

# Quantikine<sup>®</sup>

## Human I-TAC/CXCL11 Immunoassay

Catalog Number DCX110

**For the quantitative determination of human interferon gamma inducible T-cell alpha chemoattractant (I-TAC) concentrations in cell culture supernates, serum and plasma.**

*This package insert must be read in its entirety before using this product.*

**FOR RESEARCH USE ONLY.  
NOT FOR USE IN DIAGNOSTIC PROCEDURES.**

# TABLE OF CONTENTS

Contents	Page
INTRODUCTION	2
PRINCIPLE OF THE ASSAY . . . . .	3
LIMITATIONS OF THE PROCEDURE	3
REAGENTS . . . . .	4
STORAGE	4
OTHER SUPPLIES REQUIRED . . . . .	5
PRECAUTION	5
SAMPLE COLLECTION AND STORAGE . . . . .	5
REAGENT PREPARATION	6
ASSAY PROCEDURE . . . . .	7
ASSAY PROCEDURE SUMMARY	8
CALCULATION OF RESULTS . . . . .	9
TYPICAL DATA	9
TECHNICAL HINTS . . . . .	10
PRECISION	10
RECOVERY . . . . .	11
LINEARITY	11
SENSITIVITY . . . . .	11
CALIBRATION	12
SAMPLE VALUES . . . . .	12
SPECIFICITY	13
REFERENCES . . . . .	14
PLATE LAYOUT	15

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## INTRODUCTION

Interferon-inducible T cell alpha chemoattractant (I-TAC/CXCL11) is a non-ELR (lacking the Glu-Leu-Arg tripeptide motif) CXC chemokine. The cDNA encoding I-TAC (1), originally designated  $\beta$ -R1 (2), has also been reported as SCYB9B (3), H174 (4), IP-9 (5), and SCYB11 (6). I-TAC expression is significantly increased in response to IFN- $\beta$  or IFN- $\gamma$ , and is presumed to be involved in pathologies characterized by the presence of activated T cells.

The I-TAC gene consists of four exons interrupted by three introns of 585, 98, and 230 base pairs (7). I-TAC cDNA encodes a 94 amino acid (aa) residue precursor protein with a 21 aa residue putative signal sequence that is cleaved to generate the mature protein (1). I-TAC is 36% and 37% identical in primary structure to the other non-ELR CXC chemokines IP-10 and MIG, respectively (1). Mouse I-TAC exhibits 68% aa sequence homology with human I-TAC (8). In humans, I-TAC, IP-10 and MIG all map to the same locus on chromosome 4 (1, 3, 9).

I-TAC exclusively utilizes CXC chemokine receptor 3 (CXCR3) (1, 10), a G protein-coupled receptor expressed primarily on activated T cells, yet also found on endothelial cells (11). Among activated T cells, CXCR3 is more highly expressed on the Th1 subset (12). IP-10 and MIG also bind CXCR3, but with lower affinity (1, 10) and less potency (1) than I-TAC. Furthermore, there is evidence that I-TAC is able to bind both the free and coupled forms of the receptor, whereas IP-10 and MIG bind only the coupled receptor (10).

The I-TAC promoter region contains the consensus sequences for the interferon-responsive DNA regulatory elements ISRE, GAS, and the cytokine-responsive binding protein NF- $\kappa$ B (7). I-TAC expression is strongly upregulated in response to IFN- $\gamma$ . This effect is dramatically enhanced by addition of IL-1 $\beta$  (1) or TNF- $\alpha$  (7), though these pro-inflammatory cytokines alone fail to induce expression. I-TAC mRNA can be induced in astrocytes and monocytes (1), bronchial epithelial cells (13), intestinal epithelial cells (14), endothelial cells (15, 16), keratinocytes (5, 17), macrophages (15), and neutrophils (18). I-TAC expression is suppressed by the Th2 cytokines IL-4 and IL-10 (18) and also by peroxisome proliferator-activated receptor  $\gamma$  activators (19). I-TAC recruits activated Th1 lymphocytes to sites of inflammation. It is an antagonist of CC chemokine receptor 3 (CCR3) (20), a chemokine receptor expressed on eosinophils and Th2 lymphocytes, suggesting that I-TAC may act to further polarize T cell recruitment at sites of expression. There is evidence that I-TAC may be influential in the migration of different subsets of mature thymocytes during thymus lymphopoiesis (21). I-TAC also exhibits an angiostatic effect, apparently via CXCR3 expressed on endothelial cells in the S/G2-M phase of the cell cycle (11). I-TAC has been implicated in allergic contact dermatitis (22), atherosclerosis (15), mycosis fungoides (a type of T cell lymphoma) (23), and immune-mediated disorders of the central nervous system such as multiple sclerosis (24). In mice, the I-TAC gene is required for acute allograft rejection (25).

