

# RayBio® Human Cytokine Antibody Array

Patent Pending Technology

## User Manual (Revised December 21, 2009)

- RayBio® Human Cytokine Antibody Array 1 (Cat# AAH-CYT-1)
- RayBio® Human Cytokine Antibody Array 3 (Cat# AAH-CYT-3)
- RayBio® Human Cytokine Antibody Array 4 (Cat# AAH-CYT-4)
- RayBio® Human Cytokine Antibody Array 5 (Cat# AAH-CYT-5)
- RayBio® Human Cytokine Antibody Array 6 (Cat# AAH-CYT-6)
- RayBio® Human Cytokine Antibody Array 7 (Cat# AAH-CYT-7)
- RayBio® Human Cytokine Antibody Array 8 (Cat# AAH-CYT-8)
- RayBio® Human Cytokine Antibody Array 9 (Cat# AAH-CYT-9)
- RayBio® Human Cytokine Antibody Array 10 (Cat# AAH-CYT-10)
- RayBio® Human Inflammation Antibody Array 1 (Cat# AAH-INF-1)
- RayBio® Human Inflammation Antibody Array 2 (Cat# AAH-INF-2)
- RayBio® Human Inflammation Antibody Array 3 (Cat# AAH-INF-3)
- RayBio® Human Angiogenesis Antibody Array 1 (Cat# AAH-ANG-1)
- RayBio® Human Atherosclerosis Antibody Array 1 (Cat# AAH-ATH-1)
- RayBio® Human Matrix Metalloproteinase Antibody Array 1 (Cat# AAH-MMP-1)
  - RayBio® Human Chemokine Antibody Array 1 (Cat# AAH-CHE-1)
  - RayBio® Human Growth Factor Antibody Array 1 (Cat# AAH-GF-1)
- RayBio® Custom Human Cytokine Antibody Array (Cat# AAH-CUST)
- RayBio® Human Cytokine Antibody Array Service (Cat# AAH-SERV)

***Please read manual carefully before starting experiment***

---

# RayBio® Human Cytokine Antibody Array Protocol

## **TABLE OF CONTENTS**

I.	Introduction.....	2
	How It Works.....	5
II.	Materials Provided.....	6
	Additional Materials Required.....	6
III.	Overview and General Considerations.....	7
	A. Preparation of Samples.....	7
	B. Handling Array Membrane.....	7
	C. Incubation.....	8
IV.	Protocol.....	8
	A. Blocking and Incubation.....	8
	B. Detection.....	10
V.	Interpretation of Results.....	11
VI.	Troubleshooting Guide.....	21
VII.	Selected References Using RayBiotech Products...	22

Cytokine protein arrays are RayBiotech patent-pending technology.

RayBio® is the trademark of RayBiotech, Inc.

## I. Introduction

All cell functions, including cell proliferation, cell death and differentiation, as well as maintenance of health status and development of disease, are controlled by a multitude of genes and signaling pathways. New techniques such as cDNA microarrays have enabled us to analyze global gene expression<sup>1-3</sup>. However, almost all cell functions are executed by proteins, which cannot be studied simply through DNA and RNA techniques. Experimental analysis clearly shows a disparity between the relative expression levels of mRNA and their corresponding proteins<sup>4</sup>. Therefore, analysis of the protein profile is critical. Currently, two-dimensional polyacrylamide SDS page coupled with mass spectrometry is the mainstream approach to analyzing multiple protein expression levels<sup>5,6</sup>. However, the requirement of sophisticated devices and the lack of quantitative measurements greatly limit their broad application. Thus, effective study of multiple protein expression levels has been complicated, costly and time-consuming until now.

Our RayBio® Human Cytokine Antibody Arrays are the first commercially available protein array system<sup>7-11</sup>. By using the RayBiotech system, scientists can rapidly and accurately identify the expression profiles of multiple cytokines in several hours inexpensively.

The RayBiotech kit provides a simple array format, and highly sensitive approach to simultaneously detect multiple cytokine expression levels from conditioned media, patient's sera, cell lysate, tissue lysates and other sources.

Traditionally, cytokines are detected by using ELISA. However, RayBiotech's approach has several advantages over ELISA. First, and most importantly, our approach can detect many cytokines simultaneously. Secondly, sensitivity is greatly increased. As little as 4 pg/ml of MCP-1 can be detected using the protein array format. In contrast, at least 40 pg/ml of MCP-1 is required to produce a clear signal in an ELISA assay. Furthermore, the detection range is much greater than ELISA. For example, the detection range of IL-2 varies from 25 to 250,000 pg/ml using RayBiotech technology, whereas the detection range varies only within 100-1000 fold in a typical

ELISA. Therefore, the detection range is greater with protein array compared with ELISA. The variation is lower than ELISA as well. As determined by densitometry, the variation between two spots ranged from 0 to 10% in duplicated experiments. In contrast, variation (about 20%) in ELISA is much higher. Finally, the system is much quicker and can be much easier to adapt to high-throughput technique.

Pathway-specific array systems allow investigators to focus on the specific problem and are becoming an increasingly powerful tool in cDNA microarray systems. RayBiotech's first protein array system, known as RayBio® Human Angiogenesis antibody array, is particularly useful in comparison with the human cytokine cDNA microarray system. Besides the ability to detect protein expression, RayBiotech's system is a more accurate reflection of active cytokine levels because it only detects secreted cytokines, and no amplification step is needed. Furthermore, it is much simpler, faster, environmentally friendlier, and more sensitive.

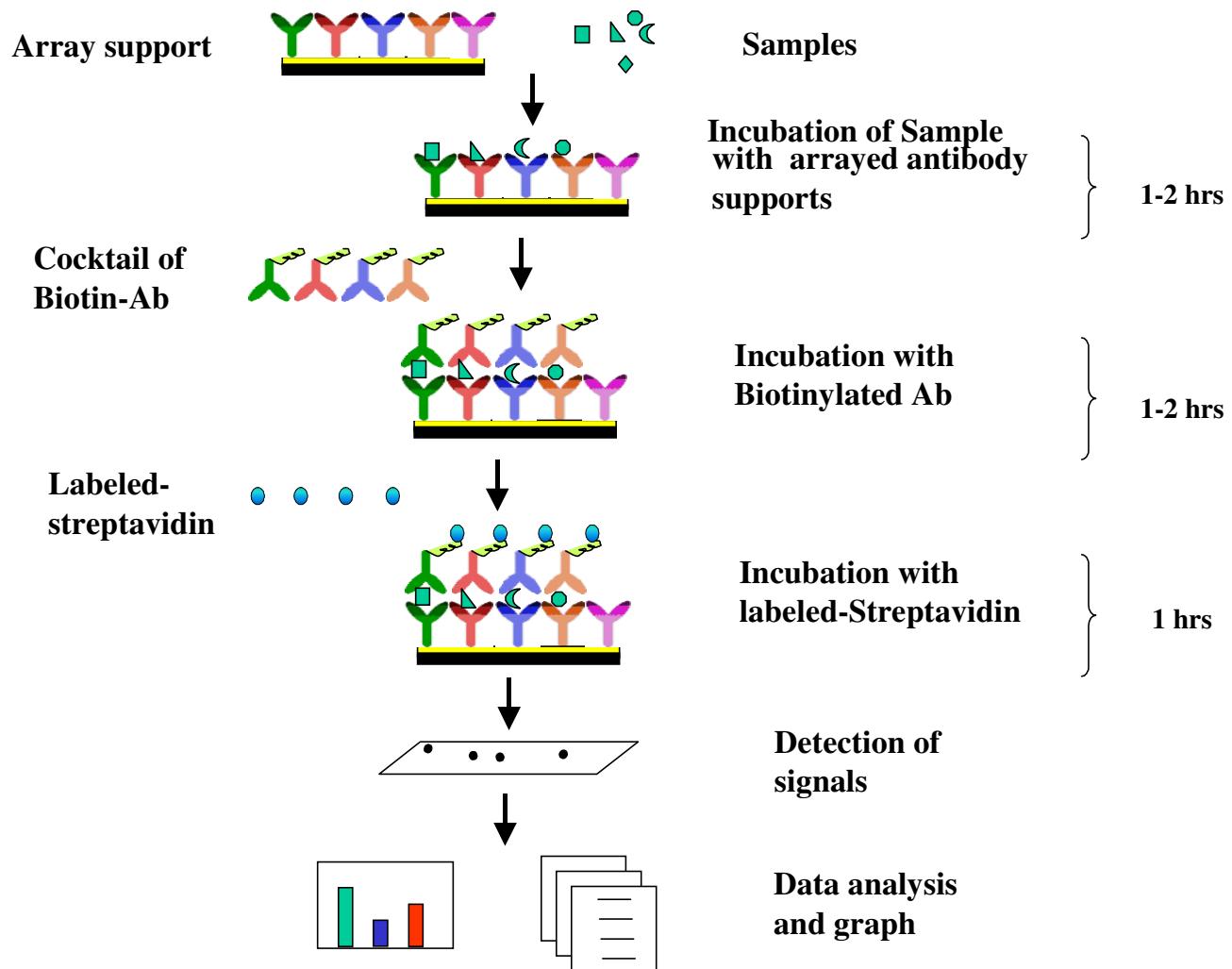
Simultaneous detection of multiple cytokines undoubtedly provides a powerful tool to study cytokines. Cytokines play an important role in innate immunity, apoptosis, angiogenesis, cell growth and differentiation<sup>12</sup>. Cytokines are involved in most disease processes, including cancer and cardiac diseases. The interaction between cytokines and the cellular immune system is a dynamic process. The interactions of positive and negative stimuli, and positive as well as negative regulatory loops are complex and often involve multiple cytokines.

