



PRODUCT INFORMATION & MANUAL

Human CXCL1/GRO α Valukine™ ELISA

Catalog Number: VAL139

For the quantitative determination of natural and recombinant
Human CXCL1/GRO α concentrations

For research use only.
Not for diagnostic or therapeutic procedures.

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Please refer to the kit label for expiry date.
Novus kits are guaranteed for 3 months from date of receipt

Version 202308.4

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I. BACKGROUND

CXCL1, also known as KC, GRO alpha, and CINC-1, is an approximately 8 kDa proinflammatory chemokine that plays a key role in neutrophil migration and activation (1). Mature human CXCL1 shares 64% and 67% aa sequence identity with mouse and rat CXCL1, respectively (2). It is produced by many cell types in inflammatory sites and during chronic inflammatory diseases (1). CXCL1 can associate into bioactive dimers and primarily signals through CXCR2/IL-8 RB but can also bind with lower affinity to CXCR2/IL-8 RA (3-5). It induces neutrophil migration, extravasation, respiratory burst, and degranulation and also induces T cells to produce proinflammatory IL-17 (4, 6, 7). CXCL1 additionally binds to Syndecan-1 on epithelial cells which acts as a sink for CXCL1 activity until Syndecan-1 cleavage by MMP-7 (8). CXCL1 is up-regulated in spinal cord astrocytes by inflammatory stimuli or tumor cell injection, and it exacerbates pain sensation by potentiating excitatory NMDA neurotransmission (9, 10). In the circulatory system, CXCL1 interacts with CXCR2 on endothelial cells to promote lymphatic tube formation and angiogenesis (11,12). It promotes the hypertrophic differentiation of chondrocytes resulting in cartilage matrix deposition, calcification, and remodeling (13). It interacts with both CXCR1 and CXCR2 on adipose stromal cells and promotes their recruitment to prostate tumors in obese patients (14). It also binds CXCR2 on ovarian cancer cells, leading to cleavage of cell surface HB-EGF, transactivation of EGF R, and cell proliferation (15).

II. OVERVIEW

A. PRINCIPLE OF THE ASSAY

This assay employs the quantitative sandwich enzyme immunoassay technique. An antibody specific for human CXCL1/GRO α has been pre-coated onto a microplate. Standards and samples are pipetted into the wells and any CXCL1/GRO α present is bound by the immobilized antibody. After washing away any unbound substances, a biotinylated detection antibody specific for human CXCL1/GRO α are pipetted into the wells. After washing away any unbound substances, Streptavidin-HRP are pipetted into the wells. Following a wash to remove any unbound reagent, TMB substrate solution (Chromogenic agent) is added to the wells and color develops in proportion to the amount of CXCL1/GRO α bound in the initial step. The color development is stopped, and the intensity of the color is measured.

B. LIMITATIONS OF THE PROCEDURE

- **FOR RESEARCH USE ONLY NOT FOR USE IN DIAGNOSTIC PROCEDURES.**
- This kit is suitable for cell culture supernate and human serum.
- The kit should not be used beyond the expiration date on the kit label.
- Do not mix or substitute reagents with those from other lots or sources.
- If samples generate values higher than the highest standard, dilute the samples with Calibrator Diluent (1 \times) and repeat the assay.
- Any variation in operator, pipetting technique, washing technique, incubation time or temperature, and kit age can cause variation in binding.

III. ADVANTAGES

A. PRECISION

Intra-assay Precision (Precision within an assay)

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

Inter-assay Precision (Precision between assays)

Three samples were tested in twenty separate assays to assess inter-assay precision.

	Intra-assay Precision			Inter-assay Precision		
Sample	1	2	3	1	2	3
Mean (pg/mL)	95.7	303.0	926.7	93.5	298.4	942.9
Standard Deviation	4.3	9.7	61.7	5.3	11.0	58.7
CV%	4.5	3.2	6.7	5.7	3.7	6.2

B. RECOVERY

The recovery of human CXCL1/GRO α spiked to different levels throughout the range of the assay in cell culture media was evaluated. The recovery ranged from 111.7-124.8% with an average of 115.0%.

The recovery of Human CXCL1/GRO α spiked to different levels throughout the range of the assay in human serum was evaluated. The recovery ranged from 81.8-98.1% with an average of 91.7%.

