

Human IL-29 Sandwich ELISA Kit Datasheet

Please read it entirely before use

Catalogue Number: KE00241

Size: 96T

Sensitivity: 3.3 pg/mL

Range: 62.5 -4000 pg/mL

Usage: For the quantitative detection of human IL-29 concentrations in serum, plasma, cell culture supernatant.

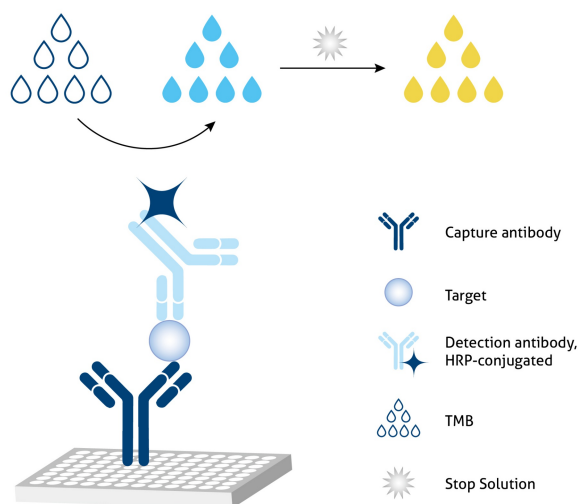
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1. Background

Interleukin-29 (IL-29) is a cytokine belonging to the Type III interferon family, which has another two subfamilies IL-28A and IL-28B. They are also known as IFN- λ 1, IFN- λ 2 and IFN- λ 3, respectively. IL-29 is produced predominantly by maturing dendritic cells and macrophages. IL-29 plays an important role in the immune response against pathogens and especially against viruses by mechanisms similar to type I interferons. IL-29 receptor signals through JAK-STAT pathways leading to activated expression of interferon-stimulated genes and production of antiviral proteins.

2. Principle



Sandwich ELISA structure (Detection antibody labeled with HRP)

A capture antibody is pre-coated onto the bottom of wells which binds to analyte of interest. A detection antibody labeled with HRP also binds to the analyte. TMB acts as the HRP substrate and the solution color will change from colorless to blue. A stop solution containing sulfuric acid turns solution yellow. The color intensity is proportional to the quantity of bound protein which is measurable at 450 nm with the correction wavelength set at 630 nm.

3. Required Materials

- 3.1 A microplate reader capable of measuring absorbance at 450 nm with the correction wavelength set at 630 nm.
- 3.2 Calibrated, adjustable precision pipettes and disposable plastic tips. A manifold multi-channel pipette is recommended for large assays.
- 3.3 Plate washer: automated or manual.
- 3.4 Absorbent paper towels.
- 3.5 Glass or plastic tubes to prepare standard and sample dilutions.
- 3.6 Beakers and graduated cylinders.
- 3.7 Log-log or semi-log graph paper or computer and software for ELISA data analysis. A four-parameter logistic (4-PL) curve-fit is recommended.

4. Kit Components and Storage

| | | |
|--|-----------|---|
| Microplate - antibody coated 96-well microplate (8 well × 12 strips) | 1 plate | Unopened Kit: Store at 2-8°C for 6 months or -20°C for 12 months. Opened Kit: All reagents stored at 2-8°C for 7 days. Please use a new standard for each assay. |
| Protein standard - 8000 pg/bottle; lyophilized | 2 bottles | |
| Detection antibody, HRP-conjugated (100×) - 120 µL/vial* | 1 vial | |
| Sample Diluent PT 4B1 - 30 mL/bottle. For human serum and plasma | 1 bottle | |
| Sample Diluent PT 1-ef - 30 mL/bottle. For cell culture supernatant | 1 bottle | |
| Detection Diluent - 30 mL/bottle | 1 bottle | |
| Wash Buffer Concentrate (20×) - 30 mL/bottle | 1 bottle | |
| Tetramethylbenzidine Substrate (TMB) - 12 mL/bottle | 1 bottle | |
| Stop Solution - 12 mL/bottle | 1 bottle | |
| Plate Cover Seals | 4 pieces | |

* Centrifugation immediately before use

5. Safety Notes

- 5.1 Avoid any skin and eye contact with Stop Solution and TMB. In case of contact, wash thoroughly with water.
- 5.2 Do not use the kit after the expiration date.
- 5.3 Do not mix or substitute reagents or materials from other kit lots or other sources.
- 5.4 Be sure to wear protective equipment such as gloves, masks and goggles during the experiment.
- 5.5 When using an automated plate washer, adding a 30 second soak period following the addition of Wash Buffer to improve assay precision