Without doubt, simultaneous detection of multiple cytokines provides a powerful tool to study cytokines.

1. LPS induces the interaction of a transcription factor, LPS-induced TNF- $\alpha$  factor, and STAT6(B) with effects on multiple cytokines. Tang X, Marciano DL, Leeman SE, Amar S. **PNAS**. April 5, 2005 vol. 102 no. 14 5132-5137
2. HIV-1-mediated apoptosis of neuronal cells: Proximal molecular mechanisms of HIV-1-induced encephalopathy. Xu Y, Kulkoshy J, Pomerantz RJ. **PNAS**. 2004 May 4, 2004 Vol. 101 No. 18.

3. Synergistic increases in intracellular Ca(2+), and the release of MCP-1, RANTES, and IL-6 by astrocytes treated with opiates and HIV-1 Tat. **GLIA**. 2005 Apr 15;50(2):91-106.
4. Bone Marrow Stroma Influences Transforming Growth Factor- $\beta$  Production in Breast Cancer Cells to Regulate c-myc Activation of the Preprotachykinin-I Gene in Breast Cancer Cells. Oh HS, Moharita A, Rameshwar P. **Cancer Res**. 64, 6327-6336.
5. Recombinant Herpes Simplex Virus Type 1 (HSV-1) Codelivering Interleukin-12p35 as a Molecular Adjuvant Enhances the Protective Immune Response against Ocular HSV-1 Challenge **J Virol**. Mar. 2005 Vol. 79, No. 6.
6. Dysregulated Inflammatory Response to *Candida albicans* in a C5-Deficient Mouse Strain. Alaka Mullick, Miria Elias, Serge Picard, Philippe Gros. **Infect Immunity**, Oct. 2004, p. 5868-5876.
7. Leukotriene B<sub>4</sub> Strongly Increases Monocyte Chemoattractant Protein-1 in Human Monocytes Li Huang, Annie Zhao, Frederick Wong, Julia M. Ayala, Jisong Cui **Arterioscler Thromb Vascul Biol**. 2004;24:1783-1788
8. Human CD1d-unrestricted NKT cells release chemokines upon Fas engagement. Giroux M and François Denis. Yan Xu, Joseph Kulkoshy, Roger j. Pomerantz. **Blood**. prepublished online September 2, 2004; DOI 10.1182/blood-2004-04-1537
9. Monitoring the response of orthotopic bladder tumors to granulocyte macrophage colony-stimulating factor therapy using the prostate-specific antigen gene as a reporter. Wu Q, Esuvaranathan K, Mahendran R. **Clin Cancer Res**. 2004 Oct 15; 10(20):6977-84.
10. Neuroglial activation and neuroinflammation in the brain of patients with autism (p NA). Vargas DL, Nascimbene C, Krishnan C, Zimmerman AW, Pardo CA. **Ann Neurology**. 2005 Jan 1; DOI: 10.1002/ana.20315
11. Cytokine profiling of macrophages exposed to *Porphyromonas gingivalis*, its LPS or its FimA. Zhou Q, Desta T, Graves DT, Amar S. **Infect Immunity** (IAI). 2005 Feb;73(2):935-43.

## Here's how it works



## II. Materials Provided

Upon receipt, all components of the RayBio® Human Cytokine Antibody Array kit should be stored at -20°C to -80°C. At -20°C to -80°C the kit will retain complete activity for up to 6 months. Once thawed, the array membranes and Blocking Buffer should be kept at -20°C and all other component should be stored at 4°C. After thawing the reagents, the kit must be used within three months, and please use the kit within six months of purchase.

COMPONENT	2 SAMPLE KIT	4 SAMPLE KIT	8 SAMPLE KIT
Human Antibody Arrays	2 membranes	4 membranes	8 membranes
Blocking Buffer		1 vial (25 ml)	2 vials (25 ml/ea)
Biotinylated Antibody Cocktail	1 vial	2 vials	4 vials
1,000X HRP-Streptavidin Concentrate			1 vial (50 µl)
20X Wash Buffer I Concentrate		1 vial (10 ml)	1 vial (20 ml)
20X Wash Buffer II Concentrate		1 vial (10 ml)	1 vial (20 ml)
2X Cell Lysis Buffer Concentrate		1 vial (10 ml)	1 vial (16 ml)
Detection Buffer C		1 vial (1.5 ml)	1 vial (2.5 ml)
Detection Buffer D		1 vial (1.5 ml)	1 vial (2.5 ml)
8-Well Incubation Tray w/ Lid			1 tray
<b>Other Kit Components:</b> Plastic Sheets, Array Map Template, User Manual			

## Additional Materials Required

- Small plastic boxes or containers
- Orbital shaker
- Plastic sheet protector or SaranWrap
- Kodak X-Omat AR film (REF 165 1454) and film processor or Chemiluminescence imaging system

### **III. Overview and General Considerations**

#### **A. Preparation of Samples**

- Use serum-free conditioned media if possible.
- If serum-containing media is required, use an uncultured media aliquot as a negative control sample, since many types of sera contain cytokines.
- For cell lysates and tissue lysates, we recommend using RayBio® Cell Lysis Buffer to extract proteins from cell or tissue (e.g. using homogenizer). Dilute 2X RayBio® Cell Lysis Buffer with H<sub>2</sub>O (we recommend adding proteinase inhibitors to Cell Lysis Buffer before use). After extraction, spin the sample down and save the supernatant for your experiment. Determine protein concentration.
- We recommend using per membrane:
  - 1 ml of Conditioned media (undiluted), or
  - 1 ml of 2-fold to 5-fold diluted sera or plasma, or
  - 50-500 µg of total protein for cell lysates and tissue lysates (use ~200-250 µg of total protein for first experiment) **Dilute the lysate at least 10 fold with 1 X blocking buffer.**

*Note: The amount of sample used depends on the abundance of cytokines. More of the sample can be used if the signals are too weak. If the signals are too strong, the sample can be diluted further.*

○

*If you experience high background, you may further dilute your sample.*

#### **B. Handling Array Membranes**

- Always use forceps to handle membranes, and grip the membranes by the edges only.
- Never allow the array membranes to dry during experiments.