C. SENSITIVITY

The minimum detectable dose (MDD) of human CXCL1/GRO α is typically less than 2.03 pg/mL.

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

D. CALIBRATION

This immunoassay is calibrated against a highly purified *E. Coli*-expressed recombinant human CXCL1/GRO α produced at R&D Systems.

E. LINEARITY

To assess the linearity of the assay, different samples were containing or spiked with high concentrations of human CXCL1/GRO α and diluted with Calibrator Diluent (1 \times) to produce samples with values within the dynamic range of the assay.

Dilution	Average % of Expected	Range (%)
1:2	99.1	89.8-108.6
1:4	103.8	91.1-116.2
1:8	107.1	88.6-121.2
1:16	106.9	82.9-124.6

F. SAMPLE VALUES

Serum - Four human serum samples were evaluated for the presence of human CXCL1/GRO α in this assay. All samples measured ranged from 91.6 to 187.6 pg/mL with an average of 134.6 pg/mL.

G. SPECIFICITY

The following factors prepared at 50 ng/mL were assayed and exhibited no cross-reactivity or interference.

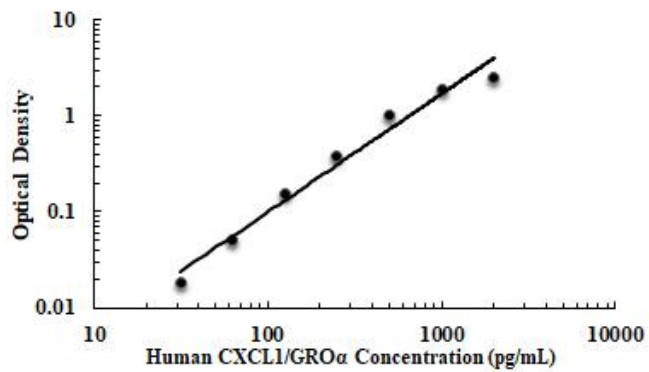
Recombinant human	Recombinant Mouse
ENA-78	KC
GRO β	MIP-1 α
IL-8	MIP-1 β
MCP-1	MIP-2
MIP-1 α	
MIP-1 β	
NAP-2	Recombinant Rat
RANTES	CINC-1

A sample containing 50 ng/mL of recombinant human GRO γ measured 0.144 ng/mL (0.3% cross reactivity)

IV. EXPERIMENT

EXAMPLE STANDARD

The standard curve is provided for demonstration only. A standard curve should be generated for each set of samples assayed.



pg/mL	OD	Average	Corrected
0	0.078 0.081	0.080	-
31.3	0.097 0.099	0.098	0.018
62.5	0.129 0.132	0.131	0.051
125	0.228 0.238	0.234	0.154
250	0.476 0.458	0.467	0.387
500	1.108 1.094	1.101	1.021
1000	1.972 1.973	1.973	1.893
2000	2.560 2.606	2.583	2.503

V. KIT COMPONENTS AND STORAGE

A. MATERIALS PROVIDED

Parts	Description	Size
Human CXCL1/GRO α Microplate	96 well polystyrene microplate (12 strips of 8 wells) coated with a Mouse antibody against Human CXCL1/GRO α .	1 plate
Human CXCL1/GRO α Standard	Recombinant Human CXCL1/GRO α in a buffered protein base; lyophilized. Refer to the vial label for reconstitution volume.	2 vials
Human CXCL1/GRO α Detection Antibody	Biotinylated CXCL1/GRO α antibody, lyophilized. Refer to the vial label for reconstitution volume.	1 vial
Calibrator Diluent (1 \times)	Buffered diluent used to dilute standard and samples.	2 vials
Detection Antibody Diluent (4 \times)	Concentrated buffered diluent used to dilute Detection Antibody.	1 vial
Streptavidin-HRP B (40 \times)	40 \times Streptavidin conjugated to horseradish peroxidase.	1 vial
Reagent Diluent (10 \times)	A 10 \times concentrated buffered protein base used to dilute Detection Antibody and HRP.	1 vial
Wash Buffer Concentrate (25 \times)	A 25 \times concentrated solution of buffered surfactant with preservatives.	1 vial
TMB Substrate	TMB ELISA Substrate Solution	1 vial
Stop Solution	2 N sulfuric acid.	1 vial
Plate Sealers	Adhesive strip.	3 strips