The Quantikine I-TAC Immunoassay is a 4.5 hour solid-phase ELISA designed to measure human I-TAC in cell culture supernates, serum and plasma. It contains *E. coli*-expressed recombinant human I-TAC and has been shown to accurately quantitate the recombinant factor. Results obtained using natural human I-TAC showed linear curves that were parallel to the standard curves obtained using the Quantikine kit standards. These results indicate that the Quantikine I-TAC kit can be used to determine relative mass values for naturally occurring I-TAC.

## **PRINCIPLE OF THE ASSAY**

This assay employs the quantitative sandwich enzyme immunoassay technique. A monoclonal antibody specific for I-TAC has been pre-coated onto a microplate. Standards and samples are pipetted into the wells and any I-TAC present is bound by the immobilized antibody. After washing away any unbound substances, an enzyme-linked polyclonal antibody specific for I-TAC is added to the wells. Following a wash to remove any unbound antibody-enzyme reagent, a substrate solution is added to the wells and color develops in proportion to the amount of I-TAC bound in the initial step. The color development is stopped and the intensity of the color is measured.

## **LIMITATIONS OF THE PROCEDURE**

- FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES.
- The kit should not be used beyond the expiration date on the kit label.
- If samples generate values higher than the highest standard, dilute the samples with the appropriate Calibrator Diluent and repeat the assay.
- Any variation in standard diluent, operator, pipetting technique, washing technique, incubation time or temperature, and kit age can cause variation in binding.
- This assay is designed to eliminate interference by soluble receptors, binding proteins, and other factors present in biological samples. Until all factors have been tested, the possibility of interference cannot be excluded.

## REAGENTS

**I-TAC Microplate** (Part 891030) - 96 well polystyrene microplate (12 strips of 8 wells) coated with a mouse monoclonal antibody against I-TAC.

**I-TAC Conjugate** (Part 891031) - 21 mL of polyclonal antibody against I-TAC conjugated to horseradish peroxidase, with preservatives.

**I-TAC Standard** (Part 891032) - 40 ng of recombinant human I-TAC in a buffer with preservative, lyophilized.

**Assay Diluent RD1-68** (Part 895528) - 11 mL of a buffer with preservative.

**Calibrator Diluent RD5-21** (Part 895348) - 21 mL of a buffered protein base with preservative.  
*For cell culture samples.*

**Calibrator Diluent RD6P** (Part 895118) - 21 mL of animal serum with preservative.  
*For serum/plasma samples.*

**Wash Buffer Concentrate** (Part 895003) - 21 mL of a 25-fold concentrated solution of buffered surfactant with preservatives.

**Color Reagent A** (Part 895000) - 12.5 mL of stabilized hydrogen peroxide.

**Color Reagent B** (Part 895001) - 12.5 mL of stabilized chromogen (tetramethylbenzidine).

**Stop Solution** (Part 895032) - 6 mL of 2 N sulfuric acid.

**Plate Covers** - 4 adhesive strips.

## STORAGE

<b>Unopened Kit</b>	Store at 2 - 8° C. Do not use past kit expiration date.	
<b>Opened/ Reconstituted Reagents</b>	Diluted Wash Buffer	May be stored for up to 1 month at 2 - 8° C.*
	Stop Solution	
	Assay Diluent RD1-68	
	Calibrator Diluent RD5-21	
	Calibrator Diluent RD6P	
	Conjugate	
	Unmixed Color Reagent A	
	Unmixed Color Reagent B	
	Standard	
Microplate Wells	Return unused wells to the foil pouch containing the desiccant pack, reseal along entire edge of zip-seal. May be stored for up to 1 month at 2 - 8° C.*	

\*Provided this is within the expiration date of the kit.

