPII				DSPG3			
I	EMR2				FCAR				FCRL1			
J	FCRL2				Gas6				GPR56			
K	GPVI				Hepsin				ILT2			
L	Jagged 2				Kirrel3				KLF4			
M	LAIR1				LAMP				LAMP1			
N	MDGA1				MIS RII				Neurexin 3 beta			

QAH-CYT-15

(hCYT-15 Map) Each antibody is printed in quadruplicate horizontally												
	1	2	3	4	1	2	3	4	1	2	3	4
A	POS1				POS2				AMIGO			
B	Aminopeptidase LRAP				Amnionless				Arylsulfatase A			
C	Bcl-w				CD109				CD157			
D	CD34				CD83				CLEC-1			
E	CLEC10A				CMG-2				CREG			
F	Cystatin SN				Cytokeratin-8				Dectin-1			
G	Desmocollin-3				Endoglycan				Galectin-4			
H	HAPLN1				Jagged 1				Langerin			
I	Lumican				Matriptase				MEP1B			
J	Nectin-3				OX40				OX40 Ligand			
K	p27				Pappalysin-2				Plexin B3			
L	Plexin D1				proGRP				PSA-total			
M	Reg1B				RGM-A				ROBO2			
N	Spinesin				TWEAK R				ULBP-3			

X. Array Data Analysis Tool

The RayBio Analysis Tools are array specific, Excel-based program that perform sophisticated data analysis on the raw numerical data extracted from the array scan. All RayBiotech array analysis tools are now free to download! Just like the GAL file, you can find this analysis tool on the product web page under the 'Files' tab.

Key features:

- Simplicity: Easy to operate and requires no professional training. With a simple copy and paste process, the cytokine expression levels are determined per sample.
- Outlier Marking & Removing: The software can automatically mark and remove the outlier spots for more accurate data analysis
- Normalization: The program allows for intra- and inter-slide normalization for large numbers of samples.
- Two Positive Controls: The program utilizes the two positive controls in each array for normalization.
- User Intervention: The program allows for user manual handling of outliers and other analytical data.
- Analyze Multiple Slide: The data for multiple slides can be inputted for easy slide-to-slide comparison.

XI. Troubleshooting Guide

Problem	Cause	Recommendation
Weak Signal	Inadequate detection	Increase laser power and PMT parameters
	Inadequate reagent volumes or improper dilution	Check pipettes and ensure correct preparation
	Short incubation time	Increase incubation time or change sample incubation step to overnight
	Too low protein concentration in sample	Lessen dilution or do not dilute sample. Concentrate sample if necessary.
	Improper storage of kit	Store kit as suggested temperature. Don't freeze/thaw the slide.
Uneven signal	Bubble formed during incubation	Decrease amount of rocking/shaking during incubations. check for bubble formation and remove bubbles.
	Arrays are not completed covered by reagent	Completely cover arrays with solution for all required steps.
	Reagent evaporation	Cover the incubation chamber with adhesive film during incubation
High background	Overexposure	Lower the PMT or sigmal gain.
	Dark spots	Completely remove wash buffer in each wash step.
	Insufficient wash	Increase wash time and use more wash buffer
	Dust	Work in clean environment
	Slide is allowed to dry out	Don't dry out slides during experiment.