B. STORAGE

Unopened Kit	Store at 2-8 °C. Do not use past kit expiration date.	
Opened/ Reconstituted Reagents	Streptavidin-HRP B	May be stored for up to 1 month at 2-8 °C.*
	Diluted Wash Solution	
	TMB Substrate	
	Stop Solution	
	Standard	Prepare fresh for each assay.
	Detection Antibody	Aliquot and store for up to 1 month at -20 °C in a manual defrost freezer. *
	Reagent Diluent (10×)	May be stored for up to 1 month at 2-8 °C.* Use and discard diluted Reagent Diluent (1×). Prepare fresh for each assay.
	Calibrator Diluent (1×)	May be stored for up to 1 month at 2-8 °C.*
	Detection Antibody Diluent (4×)	May be stored for up to 1 month at 2-8 °C.* Use and discard diluted Detection Antibody Diluent (1×). Prepare fresh for each assay.
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.

C. 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.
- ◆ Squirt bottle, manifold dispenser, or automated microplate washer.
- ◆ 500 mL graduated cylinder.

D. PRECAUTION

- ◆ Some components in this kit contain a preservative which may cause an allergic skin reaction. Avoid breathing mist.
- ◆ The Stop Solution provided with this kit is an acid solution. Wear protective gloves, clothing, eye, and face protection. Wash hands thoroughly after handling.

VI. PREPARATION

A. SAMPLE COLLECTION AND STORAGE

Cell Culture Supernates - Remove particulates by centrifugation and assay immediately or aliquot and store samples at $\leq -20^{\circ}\text{C}$. Avoid repeated freeze-thaw cycles. Samples may require dilution with Calibrator Diluent (1 \times).

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

B. SAMPLE PREPARATION

Serum samples recommend a 2-fold dilution. A suggested 2-fold dilution is 100 μL of sample + 100 μL of Calibrator Diluent (1 \times). Optimal dilutions should be determined by the end user.

C. REAGENT PREPARATION

Note: *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 (25 \times) into deionized or distilled water to prepare 500 mL of Wash Buffer.

Detection Antibody Diluent (1 \times) - Use deionized or distilled water to prepare Detection Antibody Diluent (1 \times).

Reagent Diluent (1 \times) - Use deionized or distilled water to prepare Reagent Diluent (1 \times).

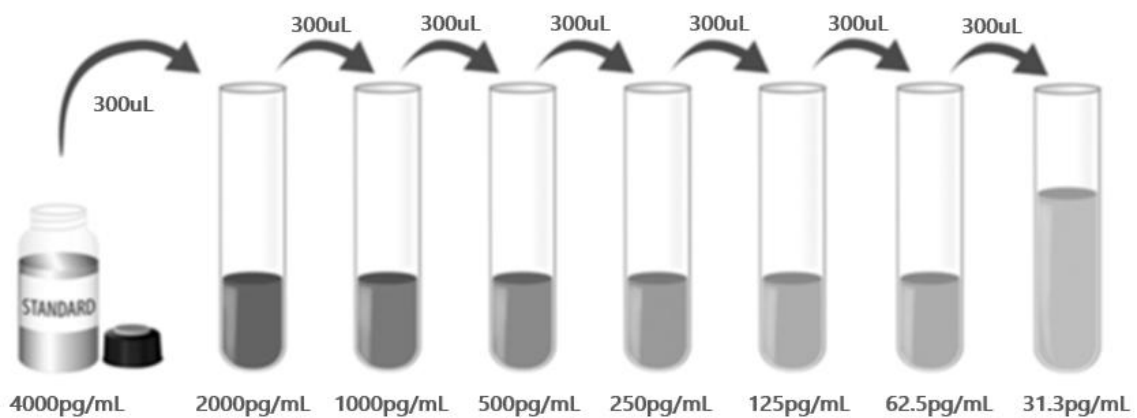
Detection Antibody (1 \times) - **Centrifuge briefly before opening. Reconstitution volume refer to vial label to prepare Detection Antibody (100 \times).** Allow the Detection Antibody to sit for a minimum of 15 minutes with gentle agitation prior to making dilutions. Aliquot and store if needed. Dilute to Detection Antibody (1 \times) with Detection Antibody Diluent (1 \times). Prepare at least 15 minutes prior to use.

