ClinMax™ Human Perforin ELISA Kit

Catalog Number：CEA-B034

Assay Tests: 96 tests

For Research Use Only. Not For Use in Diagnostic or Therapeutic Procedures

IMPORTANT: Please carefully read this user guide before performing your experiment.

Product information

This kit is specifically designed for the accurate quantitation of Human Perforin from cell culture supernates, serum and plasma.

The principle of this assay employs a quantitative sandwich enzyme immunoassay approach. Initially, a microplate is coated with a capture antibody. Then, samples and biotinylated capture antibody are added to the wells. After the removal of any unbound materials through washing, streptavidin-HRP (SA-HRP) conjugate is added to the wells. Streptavidin has a very high affinity for biotin, so it binds to the biotinylated capture antibody that is already bound to the target antigen. After washing, a substrate specific to HRP is added to the wells. HRP catalyzes a reaction that converts the substrate into a detectable signal, often a color change or luminescence, depending on the substrate used. This enzymatic reaction amplifies the signal, allowing for higher sensitivity in detecting the target analyte. The intensity of the signal is measured using a spectrophotometer.

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NOTE:

1. This kit is for research use only and is not for use in diagnostic or therapeutic applications.
2. Please do not use the kit after the expiration date indicated on the kit label.
3. Do not mix or substitute reagents with those from other lots or sources.

Manufactured and distributed by
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Contents

The kit contains sufficient reagents for 96 wells.

Catalog	Contents	Amount
CEA034-C01	Pre-coated Anti-Perforin Antibody Microplate	1 plate
CEA034-C02	Human Perforin Standard	500 µL×2
CEA034-C03	Biotin-Anti-Perforin Antibody Con. Solution	150 µL
CEA034-C04	Biotin-Antibody Dilution Buffer	15 mL
CEA034-C05	Streptavidin-HRP Con. Solution	1.5 mL
CEA034-C06	Streptavidin-HRP Dilution Buffer	15 mL
CEA034-C07	20× Washing Buffer	50 mL
CEA034-C08	Sample Dilution Buffer	15 mL×2
CEA034-C09	Substrate Solution	12 mL
CEA034-C10	Stop Solution	6 mL

Storage

Keep the unopened kit stored at 2-8 °C. Avoid using the kit beyond its expiration date.
For opened kit and reconstituted reagents, with the exception of the content listed in following table, others can be stored for up to 30 days at 2-8 °C.

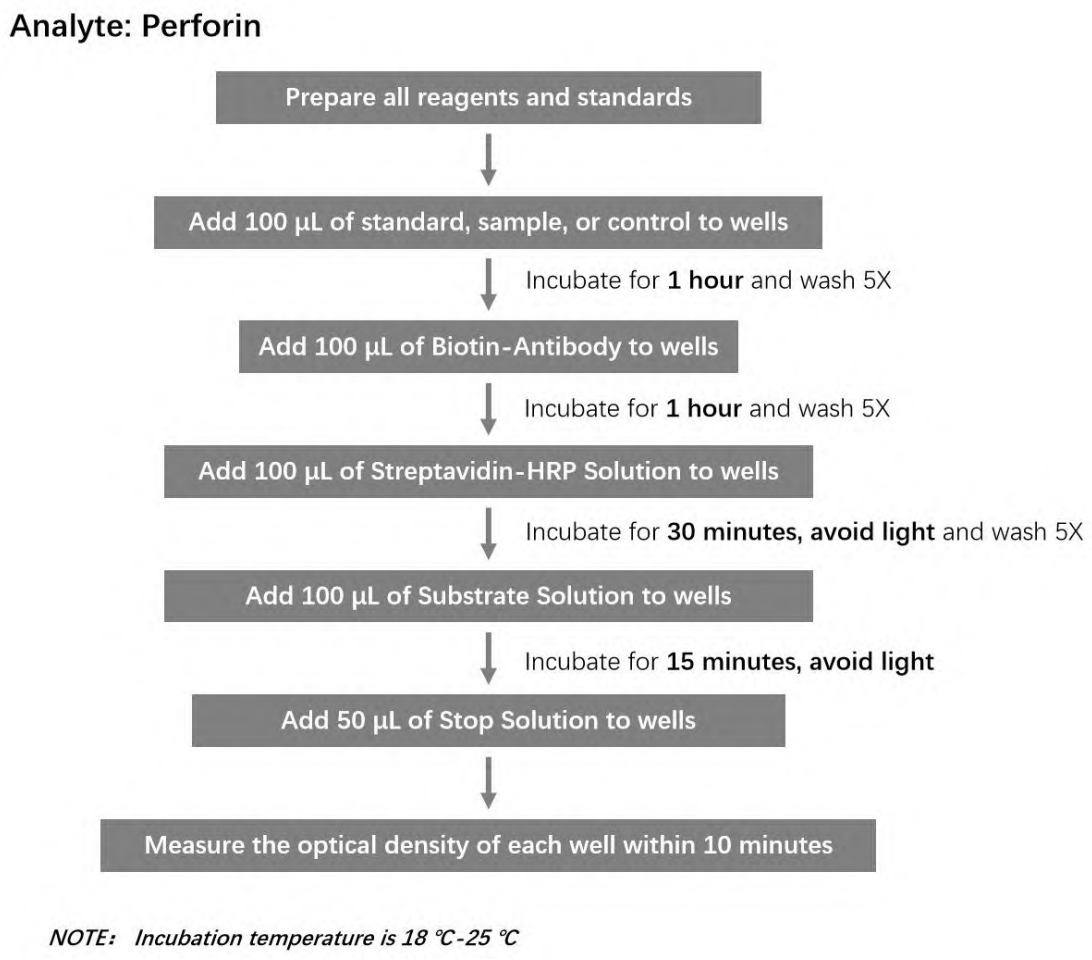
Contents	Storage conditions
Pre-coated Anti-Perforin Antibody Microplate	Return unused wells to the foil pouch, reseal along entire edge of zip-seal. May be stored for up to 1 month at 2-8°C.