## OTHER SUPPLIES REQUIRED

- Microplate reader capable of measuring absorbance at 450 nm, with the correction wavelength set at 540 nm or 570 nm.
- Pipettes and pipette tips.
- Deionized or distilled water.
- Multi-channel pipette, squirt bottle, manifold dispenser, or automated microplate washer.
- 500 mL graduated cylinder.
- **12 mm x 75 mm polypropylene tubes.**

## PRECAUTION

The Stop Solution provided with this kit is an acid solution. Wear eye, hand, face, and clothing protection when using this material.

## SAMPLE COLLECTION AND STORAGE

**Cell culture supernates** - Remove particulates by centrifugation and assay immediately or aliquot and store samples at  $\leq -20^{\circ}$  C. Avoid repeated freeze-thaw cycles.

**Serum** - Use a serum separator tube (SST) and allow samples to clot for 30 minutes before centrifugation for 15 minutes at approximately 1000 x g. Remove serum and assay immediately or aliquot and store samples at  $\leq -20^{\circ}$  C. Avoid repeated freeze-thaw cycles.

**Plasma** - Collect plasma using heparin or EDTA as an anticoagulant. Centrifuge for 15 minutes at 1000 x g within 30 minutes of collection. Assay immediately or aliquot and store samples at  $\leq -20^{\circ}$  C. Avoid repeated freeze-thaw cycles.

**Note:** *Grossly hemolyzed samples are not suitable for use in this assay. Citrate plasma has not been validated for use in this assay. Heparin appears to release erythrocyte-bound I-TAC, resulting in higher measured concentrations than matched EDTA plasma or serum samples. See the Sample Values section.*

## REAGENT PREPARATION

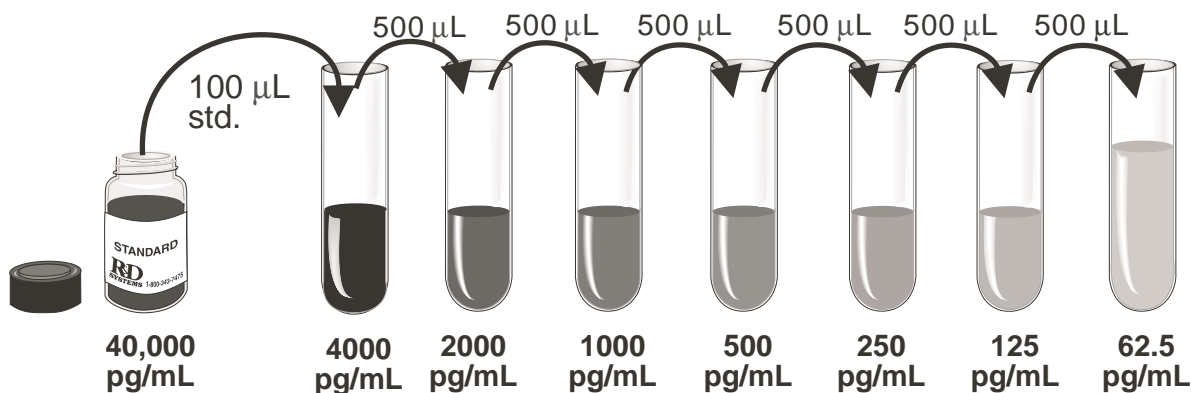
**Bring all reagents to room temperature before use.**

**Wash Buffer** - If crystals have formed in the concentrate, warm to room temperature and mix gently until the crystals have completely dissolved. Dilute 20 mL of Wash Buffer Concentrate into deionized or distilled water to prepare 500 mL of Wash Buffer.

**Substrate Solution** - Color Reagents A and B should be mixed together in equal volumes within 15 minutes of use. Protect from light. 200  $\mu\text{L}$  of the resultant mixture is required per well.

**I-TAC Standard** - Reconstitute the I-TAC Standard with 1 mL of deionized or distilled water. This reconstitution produces a stock solution of 40,000 pg/mL. Mix the standard to ensure complete reconstitution and allow the standard to sit for a minimum of 15 minutes with gentle agitation prior to making dilutions.

**Use polypropylene tubes.** Pipette 900  $\mu\text{L}$  of Calibrator Diluent RD5-21 (*for cell culture supernate samples*) or Calibrator Diluent RD6P (*for serum/plasma samples*) into the 4000 pg/mL tube. Pipette 500  $\mu\text{L}$  of the appropriate Calibrator Diluent into the remaining tubes. Use the stock solution to produce a dilution series (below). Mix each tube thoroughly before the next transfer. The 4000 pg/mL standard serves as the high standard. The appropriate Calibrator Diluent serves as the zero standard (0 pg/mL).