6. Sample Collection and Storage

- 6.1 Serum: Allow blood samples to clot for 30 minutes, followed by centrifugation for 15 minutes at 1000xg. Clear serum can be assayed immediately or aliquoted and stored at -20°C. Avoid repeated freeze-thaw cycles.
- 6.2 Plasma: Use EDTA, heparin, or citrate as an anticoagulant for plasma collection. Centrifuge for 15 minutes at 1000xg within 30 minutes of collection. The plasma can be assayed immediately or aliquoted and stored at -20°C. Avoid repeated freeze-thaw cycles.
- 6.3 Cell Culture Supernatant: Remove particulates by centrifugation for 5 minutes at 500xg and assay immediately or aliquot and store samples at ≤ -20°C. Avoid repeated freeze-thaw cycles.

7. Regent Preparation

7.1 Wash Buffer (1X): If crystals have formed in the concentrate, warm to room temperature and mix gently until the crystals have completely dissolved. Add 30 mL of Wash Buffer Concentrate(20X) to 570 mL deionized or distilled water to prepare 1X Wash Buffer.

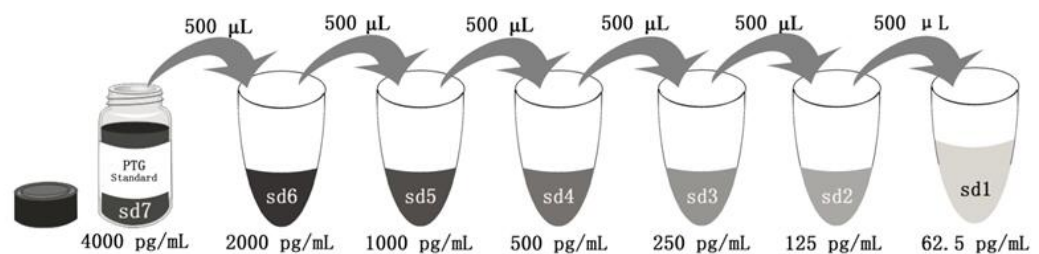
7.2 Detection Antibody, HRP-conjugated (1X): Dilute 100X Detection Antibody, HRP-conjugated 1:100 using Detection Diluent prior to assay. Suggested 1:100 dilution: 10 μ L 100X Detection Antibody, HRP-conjugated + 990 μ L Detection Diluent (Centrifuge the 100 X Detection Antibody solution, HRP-conjugated for a few seconds prior to use)

7.3 Sample Dilution: Different samples should be diluted with corresponding Sample Diluent, samples may require further dilution if the readout values are higher than the highest standard OD reading. Variations in sample collection, processing and storage may affect the results of the measurement.

Recommended Dilution for different sample types: 1:2 is recommended for human serum and plasma; 1:2 or 1:4 is recommended for cell culture supernatant.

7.4 Standard Serial Dilution:

For human serum and plasma samples, add 2 mL Sample Diluent PT 4B1 in protein standard; For cell culture supernatant, add 2 mL Sample Diluent PT 1-ef in protein standard.



| | | | | | | | |
|--|--------------|-------------|-------------|-------------|-------------|-------------|-------------|
| Add # μ L of Standard diluted in the previous step | — | 500 μ L | 500 μ L | 500 μ L | 500 μ L | 500 μ L | 500 μ L |
| # μ L of Sample Diluent PT 4B1 or PT 1-ef | 2000 μ L | 500 μ L | 500 μ L | 500 μ L | 500 μ L | 500 μ L | 500 μ L |
| | "sd7" | "sd6" | "sd5" | "sd4" | "sd3" | "sd2" | "sd1" |

8. Assay Procedure Summary

Bring all reagents to room temperature before use (Detection antibody, HRP-conjugated can be used immediately). 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.

8.1 Take out the required number of microplate strips and return excess strips to the foil pouch containing the drying reagent pack and reseal; store at 4°C immediately. Microplate strips should be used in one week.

8.2 Preset the layout of the microplate, including control group, standard group and sample group, add 100 µL of each standard and sample to the appropriate wells. (Make sure sample addition is uninterrupted and completed within 5 to 10 minutes, it is recommended to assay all standards, controls, and samples in duplicate).

8.3 Seal plate with cover seal, pressing it firmly onto top of microwells. Incubate the plate for 2 hours at 37°C.

8.4 Wash

1) Gently remove the cover seal. Discard the liquid from wells by aspirating or decanting. Remove any residual solution by tapping the plate a few times on fresh paper towels.