NOTE: Streptavidin-HRP Con. Solution and Substrate Solution should avoid light.

Required materials not supplied.

Instrument	Microplate reader capable of measuring absorbance at 450 nm
Reagents	Deionized or distilled water
Consumables	50 mL and 500 mL graduated cylinders
	Pipettes and pipette tips
	Tubes to prepare standard dilutions.

Workflow



NOTE: Incubation temperature is 18 ℃-25 ℃

Prepare the working buffers and standard dilutions.

IMPORTANT: Bring all reagents to room temperature before use. If crystals have formed in buffer solution, place the buffer solution in an 37°C incubator until the crystals have completely dissolved and bring the solution back to room temperature before use.

Prepare the working buffers.

1. 1×Washing Buffer: Dilute 50 mL 20×Washing Buffer with deionized or distilled water to 1000 mL.
2. Biotin-Anti-Perforin Antibody Solution: Add 120 µL of Biotin-Anti-Perforin Antibody Con. Solution to 12 mL Biotin-Antibody Dilution Buffer, thoroughly mix. The solution was freshly prepared just before use.
3. Perforin Streptavidin-HRP Solution: Add 1.2 mL of Perforin Streptavidin-HRP Con. Solution to 12 mL of Streptavidin-HRP Dilution Buffer, thoroughly mix. The solution was freshly prepared just before use.

Prepare the standard serial dilutions.

1. Label 6 tubes, one for each standard point: Std.-1, Std.-2, Std.-3, Std.-4, Std.-5, Std.-6, .
2. Add 500 µL of the liquid from Human Perforin Standard（CEA034-B02）to tube Std.-1, (Std.-1 =2000 pg/mL).
3. Prepare serial dilutions for the standard curve as follows: Add 250 µL of Sample Dilution Buffer to each tube (Std.-2, Std.-3, Std.-4, Std.-5, Std.-6).
4. Transfer 250 µL of liquid from Std.-1 to the tube Std.-2, and thoroughly mix (Std.-2 = 1000 pg/mL).
5. Continue to transfer 250 µL of liquid from previous dilution tube to the next dilution tube until add liquid to tube Std.-6.
6. Sample Dilution Buffer serves as zero standard (blank).

PROCEDURE OF ASSAY

1. Add 100 µL of Perforin Standard, sample, or control to wells, Seal the plate with microplate sealing film. Incubate at room temperature (18-25 °C) for **1 hour.**
2. Aspirate each well and add 300 µL of 1×Washing Buffer to each well, gently tap the plate for **1 minute**. Remove any remaining Washing Buffer by aspirating or decanting. Invert the plate and blot it against clean paper towels. Repeat the wash process four times for a total of five washes.
3. Add 100 µL Biotin-Anti-Perforin Antibody Solution to each well, Seal the plate with microplate sealing film. Incubate at room temperature (18-25 °C) for **1 hour.**
4. Repeat step 2.
5. Add 100 µL of Perforin Streptavidin-HRP Solution to each well. Seal the plate with microplate sealing film. Incubate at room temperature (18-25 °C) for **30 minutes, avoid light.**
6. Repeat step 2.
7. Add 100 µL of Substrate Solution to each well. Seal the plate with microplate sealing film and incubate at room temperature (18-25 °C) for **15 minutes, avoid light**.
8. Add 50 µL of Stop Solution to each well. Tap the plate gently to ensure thorough mixing. ***Note:*** *the color in the wells should change from blue to yellow.*
9. Read the absorbance at 450nm and 630nm using Microplate reader within 10minutes. **Note:** To reduce the background noise, subtract the readings at 630nm from the readings at 450nm.