## ASSAY PROCEDURE

**Bring all reagents and samples to room temperature before use. It is recommended that all samples and standards be assayed in duplicate.**

1. Prepare all reagents, working standards and samples as directed in the previous sections.
2. Remove excess microplate strips from the plate frame, return them to the foil pouch containing the desiccant pack, reseal.
3. Add 100  $\mu\text{L}$  of Assay Diluent RD1-68 to each well.
4. Add 100  $\mu\text{L}$  of Standard or sample per well. Cover with the adhesive strip provided. Incubate for 2 hours at room temperature. A plate layout is provided to record standards and samples assayed.
5. Aspirate each well and wash, repeating the process three times for a total of four washes. Wash by filling each well with Wash Buffer (400  $\mu\text{L}$ ) using a squirt bottle, multi-channel pipette, manifold dispenser or autowasher. Complete removal of liquid at each step is essential to good performance. After the last wash, remove any remaining Wash Buffer by aspirating or decanting. Invert the plate and blot it against clean paper towels.
6. Add 200  $\mu\text{L}$  of I-TAC Conjugate to each well. Cover with a new adhesive strip. Incubate for 2 hours at room temperature.
7. Repeat the aspiration/wash as in step 5.
8. Add 200  $\mu\text{L}$  of Substrate Solution to each well. Incubate for 30 minutes at room temperature. **Protect from light.**
9. Add 50  $\mu\text{L}$  of Stop Solution to each well. If color change does not appear uniform, gently tap the plate to ensure thorough mixing.
10. Determine the optical density of each well within 30 minutes, using a microplate reader set to 450 nm. If wavelength correction is available, set to 540 nm or 570 nm. If wavelength correction is not available, subtract readings at 540 nm or 570 nm from the readings at 450 nm. This subtraction will correct for optical imperfections in the plate. Readings made directly at 450 nm without correction may be higher and less accurate.

## ASSAY PROCEDURE SUMMARY

1. Prepare reagents, standards, and samples as instructed.



2. Add 100  $\mu\text{L}$  Assay Diluent RD1-68 to each well.



3. Add 100  $\mu\text{L}$  Standard or sample to each well.  
Incubate 2 hr. RT



4. Aspirate and wash 4 times.



5. Add 200  $\mu\text{L}$  Conjugate to each well.  
Incubate 2 hr. RT



6. Aspirate and wash 4 times.



7. Add 200  $\mu\text{L}$  Substrate Solution to each well.  
Incubate 30 min. RT. **Protect from light.**



8. Add 50  $\mu\text{L}$  Stop Solution to each well.  
Read at 450 nm within 30 min.  
 $\lambda$  correction 540 or 570 nm

# CALCULATION OF RESULTS

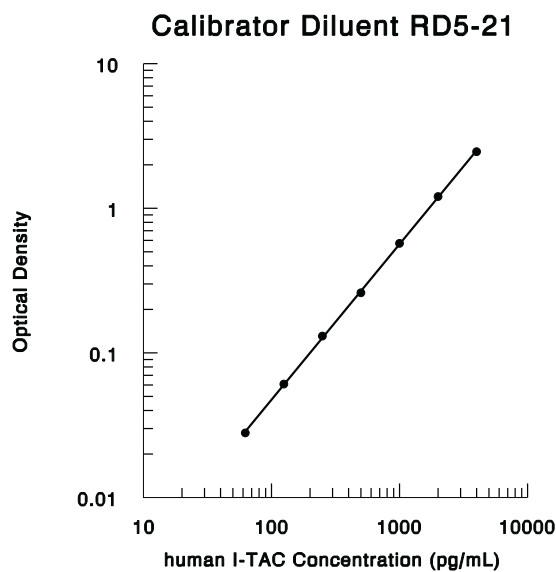
Average the duplicate readings for each standard, control, and sample and subtract the average zero standard optical density.

Plot the optical density for the standards versus the concentration of the standards and draw the best curve. The data can be linearized by using log/log paper and regression analysis may be applied to the log transformation.