## **C. Incubation**

- Completely cover the membranes with sample or buffer during incubation, and cover the eight-well tray with lid to avoid drying.
- Avoid foaming during incubation steps.
- Perform all incubation and wash steps under gentle rotation.
- Several incubation steps such as step 2 (blocking), step 3 (sample incubation), step 8 (biotin-Ab incubation) and step 11 (HRP-streptavidin incubation) may be done at 4°C for overnight, but make sure to cover the 8 well plate tightly to prevent evaporation.

## **IV. Protocol**

### **A. Blocking and Incubation**

1. Place each membrane into the provided eight-well tray ("—" mark is on the side printed with antibodies).
2. Add 2 ml Blocking Buffer and incubate at room temperature for 30 min to block membranes. Make sure there are no bubbles between the membranes.
3. Decant Blocking Buffer from each container, and incubate membranes with 1 ml of sample at room temperature for 1 to 2 hours. Dilute sample using Blocking Buffer if necessary.

*Note: Incubation may be done at 4°C for overnight.*

4. Decant the samples from each container, and wash 3 times with 2 ml of 1X Wash Buffer I at room temperature with shaking. Please allow 5 min per wash. Dilute 20X Wash Buffer I with H<sub>2</sub>O.
5. Wash 2 times with 2 ml of 1X Wash Buffer II at room temperature with shaking. Allow 5 min per wash. Dilute 20X Wash Buffer II with H<sub>2</sub>O.

6. Prepare working solution for primary antibody.

Add 100  $\mu$ l of Blocking Buffer to the Biotin-Conjugated Anti-Cytokines tube. Mix gently and transfer all mixture to a tube containing 2 ml of Blocking Buffer.

*Note: the diluted biotin-conjugated antibodies can be stored at 4°C for 2-3 days.*

7. Add 1 ml of diluted biotin-conjugated antibodies to each membrane. Incubate at room temperature for 1-2 hours.

*Note: incubation may be done at 4°C for overnight.*

8. Wash as directed in steps 4 and 5.

9. Add 2 ml of **1,000** fold diluted HRP-conjugated streptavidin (e.g. add 2  $\mu$ l of HRP-conjugated streptavidin to **1998**  $\mu$ l Blocking Buffer) to each membrane.

*Note: mix the tube containing 1,000X HRP-Conjugated Streptavidin well before use since precipitation may form during storage.*

10. Incubate at room temperature for 2 hours.

*Note: incubation may be done at 4°C for overnight.*

11. Wash as directed in steps 4 and 5.

## **B. Detection**

\* **Do not let the membrane dry out during detection. The detection process must be completed within 40 minutes without stopping.**

1. Proceed with the detection reaction.

Add 250 µl of 1X Detection Buffer **C** and 250 µl of 1X Detection Buffer **D** for one membrane; mix both solutions; Drain off excess wash buffer by holding the membrane vertically with forceps. Place membrane protein side up (“-“ mark is on the protein side top left corner) on a clean plastic sheet (provided in the kit). Pipette the mixed Detection Buffer on to the membrane and incubated at room temperature for 2 minute. Ensure that the detection mixture is completely and evenly covers the membrane without any air bubbles.

2. Drain off any excess detection reagent by holding the membrane vertically with forceps and touching the edge against a tissue. Gently place the membrane, protein side up, on a piece of plastic sheet (“-“ mark is on the protein side top left corner). Cover another piece of plastic sheet on the array. Gently smooth out any air bubble. Avoid using pressure on the membrane.

3. Expose the array to x-ray film (we recommend to use Kodak X-Omat AR film) and detect the signal using film developer, or the signal can be detected directly from the membrane using a chemiluminescence imaging system.

Expose the membranes for 40 Seconds. Then re-expose the film according to the intensity of signals. If the signals are too strong (background too high), reduce exposure time (e.g. 5-30 seconds). If the signals are too weak, increase exposure time (e.g. 5-20 min or overnight. Or re-incubate membranes overnight with 1X HRP-conjugated streptavidin, and redo detection in the second day).

4. Save membranes in -20°C to -80°C for future references.

## V. Interpretation of Results:

The following figure shows RayBio® Human Cytokine Antibody Array membranes probed with conditioned media from two different cell lines. Membranes were exposed to Kodak X-Omat film at room temperature for 1 minute. The biotin-conjugated IgG produces positive signals, which can be used to identify the orientation and to compare the relative expression levels among the different membranes.

One important parameter is background. To obtain the best results, we suggest that several exposures be attempted. We also strongly recommend using a negative control in which the sample is replaced with an appropriate mock buffer according to the array protocol, particularly during your first experiment.

Typical results using RayBio® Cytokine Antibody arrays



By comparing the signal intensities, relative expression levels of cytokines can be made. The intensities of signals can be quantified by densitometry. Positive control can be used to normalize the results from different membranes being compared. The signals also can be detected and quantitated by using a chemiluminescence imaging device.

The **RayBio® Analysis Tool** is a program specifically designed for analysis of RayBio® Cytokine Antibody Arrays. This tool will not only assist in compiling and organizing your data, but also reduces your calculations to a “copy and paste.” Call RayBiotech, Inc. at 770-729-2992 for ordering information.

## RayBio® Human Cytokine Antibody Array 1 Map

	a	b	c	d	e	f	g	h
1	Pos	Pos	Neg	Neg	GCSF	GM-CSF	GRO	GRO-α
2	Pos	Pos	Neg	Neg	GCSF	GM-CSF	GRO	GRO-α
3	IL-1α	IL-2	IL-3	IL-5	IL-6	IL-7	IL-8	IL-10
4	IL-1α	IL-2	IL-3	IL-5	IL-6	IL-7	IL-8	IL-10
5	IL-13	IL-15	IFN-γ	MCP-1	MCP-2	MCP-3	MIG	RANTES
6	IL-13	IL-15	IFN-γ	MCP-1	MCP-2	MCP-3	MIG	RANTES
7	TGF-β1	TNF-α	TNF-β	Blank	Blank	Blank	Blank	Pos
8	TGF-β1	TNF-α	TNF-β	Blank	Blank	Blank	Blank	Pos

Note: GRO detects CXCL1, CXCL2, CXCL3; GRO-α detects only CXCL1

Note: TGF-β1 detects only active form

## RayBio® Human Cytokine Antibody Array 3 Map

	a	b	c	d	e	f	g	h	i	j	k	l
1	Pos	Pos	Neg	Neg	ENA-78	GCSF	GM-CSF	GRO	GRO-α	I-309	IL-1α	IL-1 β
2	Pos	Pos	Neg	Neg	ENA-78	GCSF	GM-CSF	GRO	GRO-α	I-309	IL-1α	IL-1 β
3	IL-2	IL-3	IL-4	IL-5	IL-6	IL-7	IL-8	IL-10	IL-12 p40p70	IL-13	IL-15	IFN-γ
4	IL-2	IL-3	IL-4	IL-5	IL-6	IL-7	IL-8	IL-10	IL-12 p40p70	IL-13	IL-15	IFN-γ
5	MCP-1	MCP-2	MCP-3	MCSF	MDC	MIG	MIP-1 δ	RANTES	SCF	SDF-1	TARC	TGF- β1
6	MCP-1	MCP-2	MCP-3	MCSF	MDC	MIG	MIP-1 δ	RANTES	SCF	SDF-1	TARC	TGF- β1
7	TNF-α	TNF-β	EGF	IGF-I	Angiogenin	Oncostatin M	Thrombopoietin	VEGF	PDGF BB	Leptin	Neg	Pos
8	TNF-α	TNF-β	EGF	IGF-I	Angiogenin	Oncostatin M	Thrombopoietin	VEGF	PDGF BB	Leptin	Neg	Pos