2) Wash 4 times with 1X Wash Buffer, using at least 350-400 µL per well. Following the last wash, firmly tap plates on fresh towels 10 times to remove residual Wash Buffer. Avoid getting any towel fibers in the wells or wells drying out completely.

8.5 Add 100 µL of 1X Detection antibody, HRP-conjugated solution (refer to Reagent Preparation 7.2) to each well. Seal plate with cover seal and incubate for 1 hour at 37°C.

8.6 Repeat wash step in 8.4.

8.7 Signal development: Add 100 µL of TMB substrate solution to each well, protected from light. Incubate for 15 to 20 minutes. Substrate Solution should remain colorless until added to the plate.

8.8 Quenching color development: Add 100 µL of Stop Solution to each well in the same order as addition of the TMB substrate. Mix by tapping the side of the plate gently. NB: Avoid skin and eye contact with the Stop solution.

8.9 Read results: Immediately after adding Stop solution read the absorbance on a microplate reader at a wavelength of 450 nm. If possible, perform a double wavelength readout (450 nm and 630 nm).

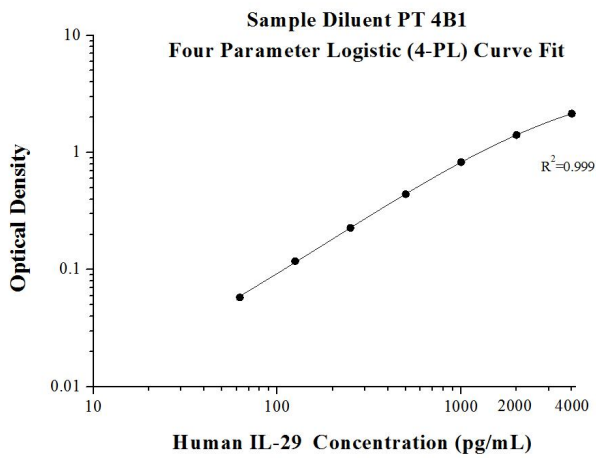
8.10 Data analysis: Calculate the average of the duplicate readings (OD value) for each standard and sample, and subtract the average of the zero standard absorbance. Construct a standard curve by plotting the mean absorbance for each standard on the y-axis against the concentration on the x-axis, use four-parameter logistic curve-fit (4-PL) analysis to do this. If the samples have been diluted, the OD readout from the standard curve must be multiplied by the dilution factor used.

| Step | Reagent | Volume | Incubation | Wash | Notes |
|------|--|--------|------------|-------------|------------------------------|
| 1 | Standard and Samples | 100 µL | 120 min | 4 times | Cover Wells incubate at 37°C |
| 2 | Diluent Detection antibody, HRP-conjugated Solution | 100 µL | 60 min | 4 times | Cover Wells incubate at 37°C |
| 3 | TMB Substrate | 100 µL | 15-20 min | Do not wash | Incubate in the dark at 37°C |
| 4 | Stop Solution | 100 µL | 0 min | Do not wash | - |
| 5 | Read plate at 450 nm and 630 nm immediately after adding Stop solution. DO NOT exceed 5 minutes. | | | | |

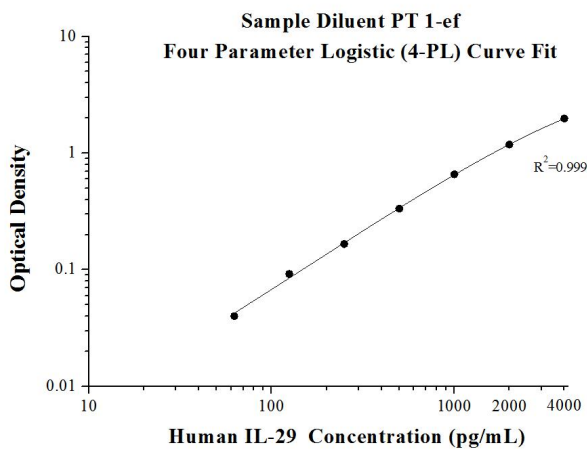
9. Validation Data

9.1 Standard curve

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 | 0.106 0.09 | 0.098 | - |
| 62.5 | 0.159 0.152 | 0.156 | 0.058 |
| 125 | 0.216 0.216 | 0.216 | 0.118 |
| 250 | 0.337 0.312 | 0.325 | 0.227 |
| 500 | 0.554 0.523 | 0.539 | 0.441 |
| 1000 | 0.941 0.911 | 0.926 | 0.828 |
| 2000 | 1.527 1.493 | 1.51 | 1.412 |
| 4000 | 2.278 2.213 | 2.246 | 2.148 |