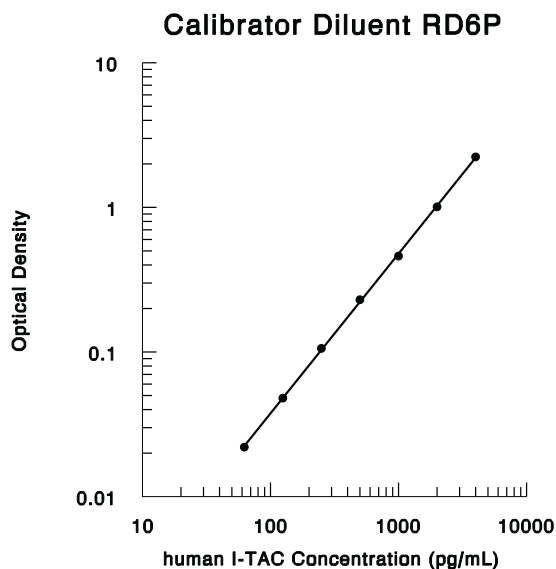
To determine the I-TAC concentration of each sample, first find the absorbance value on the y-axis and extend a horizontal line to the standard curve. At the point of intersection, extend a vertical line to the x-axis and read the corresponding I-TAC concentration. If the samples have been diluted, the concentration read from the standard curve must be multiplied by the dilution factor.

## TYPICAL DATA

These standard curves are provided for demonstration only. A standard curve should be generated for each set of samples assayed.



pg/mL	O.D.	Average	Corrected
0	0.053 0.057 0.079	0.055	-
62.5	0.086 0.114	0.083	0.028
125	0.118 0.178	0.116	0.061
250	0.194 0.309	0.186	0.131
500	0.322 0.622	0.316	0.261
1000	0.634 1.261	0.628	0.573
2000	1.262 2.459	1.262	1.207
4000	2.587	2.523	2.468



pg/mL	O.D.	Average	Corrected
0	0.018 0.019 0.040	0.019	-
62.5	0.042 0.067	0.041	0.022
125	0.067 0.124	0.067	0.048
250	0.125 0.246	0.125	0.106
500	0.251 0.474	0.249	0.230
1000	0.485 1.019	0.480	0.461
2000	1.038 2.195	1.029	1.010
4000	2.312	2.254	2.235

## TECHNICAL HINTS

- Substrate Solution should remain colorless until added to the plate. Keep Substrate Solution protected from light. Substrate Solution should change from colorless to gradations of blue. The color developed in the wells will turn from blue to yellow upon addition of the Stop Solution.
- Stop Solution should be added to the plate in the same order as the Substrate Solution.
- When mixing or reconstituting protein solutions, always avoid foaming.
- To avoid cross-contamination, change pipette tips between additions of each standard level, between sample additions, and between reagent additions. Also, use separate reservoirs for each reagent.
- To ensure accurate results, proper adhesion of plate sealers during incubation steps is necessary.

## PRECISION

### Intra-assay Precision (Precision within an assay)

Three samples of known concentration were tested twenty times on one plate to assess intra-assay precision.

### Inter-assay Precision (Precision between assays)

Three samples of known concentration were tested in forty separate assays to assess inter-assay precision.

### Serum/Plasma Assay

Sample	Intra-assay Precision			Inter-assay Precision		
	1	2	3	1	2	3
n	20	20	20	40	40	40
Mean (pg/mL)	403	1321	2481	437	1403	2644
Standard deviation	19	62.5	147.3	35.1	107	190
CV (%)	4.7	4.7	5.9	8.0	7.6	7.2

### Cell Culture Supernate Assay

Sample	Intra-assay Precision			Inter-assay Precision		
	1	2	3	1	2	3
n	20	20	20	40	40	40
Mean (pg/mL)	353	1179	2191	409	1290	2401
Standard deviation	18	40.6	102.9	35.5	93	179
CV (%)	5.1	3.4	4.7	8.7	7.2	7.5

## RECOVERY

The recovery of I-TAC spiked to levels throughout the range of the assay in various matrices was evaluated.

Sample	Average % Recovery	Range
Cell culture media (n=4)	100	91 - 111%
Serum (n=5)	106	93 - 117%
EDTA plasma (n=5)	98	85 - 112%
Heparin plasma (n=5)	98	90 - 106%

## LINEARITY

To assess the linearity of the assay, samples containing and/or spiked with high concentrations of I-TAC were serially diluted with the appropriate Calibrator Diluent to produce samples with values within the dynamic range of the assay.