Note: GRO detects CXCL1, CXCL2, CXCL3; GRO-α detects only CXCL1

Note: IL-12 p40p70 detects both IL-12 p40 and IL-12 p70

Note: VEGF detects VEGF-165 and VEGF-121

## RayBio® Human Cytokine Antibody Array 4 Map

	a	b	c	d	e	f	g	h	i	j	k	l
1	Pos	Pos	Neg	Neg	BDNF	BLC	CKβ 8-1	Eotaxin	Eotaxin-2	Eotaxin-3	FGF-4	FGF-6
2	Pos	Pos	Neg	Neg	BDNF	BLC	CKβ 8-1	Eotaxin	Eotaxin-2	Eotaxin-3	FGF-4	FGF-6
3	FGF-7	FGF-9	Fit-3 Ligand	Fractalkine	GCP-2	GDNF	HGF	IGFBP-1	IGFBP-2	IGFBP-3	IGFBP-4	IL-16
4	FGF-7	FGF-9	Fit-3 Ligand	Fractalkine	GCP-2	GDNF	HGF	IGFBP-1	IGFBP-2	IGFBP-3	IGFBP-4	IL-16
5	IP-10	LIF	LIGHT	MCP-4	MIF	MIP-3α	NAP-2	NT-3	NT-4	Osteoprotegerin	PARC	PIGF
6	IP-10	LIF	LIGHT	MCP-4	MIF	MIP-3α	NAP-2	NT-3	NT-4	Osteoprotegerin	PARC	PIGF
7	TGF-β2	TGF-β3	TIMP-1	TIMP-2	Blank	Blank	Blank	Blank	Blank	Blank	Blank	Pos
8	TGF-β2	TGF-β3	TIMP-1	TIMP-2	Blank	Blank	Blank	Blank	Blank	Blank	Blank	Pos

## RayBio® Human Cytokine Antibody Array 5 Map

	A	B	C	D	E	F	G	H	I	J	K
1	Pos	Pos	Pos	Pos	Neg	Neg	ENA-78	GCSF	GM-CSF	GRO	GRO- $\alpha$
2	I-309	IL-1 $\alpha$	IL-1 $\beta$	IL-2	IL-3	IL-4	IL-5	IL-6	IL-7	IL-8	IL-10
3	IL-12 p40p70	IL-13	IL-15	IFN- $\gamma$	MCP-1	MCP-2	MCP-3	MCSF	MDC	MIG	MIP-1 $\beta$
4	MIP-1 $\delta$	RANTES	SCF	SDF-1	TARC	TGF- $\beta$ 1	TNF- $\alpha$	TNF- $\beta$	EGF	IGF-I	Angiogenin
5	Oncostatin M	Thrombopoietin	VEGF	PDGF-BB	Leptin	BDNF	BLC	Ck $\beta$ 8-1	Eotaxin	Eotaxin-2	Eotaxin-3
6	FGF-4	FGF-6	FGF-7	FGF-9	Flt-3 Ligand	Fractalkine	GCP-2	GDNF	HGF	IGFBP-1	IGFBP-2
7	IGFBP-3	IGFBP-4	IL-16	IP-10	LIF	LIGHT	MCP-4	MIF	MIP-3 $\alpha$	NAP-2	NT-3
8	NT-4	Osteopontin	Osteoprotegerin	PARC	PIGF	TGF- $\beta$ 2	TGF- $\beta$ 3	TIMP-1	TIMP-2	Pos	Pos

Note: GRO detects CXCL1, CXCL2, CXCL3; GRO- $\alpha$  detects only CXCL1

Note: IL-12 p40p70 detects both IL-12 p40 and IL-12 p70

Note: TGF- $\beta$ 1 detects only active form

Note: VEGF detects VEGF-165 and VEGF-121

## RayBio® Human Cytokine Antibody Array 6 (60)

	a	b	c	d	e	f	g	h	i	j	k	l	m	n
1	POS	POS	NEG	NEG	Blank	Angiogenin	BDNF	BLC	BMP-4	BMP-6	CK $\beta$ 8-1	CNTF	EGF	Eotaxin
2	POS	POS	NEG	NEG	Blank	Angiogenin	BDNF	BLC	BMP-4	BMP-6	CK $\beta$ 8-1	CNTF	EGF	Eotaxin
3	Eotaxin-2	Eotaxin-3	FGF-6	FGF-7	Fit-3 Ligand	Fractalkine	GCP-2	GDNF	GM-CSF	I-309	IFN- $\gamma$	IGFBP-1	IGFBP-2	IGFBP-4
4	Eotaxin-2	Eotaxin-3	FGF-6	FGF-7	Fit-3 Ligand	Fractalkine	GCP-2	GDNF	GM-CSF	I-309	IFN- $\gamma$	IGFBP-1	IGFBP-2	IGFBP-4
5	IGF-I	IL-10	IL-13	IL-15	IL-16	IL-1 $\alpha$	IL-1 $\beta$	IL-1ra	IL-2	IL-3	IL-4	IL-5	IL-6	IL-7
6	IGF-I	IL-10	IL-13	IL-15	IL-16	IL-1 $\alpha$	IL-1 $\beta$	IL-1ra	IL-2	IL-3	IL-4	IL-5	IL-6	IL-7
7	Leptin	LIGHT	MCP-1	MCP-2	MCP-3	MCP-4	M-CSF	MDC	MIG	MIP-1 $\delta$	MIP-3 $\alpha$	NAP-2	NT-3	PARC
8	Leptin	LIGHT	MCP-1	MCP-2	MCP-3	MCP-4	M-CSF	MDC	MIG	MIP-1 $\delta$	MIP-3 $\alpha$	NAP-2	NT-3	PARC
9	PDGF-BB	RANTES	SCF	SDF-1	TARC	TGF- $\beta$ 1	TGF- $\beta$ 3	TNF- $\alpha$	TNF- $\beta$	Blank	Blank	Blank	Blank	POS
10	PDGF-BB	RANTES	SCF	SDF-1	TARC	TGF- $\beta$ 1	TGF- $\beta$ 3	TNF- $\alpha$	TNF- $\beta$	Blank	Blank	Blank	Blank	POS

Note: TGF- $\beta$ 1 detects only active form

## RayBio® Human Cytokine Antibody Array 7 (60)

	a	b	c	d	e	f	g	h	i	j	k	l	m	n
1	POS	POS	NEG	NEG	Blank	Acrp30	AgRP	Angiopoietin-2	Amphiregulin	Axl	bFGF	b-NGF	BTC	CCL-28
2	POS	POS	NEG	NEG	Blank	Acrp30	AgRP	Angiopoietin-2	Amphiregulin	Axl	bFGF	b-NGF	BTC	CCL-28
3	CTACK	Dtk	EGF-R	ENA-78	Fas/TNFRSF6	FGF-4	FGF-9	GCSF	GITR-Ligand	GITR	GRO	GRO- $\alpha$	HCC-4	HGF
4	CTACK	Dtk	EGF-R	ENA-78	Fas/TNFRSF6	FGF-4	FGF-9	GCSF	GITR-Ligand	GITR	GRO	GRO- $\alpha$	HCC-4	HGF
5	ICAM-1	ICAM-3	IGFBP-3	IGFBP-6	IGF-I SR	IL-1 R4/ST2	IL-1 RI	IL-11	IL-12 p40	IL-12 p70	IL-17	IL-2 R alpha	IL-6 R	IL-8
6	ICAM-1	ICAM-3	IGFBP-3	IGFBP-6	IGF-I SR	IL-1 R4/ST2	IL-1 RI	IL-11	IL-12 p40	IL-12 p70	IL-17	IL-2 R alpha	IL-6 R	IL-8
7	I-TAC	Lymphotactin	MIF	MIP-1 $\alpha$	MIP-1 $\beta$	MIP-3 $\beta$	MSP- $\alpha$	NT-4	Osteoprotegerin	Oncostatin M	PIGF	sgp130	sTNF RII	sTNF-RI
8	I-TAC	Lymphotactin	MIF	MIP-1 $\alpha$	MIP-1 $\beta$	MIP-3 $\beta$	MSP- $\alpha$	NT-4	Osteoprotegerin	Oncostatin M	PIGF	sgp130	sTNF RII	sTNF-RI
9	TECK	TIMP-1	TIMP-2	Thrombopoietin	TRAIL R3	TRAIL R4	uPAR	VEGF	VEGF-D	Blank	Blank	Blank	Blank	POS
10	TECK	TIMP-1	TIMP-2	Thrombopoietin	TRAIL R3	TRAIL R4	uPAR	VEGF	VEGF-D	Blank	Blank	Blank	Blank	POS