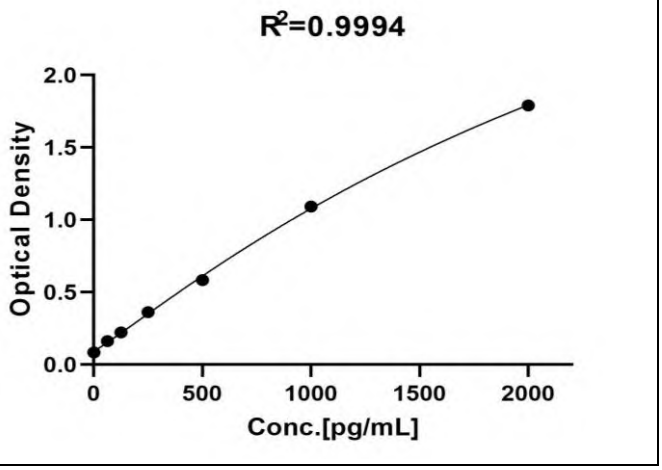
CALCULATION OF RESULTS

1. Compute the average of the duplicated readings for every standard, control, and sample. Then, subtract the average optical density (O.D.) of the zero standard(blank).
2. Establish a standard curve by processing the data using computer software capable of executing a four-parameter logistic (4-PL) curve fitting.
3. Normal range of Standard curve: $R^2 \geqslant 0.9900$.
4. If the OD value of the sample to be tested is higher than the highest standard, the sample shall be diluted with dilution buffer and assay repeated.

Typical data

Note: *For each experiment, a standard curve needs to be set for each microplate, and the specific OD value may vary depending on different laboratories, testers, or equipment. The following example data is for reference only. The sample concentration was calculated based on the results of the standard curve.*

Perforin Standard (pg/mL)	$OD_{450nm-630nm}$
2000	1.791
1000	1.093
500	0.585
250	0.363
125	0.223
62.5	0.162
Blank	0.084



PERFORMANCE CHARACTERISTICS

1. Sensitivity

The minimum detectable concentration (MDC) of Perforin is typically less than 60 pg/mL. The MDC was determined by adding two standard deviations to the mean optical density value of twenty zero standard replicates and calculating the corresponding concentration.

2. Intra-Assay Precision

Ten replicates of each of 3 samples containing different Perforin concentrations were tested in one assay. Acceptable criteria: CV < 10%.

Sample Concentration (pg/mL)	Mean (pg/mL)	SD	Numbers	CV
62.5	57.19	3.16	8	5.52%
500	497.51	27.92	8	5.61%
1000	993.59	31.35	8	3.16%

3. Inter-Assay Precision

Five samples containing different concentrations of Perforin were tested in independent assays. Acceptable criteria: CV<15%.

Sample Concentration (pg/mL)	Mean (pg/mL)	SD	Numbers	CV
62.5	57.56	3.33	9	5.78%
500	486.52	38.53	9	7.92%
1000	972.06	32.46	9	3.34%

4. Specificity

No cross-reactivity was observed when this kit was used to analyze the following recombinant cytokines at up to 1 μg/mL.

Human	IL-1β, IL-2, IL-4, IL-5, IL-6, IL-7, IL-8, IL-12 p70, IL-10、IL-10、MCP-1、GM-CSF、TNF-α、IFN-γ

TROUBLESHOOTING GUIDE

Problem	Cause	Solution
Poor standard curve	* Inaccurate pipetting	* Check pipettes
Large CV	* Inaccurate pipetting * Air bubbles in wells	* Check pipettes * Remove bubbles in wells
High background	* Plate is insufficiently washed * Contaminated wash buffer	* Review the manual for proper wash. * Make fresh wash buffer
Very low readings across the plate	* Incorrect wavelengths * Insufficient development time	* Check filters/reader * Increase development time
Samples are reading too high, but standard curve looks fine	* Samples contain cytokine levels above assay range	* Dilute samples and run again
Drift	* Interrupted assay set-up * Reagents not at room temperature	* Assay set-up should be continuous - have all standards and samples prepared appropriately before commencement of the assay * Ensure that all reagents are at room temperature before pipetting into the wells unless otherwise instructed in the antibody inserts