Streptavidin-HRP B (1 \times) - **Centrifuge briefly before opening.** Dilute to the working concentration specified on the vial label using Reagent Diluent (1 \times).

Human CXCL1/GRO α Standard - Centrifuge briefly before opening. Refer to the vial label for the reconstitution volume*. This reconstitution produces a stock solution of 4000 pg/mL. Allow the standard to sit for a minimum of 15 minutes with gentle agitation prior to making dilutions.

*If you have any question, please seek help from our Technical Support.

Pipette 300 μ L of the appropriate Calibrator Diluent (1 \times) into each tube. Use the stock solution to produce a dilution series (below). Mix each tube thoroughly before the next transfer. The 2000 pg/mL standard serves as the high standard. The Calibrator Diluent (1 \times) serves as the zero standard (0 pg/mL).



D. TECHNICAL HINTS

- 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.
- It is recommended that the samples be pipetted within 15 minutes.
- To ensure accurate results, proper adhesion of plate sealers during incubation steps is necessary.
- TMB substrate should remain colorless until added to the plate. Keep TMB Substrate Solution protected from light. TMB Substrate Solution should change from colorless to gradations of blue.
- Stop Solution should be added to the plate in the same order as the TMB substrate. The color developed in the wells will turn from blue to yellow upon addition of the Stop Solution. Wells that are green in color indicate that the Stop Solution has not mixed thoroughly with the TMB substrate.

VII. ASSAY PROCEDURE

Note: 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 and working standards 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, and reseal.
3. Add 100 μ L of standard, or prepared sample per well. Cover with the adhesive strip provided. **Incubate for 2 hours at room temperature.** A plate layout is provided for a record of standards and samples assayed. (Samples may require dilution. See Sample Preparation section.)
4. Aspirate each well and wash, repeating the process three times for a total of four washes. Wash by filling each well with Wash Solution (400 μ L) using a squirt bottle, 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.
5. Add 100 μ L of the Detection Antibody (1 \times) diluted in Detection Antibody Diluent, to each well. Cover with a new adhesive strip and **incubate 2 hours at room temperature.**
6. Repeat the aspiration/wash as in step 4.
7. Add 100 μ L of the working dilution of Streptavidin-HRP B to each well. Cover the plate and **incubate for 20 minutes at room temperature. Avoid placing the plate in direct light.**
8. Repeat the aspiration/wash as in step 4.
9. Add 100 μ L of TMB Substrate to each well. **Incubate for 20 minutes at room temperature. Avoid placing the plate in direct light.**
10. Add 50 μ L of Stop Solution to each well. Gently tap the plate to ensure thorough mixing.
11. Determine the optical density of each well within 10 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.

12. **CALCULATION OF RESULTS**

Average the duplicate readings for each standard, control, and sample and subtract the average zero standard optical density (O.D.). Create a standard curve by reducing the data using computer software capable of generating a four parameter logistic (4-PL) curve-fit. As an alternative, construct a standard curve by plotting the mean absorbance for each standard on the y-axis against the concentration on the x-axis and draw a best fit curve through the points on the graph. The data may be linearized by plotting the log of the human CXCL1/GRO α concentrations versus the log of the O.D. and the best fit line can be determined by regression analysis. This procedure will produce an adequate but less precise fit of the data.

If samples have been diluted, the concentration read from the standard curve must be multiplied by the dilution factor.

VIII. REFERENCES

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PLATE LAYOUT

Use this plate layout to record standards and samples assayed.