Note: GRO detects CXCL1, CXCL2, CXCL3; GRO- $\alpha$  detects only CXCL1

## RayBio® Human Cytokine Antibody Array 8 (54)

	a	b	c	d	e	f	g	h	i	j	k	l	m	n
1	POS	POS	NEG	NEG	BLANK	Activin A	ALCAM	B7-1(CD80)	BMP-5	BMP-7	Cardiotrophin-1	CD14	CXCL-16	DR6 (TNFRSF21)
2	POS	POS	NEG	NEG	BLANK	Activin A	ALCAM	B7-1(CD80)	BMP-5	BMP-7	Cardiotrophin-1	CD14	CXCL-16	DR6 (TNFRSF21)
3	Endoglin	ErbB3	E-Selectin	Fas Ligand	ICAM-2	IGF-II	IL-1 R II	IL-10 R beta	IL-13 R alpha 2	IL-18 BP alpha	IL-18 R beta	MMP-3	IL-2 R beta	IL-2 R gamma
4	Endoglin	ErbB3	E-Selectin	Fas Ligand	ICAM-2	IGF-II	IL-1 R II	IL-10 R beta	IL-13 R alpha 2	IL-18 BP alpha	IL-18 R beta	MMP-3	IL-2 R beta	IL-2 R gamma
5	IL-21R	IL-5 R alpha	IL-9	IP-10	LAP	Leptin R	LIF	L-Selectin	M-CSF R	MMP-1	MMP-13	MMP-9	MPIF-1	NGF R
6	IL-21R	IL-5 R alpha	IL-9	IP-10	LAP	Leptin R	LIF	L-Selectin	M-CSF R	MMP-1	MMP-13	MMP-9	MPIF-1	NGF R
7	PDGF AA	PDGF-AB	PDGF R alpha	PDGF R beta	PECAM-1	Prolactin	SCF R	SDF-1beta	Siglec-5	TGF-alpha	TGF beta 2	Tie-1	Tie-2	TIMP-4
8	PDGF AA	PDGF-AB	PDGF R alpha	PDGF R beta	PECAM-1	Prolactin	SCF R	SDF-1beta	Siglec-5	TGF-alpha	TGF beta 2	Tie-1	Tie-2	TIMP-4
9	VE-Cadherin	VEGF R2	VEGF R3	BLANK	BLANK	BLANK	BLANK	BLANK	BLANK	BLANK	BLANK	BLANK	BLANK	POS
10	VE-Cadherin	VEGF R2	VEGF R3	BLANK	BLANK	BLANK	BLANK	BLANK	BLANK	BLANK	BLANK	BLANK	BLANK	POS

Note: All MMP antibodies detect both pro and mature forms

## RayBio® Human Cytokine Antibody Array 9 (51)

	a	b	c	d	e	f	g	h	i	j	k	l
1	POS	POS	NEG	NEG	Adiposin	BCAM	CD30	CD40	FcR RIIB/C	Ferritin	FLRG	Follistatin
2	POS	POS	NEG	NEG	Adiposin	BCAM	CD30	CD40	FcR RIIB/C	Ferritin	FLRG	Follistatin
3	Furin	Galectin-7	GDF-15	Growth Hormon	IL-10 R alpha	IL-22	IL-28A	IL29	IL-31	Insulin	Luteinizing Hormone	LIMPII
4	Furin	Galectin-7	GDF-15	Growth Hormon	IL-10 R alpha	IL-22	IL-28A	IL29	IL-31	Insulin	Luteinizing Hormone	LIMPII
5	LYVE-1	Marapsin	MICA	MICB	MMP-2	MMP-7	MMP-8	MMP-10	NCAM-1	Nidogen-1	NrCAM	NRG1-beta 1
6	LYVE-1	Marapsin	MICA	MICB	MMP-2	MMP-7	MMP-8	MMP-10	NCAM-1	Nidogen-1	NrCAM	NRG1-beta 1
7	Osteopontin	PAI-I	Platelet Factor 4	PSA-total	RAGE	RANK	Resistin	SAA	Siglec-9	TACE	TIM-1	TRAIL R2
8	Osteopontin	PAI-I	Platelet Factor 4	PSA-total	RAGE	RANK	Resistin	SAA	Siglec-9	TACE	TIM-1	TRAIL R2
9	Trappin-2	TREM-1	TSH	TSLP	VCAM-1	VEGF-C	XEDAR	Blank	Blank	Blank	Blank	POS
10	Trappin-2	TREM-1	TSH	TSLP	VCAM-1	VEGF-C	XEDAR	Blank	Blank	Blank	Blank	POS

Note: All MMP antibodies detect both pro and mature forms

## RayBio® Human Cytokine Antibody Array 10 (49)

	a	b	c	d	e	f	g	h	i	j	k	l
1	POS	POS	NEG	NEG	4-1BB	ACE-2	Alpha-Fetoprotein	Angiopoietin-1	Angiostatin	ANGPTL4	Bate2 M	BCMA
2	POS	POS	NEG	NEG	4-1BB	ACE-2	Alpha-Fetoprotein	Angiopoietin-1	Angiostatin	ANGPTL4	Bate2 M	BCMA
3	beta IG-H3	CA125	CA15-3	CA19-9	Carbonic Anhydrase IX	Cathepsin S	CCL14a	CCL21	CD23	CD40 Ligand	CEA	CEACAM-1
4	beta IG-H3	CA125	CA15-3	CA19-9	Carbonic Anhydrase IX	Cathepsin S	CCL14a	CCL21	CD23	CD40 Ligand	CEA	CEACAM-1
5	Cripto-1	CRP	DAN	Decorin	DKK-1	DKK-3	DKK-4	DPPIV	E-Cadherin	EDA-A2	EG-VEGF	EpCAM
6	Cripto-1	CRP	DAN	Decorin	DKK-1	DKK-3	DKK-4	DPPIV	E-Cadherin	EDA-A2	EG-VEGF	EpCAM
7	ErbB2	Erythropoietin R	FSH	HB-EGF	hCG $\alpha$ , intact	HVEM	IL-13R1	IL-17B	IL-17C	IL-17F	IL-17R	Procalcitonin
8	ErbB2	Erythropoietin R	FSH	HB-EGF	hCG $\alpha$ , intact	HVEM	IL-13R1	IL-17B	IL-17C	IL-17F	IL-17R	Procalcitonin
9	PSA-free	S-100b	Shh N	Thyroglobulin	Ubiquitin+1	Blank	Blank	Blank	Blank	Blank	Blank	POS
10	PSA-free	S-100b	Shh N	Thyroglobulin	Ubiquitin+1	Blank	Blank	Blank	Blank	Blank	Blank	POS