| (pg/mL) | O.D | Average | Corrected |
|---------|----------------|---------|-----------|
| 0 | 0.085 0.097 | 0.091 | - |
| 62.5 | 0.128 0.134 | 0.131 | 0.04 |
| 125 | 0.178 0.187 | 0.183 | 0.092 |
| 250 | 0.239 0.274 | 0.257 | 0.166 |
| 500 | 0.417 0.433 | 0.425 | 0.334 |
| 1000 | 0.745 0.752 | 0.749 | 0.658 |
| 2000 | 1.249 1.302 | 1.276 | 1.185 |
| 4000 | 2.045 2.092 | 2.069 | 1.978 |

9.2 Precision

Intra-assay Precision (Precision within an assay) Three samples of known concentration were tested 20 times on one plate to assess intra-assay precision.

Inter-assay Precision (Precision between assays) Three samples of known concentration were tested in 24 separate assays to assess inter-assay precision.

| Intra-assay Precision | | | | | Inter-assay Precision | | | | |
|-----------------------|----|--------------|--------|-----|-----------------------|----|--------------|--------|-----|
| Sample | n | Mean (pg/mL) | SD | CV% | Sample | n | Mean (pg/mL) | SD | CV% |
| 1 | 20 | 229.0 | 9.73 | 4.2 | 1 | 24 | 212.1 | 8.67 | 4.1 |
| 2 | 20 | 878.2 | 24.62 | 2.8 | 2 | 24 | 869.2 | 25.66 | 3.0 |
| 3 | 20 | 3674.4 | 139.35 | 3.8 | 3 | 24 | 3569.6 | 145.19 | 4.1 |

9.3 Recovery

The recovery of human IL-29 spiked to three different levels throughout the range of the assay in various matrices was evaluated.

| Sample Type | | Average% of Expected | Range (%) |
|--------------------------|-----|----------------------|-----------|
| Human plasma | 1:2 | 91 | 80-105 |
| | 1:4 | 93 | 72-108 |
| Cell culture supernatant | 1:2 | 104 | 98-109 |
| | 1:4 | 101 | 90-121 |

9.4 Sample values

Human serum -Sixteen human serum samples from healthy volunteers were evaluated for human IL-29 in this assay. No medical histories were available for the donors used in this study.

| Sample Type | Mean of Detectable (pg/mL) | %Detectable | Range (pg/mL) |
|--------------------|----------------------------|-------------|---------------|
| Human serum (n=16) | 185.8 | 62.5 | 27-744 |

Cell culture supernatant

HUVEC cells were stimulated by adding human IFN-alpha at 100 ng/ml for 20 hours, followed by the addition of 30 µg/mL Poly(I:C) for 3 hours. Aliquots of the cell culture supernatant was removed and assayed for levels of human IL-29.

| Condition | (pg/mL) |
|--------------|---------|
| Unstimulated | 46.8 |
| Stimulated | 378.1 |

9.5 Sensitivity

The minimum detectable dose of human IL-29 is 3.3 pg/mL. This was determined by adding two standard deviations to the concentration corresponding to the mean O.D. of 20 zero standard replicates.

9.6 Linearity

To assess the linearity of the assay, three samples were spiked with high concentrations of human IL-29 in various matrices and diluted with the appropriate **Sample Diluent** to produce samples with values within the dynamic range of the assay.

| | | Human plasma (Sample Diluent PT 4B1) | Cell culture supernatant (Sample Diluent PT 1-ef) |
|------|----------------------|--------------------------------------|---|
| 1:2 | Average% of Expected | 104 | 78 |
| | Range (%) | 84-124 | 71-86 |
| 1:4 | Average% of Expected | 100 | 82 |
| | Range (%) | 76-124 | 74-90 |
| 1:8 | Average% of Expected | 88 | 83 |
| | Range (%) | 73-103 | 76-90 |
| 1:16 | Average% of Expected | 77 | 98 |
| | Range (%) | 69-86 | 80-115 |

10. References

1. Yoshitaka Hosokawa. et al. (2017) *Immunol Invest.* 46(6):615-624.
2. Katrin Witte et al. (2010) *Cytokine Growth Factor Rev.* 21(4):237-51.
3. Noah E Kelm. et al. (2016) *Crit Rev Oncol Hematol.* 106:91-8.
4. Jia-Min Wang. et al. (2019) *J Cell Mol Med.* 23(12):7926-7932.