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



产品信息及操作手册

人 CXCL1/GRO α Valukine™ ELISA 试剂盒

目录号: **VAL139**

适用于定量检测天然和重组人 CXCL1/GRO α 的浓度

科研专用, 不可用于临床诊断

Bio-Techne China Co. Ltd

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有效期详见试剂盒包装标签

Novus 试剂盒确保在你收货日期 3 个月内有效

版本号 202308.4

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I. 背景

CXCL1, 也称为KC、GRO α 和CINC-1, 是一种约8 kDa的促炎性趋化因子, 在中性粒细胞的迁移和激活中起着关键作用 (1)。成熟人CXCL1与小鼠和大鼠CXCL1的aa序列同源性分别为64%和67% (2)。它由炎症部位和慢性炎症性疾病中的多种细胞类型产生 (1)。CXCL1可结合成生物活性二聚体, 主要通过CXCR2/IL-8 RB传递信号, 但也可与CXCR2/IL-8 RA低亲和力结合 (3-5)。诱导中性粒细胞迁移、外渗、呼吸爆发和脱颗粒, 并诱导T细胞产生促炎性IL-17 (4, 6, 7)。CXCL1还与上皮细胞上的Syndecan-1结合, 作为CXCL1活动的接收器, 直到Syndecan-1被MMP-7分解 (8)。通过炎症刺激或肿瘤细胞注射CXCL1在脊髓星形胶质细胞中上调, 并通过增强兴奋性NMDA神经传递而加剧疼痛感 (9, 10)。在循环系统中, CXCL1与内皮细胞上的CXCR2相互作用, 以促进淋巴管的形成和血管生成 (11, 12)。它促进软骨细胞的肥大分化, 导致软骨基质沉积、钙化和重塑 (13)。它与脂肪间质细胞上的CXCR1和CXCR2相互作用, 并促进肥胖患者前列腺肿瘤的募集 (14)。它还与卵巢癌细胞上的CXCR2结合, 导致细胞表面HB-EGF断裂、EGF-R反式激活和细胞增殖 (15)。

II. 概述

A. 检测原理

本实验采用双抗体夹心ELISA法。抗人CXCL1/GRO α 抗体包被于微孔板上，样品和标准品中的人CXCL1/GRO α 会与固定在板上的抗体结合，游离的成分被洗去；接着加入生物素化的抗人CXCL1/GRO α 检测抗体进行孵育，洗涤去除未结合的物质后，加入链霉亲和素标记的辣根过氧化物酶（Streptavidin-HRP）孵育。洗涤去除未结合的试剂后，加入TMB底物溶液（显色剂）。溶液颜色与结合的目标蛋白成正比；加入终止液；用酶标仪测定吸光度。

B. 检测局限

- ◆ 仅供科研使用，不可用于体外诊断；
- ◆ 该试剂盒适用于细胞培养上清样本和人血清样本；
- ◆ 请在试剂盒有效期内使用；
- ◆ 不同试剂盒及不同批号试剂盒的组分不能混用；
- ◆ 样本值若大于标准曲线的最高值，应将样本用标准品稀释液（1 \times ）稀释后重新检测；
- ◆ 检测结果的不同可由多种因素引起，包括实验人员的操作、移液器的使用方式、洗板技术、反应时间或温度、试剂盒的效期等。

III. 优势

A. 精确度

板内精确度（同一板内不同孔间的精确度）

已知浓度的三个样本，在同一板内分别检测20次，以确定板内精确度。

板间精确度（不同板之间的精确度）

已知浓度的三个样本，在不同板中分别检测20次，以确定板间精确度。

样本	板内精确度			板间精确度		
	1	2	3	1	2	3
平均值 (pg/mL)	95.7	303.0	926.7	93.5	298.4	942.9
标准差	4.3	9.7	61.7	5.3	11.0	58.7
CV%	4.5	3.2	6.7	5.7	3.7	6.2

B. 回收率

在细胞培养基样本中掺入检测范围内不同水平的人CXCL1/GRO α ，测定其回收率。回收率范围在111.7-124.8%，平均回收率在115.0%。

在人血清样本中掺入检测范围内不同水平的人CXCL1/GRO α ，测定其回收率。回收率范围在81.8-98.1%，平均回收率在91.7%。

C. 灵敏度

人CXCL1/GRO α 的最低可测剂量（MDD）一般小于2.03 pg/mL。

MDD是根据20个重复的零标准品孔的吸光度值的平均值加两倍标准差计算得到的相对应浓度。

D. 校正

此ELISA试剂盒经由R&D Systems生产的大肠杆菌表达的高纯度重组人CXCL1/GRO α 蛋白所校正。

E. 线性

不同的样本中含有或掺入高浓度的人CXCL1/GRO α ，然后用标准品稀释液（1 \times ）将样本

稀释到检测范围内，测定其线性。

稀释倍数	平均值(%)	范围 (%)
1:2	99.1	89.8-108.6
1:4	103.8	91.1-116.2
1:8	107.1	88.6-121.2
1:16	106.9	82.9-124.6