## RayBio® Human Inflammation Antibody Array 1 Map

	A	B	C	D	E	F	G	H
1	POS	POS	NEG	NEG	EOTAXIN	EOTAXIN-2	GCSF	GM-CSF
2	POS	POS	NEG	NEG	EOTAXIN	EOTAXIN-2	GCSF	GM-CSF
3	IFN- $\gamma$	IL-1 $\alpha$	IL-1 $\beta$	IL-2	IL-3	IL-4	IL-6	IL-7
4	IFN- $\gamma$	IL-1 $\alpha$	IL-1 $\beta$	IL-2	IL-3	IL-4	IL-6	IL-7
5	IL-8	IL-10	IL-11	IL-12 p40	IL-12 p70	IL-13	I-309	TIMP-2
6	IL-8	IL-10	IL-11	IL-12 p40	IL-12 p70	IL-13	I-309	TIMP-2
7	BLANK	BLANK	BLANK	BLANK	BLANK	BLANK	NEG	POS
8	BLANK	BLANK	BLANK	BLANK	BLANK	BLANK	NEG	POS

Note: IL-12 p40 detects only IL-12 p40 and IL-12 p70 detects only IL-12 p70

## RayBio® Human Inflammation Antibody Array 2 Map

	A	B	C	D	E	F	G	H
1	POS	POS	NEG	NEG	ICAM-1	IL-6sR	IL-15	IL-16
2	POS	POS	NEG	NEG	ICAM-1	IL-6sR	IL-15	IL-16
3	IL-17	IP-10	MCP-1	MCP-2	M-CSF	MIG	MIP-1 $\alpha$	MIP-1 $\beta$
4	IL-17	IP-10	MCP-1	MCP-2	M-CSF	MIG	MIP-1 $\alpha$	MIP-1 $\beta$
5	MIP-1 $\delta$	RANTES	TGF- $\beta$ 1	TNF- $\alpha$	TNF- $\beta$	s TNF RI	s TNF RII	PDGF-BB
6	MIP-1 $\delta$	RANTES	TGF- $\beta$ 1	TNF- $\alpha$	TNF- $\beta$	s TNF RI	s TNF RII	PDGF-BB
7	BLANK	BLANK	BLANK	BLANK	BLANK	BLANK	NEG	POS
8	BLANK	BLANK	BLANK	BLANK	BLANK	BLANK	NEG	POS

Note: TGF- $\beta$ 1 detects only active form

## RayBio® Human Inflammation Antibody Array 3 Map

	A	B	C	D	E	F	G	H	I	J	K	L
1	POS	POS	NEG	NEG	EOTAXIN	EOTAXIN-2	GCSF	GM-CSF	ICAM-1	IFN-γ	I-309	IL-1α
2	POS	POS	NEG	NEG	EOTAXIN	EOTAXIN-2	GCSF	GM-CSF	ICAM-1	IFN-γ	I-309	IL-1α
3	IL-1β	IL-2	IL-3	IL-4	IL-6	IL-6sR	IL-7	IL-8	IL-10	IL-11	IL-12 p40	IL-12 p70
4	IL-1β	IL-2	IL-3	IL-4	IL-6	IL-6sR	IL-7	IL-8	IL-10	IL-11	IL-12 p40	IL-12 p70
5	IL-13	IL-15	IL-16	IL-17	IP-10	MCP-1	MCP-2	M-CSF	MIG	MIP-1α	MIP-1β	MIP-1δ
6	IL-13	IL-15	IL-16	IL-17	IP-10	MCP-1	MCP-2	M-CSF	MIG	MIP-1α	MIP-1β	MIP-1δ
7	RANTES	TGF-β1	TNF-α	TNF-β	s TNF RI	s TNF RII	PDGF-BB	TIMP-2	BLANK	BLANK	NEG	POS
8	RANTES	TGF-β1	TNF-α	TNF-β	s TNF RI	s TNF RII	PDGF-BB	TIMP-2	BLANK	BLANK	NEG	POS

Note: IL-12 p40 detects only IL-12 p40 and IL-12 p70 detects only IL-12 p70

Note: TGF-β1 detects only active form

## RayBio® Human Angiogenesis Antibody Array 1 Map

	A	B	C	D	E	F	G	H
1	POS	POS	NEG	NEG	Angiogenin	EGF	ENA-78	b FGF
2	POS	POS	NEG	NEG	Angiogenin	EGF	ENA-78	b FGF
3	GRO	IFN-γ	IGF-I	IL-6	IL-8	LEPTIN	MCP-1	PDGF-BB
4	GRO	IFN-γ	IGF-I	IL-6	IL-8	LEPTIN	MCP-1	PDGF-BB
5	PIGF	RANTES	TGF-β1	TIMP-1	TIMP-2	Thrombopoietin	VEGF	VEGF-D
6	PIGF	RANTES	TGF-β1	TIMP-1	TIMP-2	Thrombopoietin	VEGF	VEGF-D
7	BLANK	BLANK	BLANK	BLANK	BLANK	BLANK	Neg	POS
8	BLANK	BLANK	BLANK	BLANK	BLANK	BLANK	Neg	POS

Note: GRO detects CXCL1, CXCL2, CXCL3

Note: TGF-β1 detects only active form

Note: VEGF detects VEGF-165 and VEGF-121

## RayBio® Human Atherosclerosis Antibody Array 1 Map

	A	B	C	D	E	F	G	H
1	POS	POS	NEG	NEG	Eotaxin	GCSF	GDNF	GM-CSF
2	POS	POS	NEG	NEG	Eotaxin	GCSF	GDNF	GM-CSF
3	ICAM-1	IL-1sRI	IL-1sRII	IFN-γ	IL-1α	IL-1β	MCP-1	M-CSF
4	ICAM-1	IL-1sRI	IL-1sRII	IFN-γ	IL-1α	IL-1β	MCP-1	M-CSF
5	MIP-3α	PDGF-BB	RANTES	TGF-β1	TGF-β2	TGF-β3	TNF-α	TNF-β
6	MIP-3α	PDGF-BB	RANTES	TGF-β1	TGF-β2	TGF-β3	TNF-α	TNF-β
7	BLANK	BLANK	BLANK	BLANK	BLANK	BLANK	NEG	POS
8	BLANK	BLANK	BLANK	BLANK	BLANK	BLANK	NEG	POS

Note: TGF-β1 detects only active form

## RayBio® Human Chemokine Antibody Array 1 Map

	A	B	C	D	E	F	G	H	I	J	K	L
1	POS	POS	NEG	NEG	BLC	CCL28	Ckβ8-1	CTACK	CXCL16	ENA-78	Eotaxin	Eotaxin-2
2	POS	POS	NEG	NEG	BLC	CCL28	Ckβ8-1	CTACK	CXCL16	ENA-78	Eotaxin	Eotaxin-2
3	Eotaxin-3	Fractalkine	GCP-2	GRO	GROα	HCC-4	I-309	I-TAC	IL-8	IP-10	Lymphotactin	MCP-1
4	Eotaxin-3	Fractalkine	GCP-2	GRO	GROα	HCC-4	I-309	I-TAC	IL-8	IP-10	Lymphotactin	MCP-1
5	MCP-2	MCP-3	MCP-4	MDC	MIG	MIP-1α	MIP-1β	MIP-1δ	MIP-3α	MIP-3β	MPIF-1	NAP 2
6	MCP-2	MCP-3	MCP-4	MDC	MIG	MIP-1α	MIP-1β	MIP-1δ	MIP-3α	MIP-3β	MPIF-1	NAP 2
7	PARC	RANTES	SDF-1α	SDF-1β	TARC	TECK	BLANK	BLANK	BLANK	BLANK	BLANK	POS
8	PARC	RANTES	SDF-1α	SDF-1β	TARC	TECK	BLANK	BLANK	BLANK	BLANK	BLANK	POS