		Cell culture supernates (n=5)	Serum (n=5)	Heparin plasma (n=5)	EDTA plasma (n=5)
1:2	Average % of Expected	103	106	103	105
	Range (%)	96 - 109	97 - 111	100 - 107	99 - 108
1:4	Average % of Expected	105	103	106	107
	Range (%)	99 - 113	93 - 108	102 - 109	98 - 115
1:8	Average % of Expected	106	99	107	106
	Range (%)	96 - 111	90 - 104	102 - 111	103 - 111
1:16	Average % of Expected	105	95	99	104
	Range (%)	98 - 111	89 - 100	96 - 102	101 - 112

## SENSITIVITY

Fifty-six assays were evaluated and the minimum detectable dose (MDD) of I-TAC ranged from 3.4 - 39.7 pg/mL. The mean MDD was 13.9 pg/mL.

The minimum detectable dose was determined by adding two standard deviations to the mean optical density value of twenty zero standard replicates and calculating the corresponding concentration.

## CALIBRATION

This immunoassay is calibrated against a highly purified *E. coli*-expressed recombinant human I-TAC produced at R&D Systems.

## SAMPLE VALUES

**Serum/plasma** - Serum and plasma samples were evaluated for the presence of I-TAC in this assay.

**Note:** *Heparin appears to release erythrocyte-bound I-TAC, resulting in higher measured concentrations than matched EDTA plasma or serum samples.*

Sample Type	Mean of Detectable (pg/mL)	% Detectable	Range (pg/mL)
Serum (n=35)	177	17	ND - 290
EDTA plasma (n=35)	136	26	ND - 269
Heparin plasma (n=60)	440	100	100 - 1208

ND = Non-detectable

**Cell culture supernates** - Human peripheral blood mononuclear cells ( $5 \times 10^6$  cells/mL) were cultured in RPMI supplemented with 5% fetal calf serum, 50  $\mu$ M  $\beta$ -mercaptoethanol, 2 mM L-glutamine, 100 U/mL penicillin, and 100  $\mu$ g/mL streptomycin sulfate. The cells were cultured unstimulated or stimulated with 10  $\mu$ g/mL PHA. Aliquots of the culture supernate were removed and assayed for levels of natural I-TAC.

Condition	Day 1 (pg/mL)	Day 5 (pg/mL)
Unstimulated	ND	ND
Stimulated	1655	1088

ND = Non-detectable

THP-1 cells were seeded at  $5 \times 10^5$  cells/mL and grown in RPMI supplemented with 10% fetal calf serum, 50  $\mu$ M  $\beta$ -mercaptoethanol, 2 mM L-glutamine, 100 U/mL penicillin, and 100  $\mu$ g/mL streptomycin sulfate at 37° C and 5% CO<sub>2</sub>. Cells were stimulated with 1  $\mu$ g/mL rhIFN- $\gamma$  for 8 hours, 1  $\mu$ g/mL of LPS was added, and the cells were incubated for an additional 8 hours. An aliquot was removed, assayed for levels of natural I-TAC, and measured 2348 pg/mL.

## SPECIFICITY

This assay recognizes recombinant and natural human I-TAC. The factors listed below were prepared at 50 ng/mL in Calibrator Diluent RD5-21 and Calibrator Diluent RD6P and assayed for cross-reactivity. Preparations of the following factors at 50 ng/mL in a mid-range recombinant human I-TAC control were assayed for interference. No significant cross-reactivity or interference was observed.

### **Recombinant human:**

ENA-78  
BLC/BCA-1  
GCP-2  
GRO $\alpha$   
GRO $\beta$   
GRO $\gamma$   
IFN- $\gamma$

IL-8  
IL-8, endothelial cell-derived  
IP-10  
MIG  
NAP-2  
SDF-1 $\alpha$   
SDF-1 $\beta$

### **Recombinant mouse:**

BLC/BCA-1  
CRG-2 (IP-10)  
GCP-2  
I-TAC  
KC  
MIG  
SDF-1 $\alpha$

### **Recombinant porcine:**

IL-8

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# PLATE LAYOUT

Use this plate layout as a record of standards and samples assayed.

1											
2											
3											
4											
5											
6											
7											
8											
9											
10											
11											
12											
	<b>A</b>	<b>B</b>	<b>C</b>	<b>D</b>	<b>E</b>	<b>F</b>	<b>G</b>	<b>H</b>			