F. 样本预值

血清样本 - 使用本试剂盒检测了4份人血清样本中CXCL1/GRO α 的水平。4份样本的检测值范围为91.6-187.6 pg/mL，平均值为134.6 pg/mL。

G. 特异性

将以下因子配制成50ng/mL的浓度来检测没有观察到明显的交叉反应或干扰。

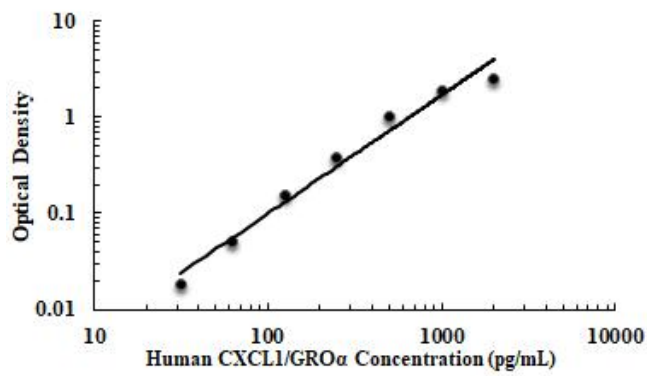
Recombinant human	Recombinant Mouse
ENA-78	KC
GRO β	MIP-1 α
IL-8	MIP-1 β
MCP-1	MIP-2
MIP-1 α	
MIP-1 β	
NAP-2	Recombinant Rat
RANTES	CINC-1

含有50 ng/mL重组人GRO γ 的样品测量值为0.144 ng/mL（0.3%交叉反应）。

IV. 实验

标准曲线实例

该标准曲线数据仅供参考，每次实验应绘制其对应的标准曲线。



pg/mL	OD	Average	Corrected
0	0.078 0.081	0.080	-
31.3	0.097 0.099	0.098	0.018
62.5	0.129 0.132	0.131	0.051
125	0.228 0.238	0.234	0.154
250	0.476 0.458	0.467	0.387
500	1.108 1.094	1.101	1.021
1000	1.972 1.973	1.973	1.893
2000	2.560 2.606	2.583	2.503

V. 试剂盒组成及储存

A. 试剂盒组成

组成	描述	规格
Human CXCL1/GRO α Microplate	包被小鼠抗人 CXCL1/GRO α 抗体的 96 孔聚苯乙烯板, 8 孔 \times 12 条	1 块板
Human CXCL1/GRO α Standard	标准品 (冻干粉), 参考瓶身标签进行重溶	2 瓶
Human CXCL1/GRO α Detection antibody	生物素化的 CXCL1/GRO α 检测抗体, 冻干粉, 参考瓶身标签进行重溶	1 瓶
Streptavidin-HRP B (40 \times)	40 \times 浓缩的链霉亲和素标记的 HRP	1 瓶
Reagent Diluent (10 \times)	浓缩的试剂稀释液 (10 \times) 用于稀释检测抗体和 HRP	1 瓶
Calibrator Diluent (1 \times)	标准品稀释液 (1 \times) 用于稀释标准品和样本	2 瓶
Detection Antibody Diluent (4 \times)	浓缩的检测抗体稀释液 (4 \times) 用于稀释检测抗体	1 瓶
Wash Buffer Concentrate (25 \times)	浓缩洗涤缓冲液 (25 \times)	1 瓶
TMB Substrate	TMB ELISA 底物溶液	1 瓶
Stop Solution	终止液	1 瓶
Plate Sealers	封板膜	3 张

B. 试剂盒储存

未开封试剂盒	2-8℃储存；请在试剂盒有效期内使用	
已打开，稀释或重溶的试剂	链霉亲和素-HRP B	2-8℃储存，最多 30 天*
	洗涤缓冲液（1×）	
	TMB 底物溶液	
	终止液	
	标准品	使用时新鲜配制*
	检测抗体	分装，-20℃储存，最多 30 天*
	标准品稀释液（1×）	2-8℃储存，最多 30 天*
	检测抗体稀释液（4×）	2-8℃储存，最多 30 天* 请每次使用新鲜配制的 1×检测抗体稀释液，多余的丢弃
	试剂稀释液（10×）	2-8℃储存，最多 30 天* 请每次使用新鲜配制的 1×试剂稀释液，多余的丢弃
	包被的微孔板条	将未用的板条放回带有干燥剂的铝箔袋内，密封：2-8℃储存，最多 30 天*