Note: GRO detects CXCL1, CXCL2, CXCL3; GROα detects only CXCL1

## RayBio® Human Matrix Metalloproteinase Antibody Array 1 Map

	A	B	C	D	E	F	G	H
1	POS	POS	NEG	NEG	MMP-1	MMP-2	MMP-3	MMP-8
2	POS	POS	NEG	NEG	MMP-1	MMP-2	MMP-3	MMP-8
3	MMP-9	MMP-10	MMP-13	TIMP-1	TIMP-2	TIMP-4	NEG	POS
4	MMP-9	MMP-10	MMP-13	TIMP-1	TIMP-2	TIMP-4	NEG	POS

Note: All MMP antibodies detect both pro and active forms

## RayBio® Human Growth Factor Antibody Array 1 Map

	A	B	C	D	E	F	G	H	I	J	K	L
1	POS	POS	NEG	NEG	AR	bFGF	b-NGF	EGF	EGF R	FGF-4	FGF-6	FGF-7
2	POS	POS	NEG	NEG	AR	bFGF	b-NGF	EGF	EGF R	FGF-4	FGF-6	FGF-7
3	GCSF	GDNF	GM-CSF	HB-EGF	HGF	IGFBP-1	IGFBP-2	IGFBP-3	IGFBP-4	IGFBP-6	IGF-I	IGF-I SR
4	GCSF	GDNF	GM-CSF	HB-EGF	HGF	IGFBP-1	IGFBP-2	IGFBP-3	IGFBP-4	IGFBP-6	IGF-I	IGF-I SR
5	IGF-II	M-CSF	M-CSF R	NT-3	NT-4	PDGF R α	PDGF R β	PDGF-AA	PDGF-AB	PDGF-BB	PIGF	SCF
6	IGF-II	M-CSF	M-CSF R	NT-3	NT-4	PDGF R α	PDGF R β	PDGF-AA	PDGF-AB	PDGF-BB	PIGF	SCF
7	SCF R	TGF-α	TGF-β	TGF-β 2	TGF-β 3	VEGF	VEGF R2	VEGF R3	VEGF-D	BLANK	BLANK	POS
8	SCF R	TGF-α	TGF-β	TGF-β 2	TGF-β 3	VEGF	VEGF R2	VEGF R3	VEGF-D	BLANK	BLANK	POS

Abbreviations: ANG, angiogenin; OSM, oncostatin M; TPO, thrombopoietin; POS, positive control; NEG, negative control. All others use standard abbreviations.

## Human Custom Antibody Array List (285 proteins)

4-1BB/TNFRSF9	CNTF	GDNF	IL-18 R alpha	MIP-1 alpha	SCF
ACE-2	Cripto-1	GITR	IL-18 R beta	MIP-1 beta	SCF R
Activin A	CRP	GITR Ligand	IL-1ra	MIP-1 delta	SDF-1 alpha
Adiponectin/Acrp30	CTACK/CCL27	GM-CSF	IL-2	MIP-3 alpha	SDF-1 beta
Adipsin/Factor D	CTLA-4	GRO	IL-2 R alpha	MIP-3 beta	sgp130
AFP	CXCL16	GRO-a	IL-2 R beta	MMP-1	Shh N
AgRP(ART)	DAN	Growth Hormom	IL-2 R gamma	MMP-2	Siglec-5
ALCAM	Decorin	HB-EGF	IL-21 R	MMP-3	Siglec-9
Angiogenin	DKK-1	HCC-4/CCL16	IL-22	MMP-7	sTNF RII
Angiopoietin-1	DKK-3	hCGa, intact	IL-28A/IFN-lambda	MMP-8	sTNT RI
Angiopoietin-2	DKK-4	HGF	IL29/IFN-lambda 1	MMP-9	TACE
Angiostatin	DPPIV/CD26	HVEM	IL-3	MMP-10	TARC
ANGPTL4	DR6	I-309	IL-31	MMP-13	TECK/CCL25
AR (amphiregulin)	Dtk	ICAM-1	IL-4	MPIF-1	TGF-alpha
Axl	E-Cadherin	ICAM-2	IL-5	MSP a Chain	TGF-beta 1
B7-1(CD80)	EDA-A2	ICAM-3	IL-5 R alpha	NAP-2	TGF-beta 2
Bate2 M	EGF	IFN-gamma	IL-6	NCAM-1	TGF-beta 3
BCAM	EGF R	IGFBP-1	IL-6 sR	NGF R	Thyroglobulin
BCMA/TNFRSF17	EG-VEGF/PK1	IGFBP-2	IL-7	Nidogen-1/Entactin	Tie-1
BDNF	ENA-78	IGFBP-3	IL-8	NRCAM	Tie-2
beta IG-H3	Endoglin	IGFBP-4	IL-9	NRG1-beta 1/HRG1-beta 1	TIM-1
Betacellulin (BTC)	Endostatin	IGFBP-5	IL-9 R	NT-3	TIMP-1
bFGF	Eotaxin	IGFBP-6	Insulin	NT-4	TIMP-2
BLC	Eotaxin-2	IGF-I	IP-10	Oncostatin M	TIMP-4
BMP-4	Eotaxin-3	IGF-I sR	I-TAC/CXCL11	Osteopontin	TNF-alpha
BMP-5	EpCAM/TROP1	IGF-II	LAP(TGF-b1)	Osteoprotegerin	TNF-beta
BMP-6	ErbB2	IL-1 alpha	Leptin R	PAI-I	TPO
BMP-7	ErbB3	IL-1 beta	LEPTIN(OB)	PARC	TRAIL R1
b-NGF	Erythropoietin R (EPO R)	IL-1 R4/ST2	LH	P-Cadherin	TRAIL R2
BTC	E-Selectin	IL-1 sRI	LIF	PDGF R alpha	TRAIL R3
CA125	Fas Ligand	IL-1 sRII	LIGHT	PDGF R beta	TRAIL R4
CA15-3	Fas/TNFRSF6	IL-10	LIMPII/SR-B2	PDGF-AA	Trappin-2/Elastin
CA19-9	FcR IIIB/C	IL-10 R alpha	Lipocalin-2/NGAL	PDGF-AB	TREM-1
Carbonic Anhydrase IX(CA9)	Ferritin	IL-10 R beta	L-Selectin	PDGF-BB	TROY
Cardiotrophin-1 (CT-1)	FGF-4	IL-11	Lymphotactin	PECAM-1	TSH
Cathepsin S	FGF-6	IL-12 p40	LYVE-1	Platelet Factor 4	TSPL
CCL14a/HCC-1	FGF-7	IL-12 p70	Marapsin/Pancreasin	PIGF	u PAR
CCL21/6ckine	FGF-9	IL-13	MCP-1	Procalcitonin/Calcitonin	Ubiquitin+1
CCL28/VIC	FLRG	IL-13 Ra1	MCP-2	Prolactin	VCAM-1
CD14	Flt-3 Ligand	IL-13 Ra2	MCP-3	PSA-free	VE-Cadherin
CD23/Fc epsilon RII	Follistatin	IL-15	MCP-4	PSA-total	VEGF
CD27	Fractalkine	IL-16	MCSF	P-selectin	VEGF R2
CD30	FSH	IL-17	M-CSF R	RAGE	VEGF R3
CD40	Furin	IL-17B	MDC	RANK	VEGF-C
CD40 Ligand	Galectin-7	IL-17C	MICA	RANTES	VEGF-D
CEA	GCP-2	IL-17F	MICB	Resistin	
CEACAM-1	GCSF	IL-17R	MIF	S-100b	
CK beta 8-1	GDF-15/MIC-1	IL-18 BPa	MIG	SAA	

RayBiotech, Inc., the protein array pioneer company, strives to research and develop new products to meet demands of the biomedical community. RayBio's patent-pending technology allows detection of 274 cytokines, chemokines and other proteins in a single experiment. Our format is simple, sensitive, reliable and cost effective. Products include: Cytokine Arrays, Chemokine Arrays, ELISA kits, Phosphotyrosine kits, EIA kits, Recombinant Proteins, Antibodies, and custom services.

