

Human Complement C5 ELISA Instructions

CAT. No: AEHY0007

CONTENT

No.	Content	Cat. No	Volume
1	CP (Coated Plate)	ER0001 CP	96 wells
2	S (Standard)	ER0001S,S1~ S7,S0	9 vials
3	DA (Detect Antibody)	ER0001 DA	6 ml/bottle
4	SH(Streptavidin-HRP)	ESH01	12 ml/bottle
5	AB (Assay Buffer 1×)	EAB01	12 ml/bottle
6	TS (TMB Substrate)	ETS01	12 ml/bottle
7	SS (Stop Solution)	ESS01	12 ml/bottle
8	WB (Wash Buffer 10×)	EWB01	50 ml/bottle
9	SF (Sealer Film)	ESF01	6 pieces

NOTE: After the kit is opened, the stabilization period of each content is 30 days, so please use the kit within 30 days after opening.

SAMPLE DILUTION

Samples such as serum, plasma require at least a 100-fold dilution into Sample Diluent. A suggested 100-fold dilution is 5 μ l of sample + 495 μ l of Sample Diluent.

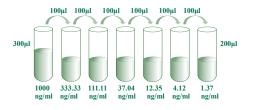
REAGENT PREPARATION

Washing Buffer (1 \times) Preparation: Pour entire contents (50 ml) of the Washing Buffer Concentrate (10 \times) into a clean 500 ml graduated cylinder. Bring to final volume of 500 ml with glass-distilled or deionized water. Transfer to a clean wash bottle and store at 2 to 25 $^{\circ}$ C.

1×DA Preparation: Mix well prior to making dilutions.

Make a 1:100 dilution of the concentrated Detect Antibody solution with **DD** (Detect Antibody Diluent) in a clean plastic tube as needed according to the Standards and Samples.

Standard Curve Preparation: S1 to S7 and S0 is ready to use for serum and plasma.



ASSAY PROCEDURE

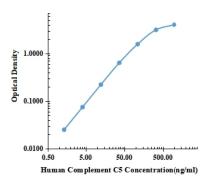
Bring all reagents and samples to room temperature before use.

- Prepare all reagents and working standards as directed in the previous sections.
- Remove excess CP (Coated Plate) strips from the plate frame, return them to the foil pouch and reseal.
- 3. Add 50 µl of AB (Assay Buffer) to each well.
- Add 50 μl or 10 μl of Standard or sample per well. Ensure reagent addition is uninterrupted and completed within 15 minutes.
- 5. Add 50 μl of DA-H (Detect Antibody) to each well.
- 6. Cover with an SF (Sealer Film). Incubate at room temperature (18 to 25°C) for 1 hour on a micro-plate shaker set at 500 rpm.
- 7. Aspirate each well and wash, repeating the process four times. Wash by filling each well with WB (Washing Buffer 300 µl). Complete removal of liquid at each step is essential to good performance. After the last wash, remove any remaining WB (Washing Buffer) by aspirating or decanting. Invert the plate and blot it against clean paper towels.
- 8. Add 100 μl of TS (TMB Substrate) to each well. Incubate for 5-30 minutes at room temperature.
- 9. Add 100 μl of SS (Stop Solution) to each well.
- Determine the optical density within 30 minutes, using microplate reader set to 450 nm corrected with 570 nm or 630 nm.



TYPICAL DATA

Human Complement C5 Typical Standard



ng/ml	O.D.		Average	Corrected
0.00	0.0061	0.0057	0.0059	
1.37	0.0318	0.0300	0.0309	0.0250
4.12	0.0813	0.0794	0.0804	0.0745
12.35	0.2275	0.2267	0.2271	0.2212
37.04	0.6468	0.6343	0.6406	0.6347
111.11	1.5860	1.5560	1.5710	1.5651
333.33	3.2300	3.0340	3.1320	3.1261
1000.00	4.1306	3.9187	4.0247	4.0188

SENSITIVITY

The minimum detectable dose (MDD) of human Complement C5 is typically less than 0.01 ng/ml (50 μ l of sample volume) or 0.18 ng/ml (10 μ l of sample volume).

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

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)

	Intra-assay Precision		Inter-assay Precision		recision	
Sample Number	S1	S2	S3	S1	S2	S3
	22	22	22	6	6	6
Average (ng/ml)	17.8	92.6	287.7	16.6	85.8	261.2
Standard deviation	0.7	4.5	15.2	0.7	4.8	16.9
Coefficient of variation (%)	3.7	4.9	5.3	4.5	5.6	6.5

RECOVERY

The spike recovery was evaluated by spiking 3 levels of human Complement C5 into health human serum sample. The un-spiked serum was used as blank in this experiment.

The recovery ranged from 91% to 118% with an overall mean recovery of 101%.

LINEARITY

To assess the linearity of the assay, five samples were spiked with high concentration of Complement C5 in human serum and diluted with Sample Diluent to produce samples with values within the dynamic range of the assay.

The linearity ranged from 94% to 114% with an overall mean recovery of 103%.

SAMPLE VALUES

Serum/Plasma - Thirty samples from apparently healthy volunteers were evaluated for the presence of human Complement C5 in this assay. No medical histories were available for the donors.

Sample Matrix	Sample Evaluated	Range (µg/ml)	Detectable %	Mean of Detectable (μg/ml)
Serum	30	1.79-58.93	100	25.41

n.d. = non-detectable. Samples measured below the sensitivity are considered to be non-detectable.

DISCLAIMER AND VERSION

For research use only. Not for use in clinical diagnostic procedures.

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