*必须在试剂盒有效期内

C. 实验所需自备试验器材

- ◆ 酶标仪（可测量450 nm检测波长的吸收值及540 nm或570 nm校正波长的吸收值）
- ◆ 高精度加液器及一次性吸头
- ◆ 蒸馏水或去离子水
- ◆ 洗瓶（喷瓶）、多通道洗板器或自动洗板机
- ◆ 500 mL量筒

D. 注意事项

- ◆ 试剂盒中的一些组分含有防腐剂，可能引起皮肤过敏反应，避免吸入。
- ◆ 试剂盒中的终止液是酸性溶液，使用时请做好眼睛、手、面部及衣服的保护。使用后请彻底洗手。

VI. 实验前准备

A. 样品收集及储存

细胞培养上清液：颗粒物应离心去除；立刻检测样本。样本收集后若不及时检测，需按一次使用量分装，冻存于-20℃冰箱内，避免反复冻融。样本可能需要用标准品稀释液（1×）稀释。

血清样本：用血清分离管(SST)分离血清。使血样室温凝集30分钟，然后1000 × g离心15分钟。吸取血清样本之后即刻用于检测，或者分装，-20℃贮存备用。避免反复冻融。

B. 样本准备工作

血清样本建议用标准品稀释液（1×）2倍稀释后进行检测，例如：100 μL样本+100 μL标准品稀释液（1×）。最佳稀释度应由最终用户确定。

C. 检测前准备工作

使用前请将所有试剂放置于室温

洗涤液：从冰箱中取出的浓缩洗涤液可能有结晶，属于正常现象；放置室温，轻摇混匀，待结晶完全溶解后再配制洗涤液。可将20 mL浓缩洗涤液用蒸馏水或去离子水稀释配制成500 mL工作浓度的洗涤液。未用完的放回4℃。

检测抗体稀释液（1×）：使用蒸馏水或去离子水稀释配制成检测抗体稀释液（1×）。

试剂稀释液（1×）：使用蒸馏水或去离子水稀释配制成试剂稀释液（1×）。

检测抗体（1×）：开盖前请瞬时离心。参考检测抗体瓶标签重溶冻干粉，制备检测抗体（100×）。轻轻震荡至少15分钟，其充分溶解。如有需要分装保存。用检测抗体稀释液（1×）稀释至检测抗体（1×），至少在使用前15分钟准备。

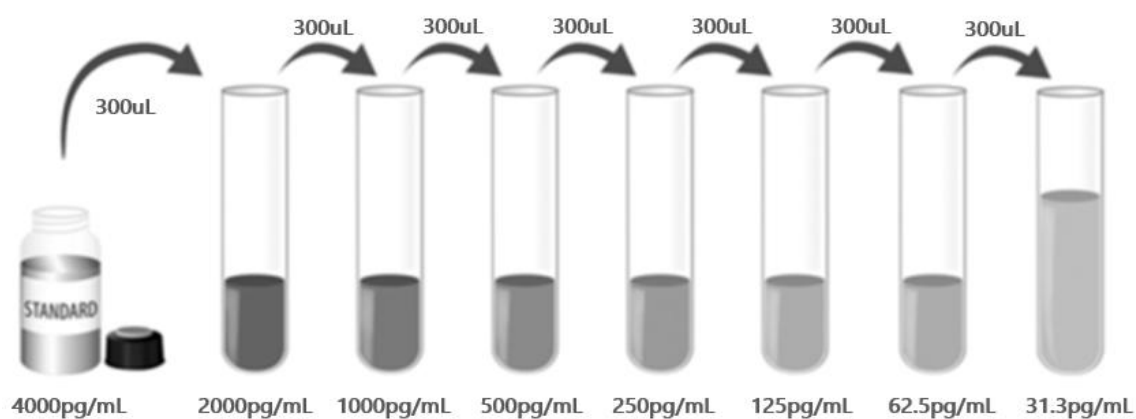
链霉亲和素- HRP B：开盖前请瞬时离心。用试剂稀释液（1×）将链霉亲和素- HRP B（40×）稀释至工作浓度链霉亲和素- HRP B（1×）。