1. Antibody arrays
2. Cytokine antibody array
  - Human cytokine antibody arrays
  - Mouse cytokine antibody arrays
  - Rat cytokine antibody arrays
- Pathway- or disease-focused antibody arrays
  - Inflammation antibody array
  - Angiogenesis antibody array
  - Chemokine antibody array
  - Growth factor antibody array
  - MMP antibody array
  - Atherosclerosis antibody array
  - Adipokine antibody arrays
- Antibody analysis tool, software
3. ELISA
4. Cell-based phosphorylation assay
5. Custom antibody arrays
6. Antibody
7. Recombinant protein
8. Cytokine protein arrays
9. Quantibody arrays for quantitative measurement of cytokine and other protein concentration.
10. Phosphorylation antibody arrays
11. Biotin label-based antibody arrays for high density antibody arrays
12. EIA
13. Peptide

RayBiotech also provides excellent custom service:

1. Antibody arrays

2. Protein arrays
3. Peptide synthesis
4. Production of recombinant protein and antibody
5. Peptide arrays
6. Phosphorylation arrays
7. ELISA
8. EIA
9. Assay development

Just simply send your samples and we will do the assay for you.

#### Technology transfer program

Have you developed technologies or reagents of interest to the scientific and research community? RayBiotech can help you commercialize your technologies, reagents and dream.

## VI. Troubleshooting guide

<b>Problem</b>	<b>Cause</b>	<b>Recommendation</b>
Weak signal or no signal	1. Taking too much time for Detection.  2. Film developer does not work properly.  3. Did not mix HRP-streptavidin well before use.  4. Sample is too dilute.  5. Other.	1. The whole Detection process must be completed in 30 min.  2. Fix film developer.  3. Mix tube containing 1,000X HRP-Conjugated Streptavidin well before use since precipitation may form during storage.  4. Increase sample volume, (e.g. using undilute sample) or using more cells (e.g. seed 2 million cells. After 1 or 2 days, change complete medium with low serum medium and collect conditioned medium 2 day later. Use about 1 to 2 ml of conditioned medium for experiment).  5. 1. Reduce blocking concentration by diluting in 1X Wash Buffer II. 2. Slightly increase HRP concentrations. 3. Slightly increase biotin-antibody concentrations. 4. Expose film for overnight to detect weak signal.
Uneven signal	1. Bubbles formed during incubation.  2. Membranes were not completely covered by solution.	1. Remove bubble during incubation.  2. Completely cover membranes with solution.
High background	1. Exposure to x-ray file is too long.  2. Membranes were allowed to dry out during experiment.  3. Sample is too concentrated.	1. Decrease exposure time.  2. Completely cover membranes with solution during experiment.  3. Use more diluted sample.

## Selected References Featuring RayBiotech Arrays

1. Classification and prediction of clinical Alzheimer's diagnosis based on plasma signaling proteins. Ray S, Britschgi M, Herbert C, Takeda-Uchimura Y, et al. **Nature Med.** 2007; 13(11):1359-1362.
2. Systemic Endocrine Instigation of Indolent Tumor Growth Requires Osteopontin. McAllister SS, Gifford AM, Greiner AL, Kelleher SP, Saelzer MP, et al. **Cell.** 2008;133:994-1005.
3. Chemokine Signaling via the CXCR2 Receptor Reinforces Senescence. Acosta JC, O'Loghlen A, Banito A, Guijarro MV, Augert A, et al. **Cell.** 2008;133: 1006-1018.
4. Conjugated Linoleic Acid Induces Human Adipocyte Delipidation: Autocrine/Paracrine Signaling by Adipokines. Brown JM, Boysen MS, Chung S, Fabiyi O, et al. **J Biol Chem.** 2004;279(25):26735-26747.
5. Towards discovery-driven translational research in breast cancer. Celis JE, Moreira JMA, Gromova I, Cabezon T, et al. **FEBS J.** 2004;272:2-15
6. Paneth Cell Cryptdins Act in Vitro as Apical Paracrine Regulators of the Innate Inflammatory Response. Lin PW, Simon PO, Gewirtz AT, Neish AS, et al. **J Biol Chem.** 2004;279(19):19902-19907.
7. HIV-1-mediated apoptosis of neuronal cells: Proximal molecular mechanisms of HIV-1-induced encephalopathy. Xu Y Kulkosky J, Pomerantz RJ. **Proc Natl Acad Sci USA.** 2004;101(18):7071-7075.
8. Tang X, Marciano DL, Leeman SE, Amar S. LPS induces the interaction of a transcription factor, LPS-induced TNF-alpha factor, and STAT6(B) with effects on multiple cytokines. **Proc Natl Acad Sci USA.** 2005;102(14):5132-5137.
9. IL-7 is a potent and proviral strain-specific inducer of latent HIV-1 cellular reservoirs of infected individuals on virally suppressive HAART. Wang F-X, Xu Y, Sullivan J, Souder E, et al. **J Clin Invest.** 2005;155:128–137.
10. Assessment of Some Tools for the Characterization of the Human Osteoarthritic Cartilage Proteome. De Ceuninck F, Marcheteau E, Berger S,

Caliez A, et al. **J Biomol Tech.** 2005;16:256–265.

11. The type 1 lysophosphatidic acid receptor is a target for therapy in bone metastases. Boucharaba A, Serre C-M, Guglielmi J, Bordet J-C, Clezardin P, Peyruchaud O. **Proc Natl Acad Sci USA.** 2006;103(25):9643-9648.
12. Airway cytokine expression measured by means of protein array in exhaled breath condensate: Correlation with physiologic properties in asthmatic patients. Matsunaga K, Yanagisawa S, Ichikawa T, Ueshima K, et al. **J Allergy Clin Immunol.** 2006; 188:84-90.
13. Neuroglial activation and neuroinflammation in the brain of patients with autism. Vargas DL, Nascimbene C, Chitra Krishnan C, Zimmerman AW, Pardo CA. **Ann Neurology.** 2005;57:67-81.
14. Age-dependent cell death and the role of ATP in hydrogen peroxide-induced apoptosis and necrosis. Miyoshi N, Oubrahim H, Chock PB, Stadtman ER. **Proc Natl Acad Sci USA.** 2006;103(6):1727–1731.
15. Moderation of the platelet releasate response by aspirin. Coppinger JA, O'Connor R, Wynne K, Flanagan M, Sullivan M, et al. **Blood.** 2007;109:4786-4792.
16. Cortez DM, Feldman MD, Mummidi S, Valente AJ, Steffensen B, et al. IL-17 stimulates MMP-1 expression in primary human cardiac fibroblasts via p38 MAPK- and ERK1/2-dependent C/EBP-beta, NF-kB, and AP-1 activation. **Am J Physiol Heart Circ Physiol.** 2007;293:H3356-H3365.
17. Microsensor Arrays for Saliva Diagnostics. Walt DR, Blicharz TM, Hayman RB, Rissin DM, et al. **Ann NY Acad Sci.** 2007;1098:389-400.

**Note:**

RayBio® is the trademark of RayBiotech, Inc.

Cytokine protein arrays are RayBiotech patent-pending technology.

This product is intended for research only and is not to be used for clinical diagnosis. Our products may not be resold, modified for resale, or used to manufacture commercial products without written approval by RayBiotech, Inc.

Under no circumstances shall RayBiotech be liable for any damages arising out of the use of the materials.

Products are guaranteed for three months from the date of purchase when handled and stored properly. In the event of any defect in quality or merchantability, RayBiotech's liability to buyer for any claim relating to products shall be limited to replacement or refund of the purchase price.

Kodak X-Omat™ is the trademark of Eastman Kodak Company.

This product is for research use only.