XII. Select Publications

1. Stechova, et al. Influence of Maternal Hyperglycaemia on Cord Blood Mononuclear Cells in Response to Diabetes-associated Autoantigens. Scandinavian Journal of Immunology. 2009. 70(2):149-158
2. Willingham, SB et al. NLRP3 (NALP3, Cryopyrin) facilitates in vivo caspase-1 activation, necrosis, and HMGB1 release via inflammasome-dependent and -independent pathways. J Immunol. 2009; 183(3):2008-15
3. El Karim et al. Neuropeptides Regulate Expression of Angiogenic Growth Factors in Human Dental Pulp Fibroblasts. Journal of Endodontics, 2009; 35(6): 829-833
4. Souquière S. et al. T-Cell tropism of simian T-cell leukaemia virus type 1 and cytokine profiles in relation to proviral load and immunological changes during chronic infection of naturally infected mandrills (Mandrillus sphinx). J Med Primatol. 2009; 38(4):279-89
5. Sharma, et al. Induction of multiple pro-inflammatory cytokines by respiratory viruses and reversal by standardized Echinacea, a potent antiviral herbal extract. Antiviral Research. 2009; 83(2)165-170.
6. Altamirano-Dimas, et al. Echinacea and anti-inflammatory cytokine responses: Results of a gene and protein array analysis. Pharmacuetical Biology. 2009; 47(6): 500-508.
7. Cheung, et al. Cordysinocan, a polysaccharide isolated from cultured Cordyceps, activates immune responses in cultured T-lymphocytes and macrophages: Signaling cascade and induction of cytokines. Journal of Ethonopharmacology. 2009; 124(1): 61-68.
8. Du, et al. P2-380: Identification and characterization of human autoantibodies that may be used for the treatment of prion diseases. Alzheimers and Dementia. 2009; 4(4): T484-T484.
9. Van Rossum et al. Granulocytosis and thrombocytosis in renal cell carcinoma: a pro-inflammatory cytokine response originating in the tumour. Neth J Med. 2009; 67(5):191-4.
10. Zhai, et al. Coordinated Changes in mRNA Turnover, Translation, and RNA Processing Bodies in Bronchial Epithelial Cells following Inflammatory Stimulation. Molecular and Cellular Biology. 2008; 28(24): 7414-7426.
11. Gao, et al. A Chinese herbal decoction, Danggui Buxue Tang, activates extracellular signal-regulated kinase in cultured T-lymphocytes. FEBS Letters, 2007; 581(26): 5087-5093.*This reference validates mulitplex ELISA results for several analytes with standard ELISA test results*.
12. Piganelli, et al: Autoreactive T-cell responses: new technology in pursuit of an old nemesis. (*Editorial Review*) Pediatric Diabetes 2007: 8: 249–251

Note: The citations listed above are for the Quantibody® product line, which is the same as the GS-Series, but include protein standards for quantitation.

XIII. Experiment Record Form

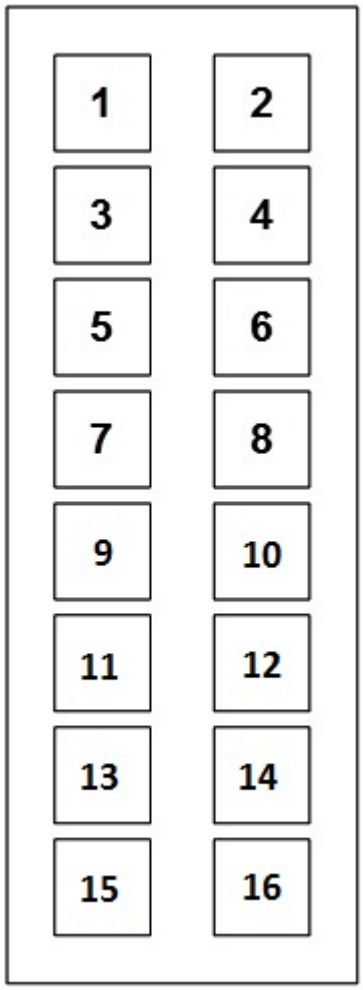
Date:_____

File Name:_____

Laser Power:_____

PMT:_____

Well No.	Sample Name	Dilution factor
1		
2		
3		
4		
5		
6		
7		
8		
9		
10		
11		
12		
13		
14		
15		
16		



XIV. How to Choose a GS-Series Array?

Species-based selection:

Human (GSH-)	Mouse (GSM-)	Rat (GSR-)	Bovine (GSB-)	Canine (GSC-)
Equine (GSE-)	Feline (GSF-)	Ovine (GSO-)	Primates (GSN-)	Porcine (GSP-)
Rabbit (GSL-)				

Function-based selection:

Adhesion Molecule Arrays	Angiogenesis Arrays	Bone Metabolism Arrays	Chemokine Arrays
Cancer Biomarker Arrays	***Custom Arrays***	Cytokine Arrays	Growth Factor Arrays
IGF Signaling Arrays	IL-1 Family Arrays	Immune Response Arrays	Inflammation Arrays
Interleukin Arrays	Isotyping Arrays	MMP Arrays	Obesity Arrays
Ophthalmic Arrays	Periodontal Disease Arrays	Receptor Arrays	Th1/Th2/Th17 Arrays

Cytokine Number-based selection:
Arrays are available in the GS-Series & Quantibody® platform to detect 660 human, 200 mouse, or 67 rat proteins. GLP-Compliant testing services are also available.

This product is for research use only.