CXCL1/GRO α 标准品：开盖前请瞬时离心。冻干标准品的重溶体积请参考瓶身标签，得到浓度为4000 pg/mL 标准品母液。轻轻震荡至少15分钟，使其充分溶解。

*如有疑问，请咨询我们的技术支持。

向各稀释管中加入300 μL标准品稀释液（1×）。将标准品母液参照下图做系列稀释，每管须充分混匀后再移液到下一管。2000 pg/mL管作标准曲线最高点，标准品稀释液（1×）

可用作标准品零点（0 pg/mL）。



D. 技术小提示

- ◆ 当混合或重溶蛋白液时，尽量避免起沫；
- ◆ 为了避免交叉污染，配制不同浓度标准品、上样、加不同试剂都需要更换枪头。另外不同试剂请分别使用不同的移液槽；
- ◆ 建议15分钟内完成一块板的上样；
- ◆ 每次孵育时，正确使用封板膜可保证结果的准确性；
- ◆ TMB底物溶液在上板前应为无色，请避光保存；加入微孔板后，将由无色变成不同深度的蓝色；
- ◆ 终止液上板顺序应同TMB底物溶液上板顺序一致；加入终止液后，孔内颜色由蓝变黄；若孔内有绿色，则表明孔内液体未混匀，请充分混合。

VII. 操作步骤

使用前请将所有试剂和样本放置于室温，建议所有的实验样本和标准品做复孔检测

1. 按照上一节的说明，准备好所有需要的试剂和标准品；
2. 从已平衡至室温的密封袋中取出微孔板，未用的板条请放回铝箔袋内，重新封口；
3. 分别将不同浓度标准品，实验样本加入相应孔中，每孔100 μL 。用封板膜封住反应孔，**室温孵育2小时**。说明书提供了一张96孔模板图，可用于记录标准品和试验样本的板内位置；（样本需要稀释，详情参见样本制备部分。）
4. 将板内液体吸去，使用洗瓶、多通道洗板器或自动洗板机洗板。每孔加洗涤液400 μL ，然后将板内洗涤液吸去。重复操作3次，共洗4次。每次洗板尽量吸去残留液体会有助于得到好的实验结果。最后一次洗板结束，请将板内所有液体吸干或将板倒置，在吸水纸拍干所有残留液体；
5. 在每个微孔内加入100 μL 配制好的检测抗体（1 \times ）。用封板膜封住反应孔，**室温孵育2小时**；
6. 重复第4步洗板操作；
7. 在每个微孔内加入100 μL 稀释好的链霉亲和素-HRP B工作液。用封板膜封住反应孔，**室温孵育20分钟，注意避光**；
8. 重复第4步洗板操作；
9. 在每个微孔内加入100 μL TMB底物溶液，**室温孵育20分钟，注意避光**；
10. 在每个微孔内加入50 μL 终止液，请轻拍微孔板，使溶液混合均匀；
11. 加入终止液后10分钟内，使用酶标仪测量450 nm的吸光度值，设定540 nm或570 nm作为校正波长。如果波长校正不可用，以450 nm的读数减去540 nm或570 nm的读数。这种减法将校正酶标板上的光学缺陷。没有校正而直接在450 nm处进行的读数可能会更高且更不准确；
12. **计算结果**：将每个标准品和样品的复孔吸光值取平均值，然后减去零标准品平均OD值（O.D.），使用计算机软件作四参数逻辑（4-PL）曲线拟合创建标准曲线。另一替代方法是，通过绘制y轴上每个标准品的平均吸光值与x轴上的浓度来构建标准曲线，并通过图上的点绘制最佳拟合曲线。数据可以通过绘制人CXCL1/GRO α 浓度的对数与O.D.的对数来线性化，并且最佳拟合线可以通过回归分析来确定。该程序将产生足够但不太精确的数据拟合。

如果样品被稀释，从标准曲线读取的浓度必须乘以稀释倍数。

VIII. 参考文献

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96 孔模板图

请使用 96 孔模板图来记录标准品及样本在板内的位置

