Mouse IgE Immunoassay

Catalog Number: EA800160

For the quantitative determination of mouse IgE concentrations in cell culture supernates, serum, and plasma.

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

MANUFACTURED AND DISTRIBUTED BY:

OriGene Technologies,Inc.

9620 Medical Center Drive Suite 200
Rockville, MD 20850,USA

Tel: 1-301-340-3188 Fax: 301-340-9254 Mail: techsupport@origene.com Web: www.origene.com

ORIGENE

TABLE OF CONTENTS

SECTION PAGE

BACKGROUND..1

PRINCIPLE OF THE ASSAY...1

TECHNICAL HINTS AND LIMITATIONS...2

PRECAUTIONS...2

KIT COMPONENTS& STORAGE CONDITIONS..3

OTHER SUPPLIES REQUIRED BUT NOT SUPPLIED...4

SPECIMEN COLLECTION & STORAGE...4

REAGENTS PREPARATION..4

ASSAY PROCEDURE 6

CALCULATION OF RESULTS...6

PERFORMANCE CHARACTERISTICS..8

REFERENCES..10

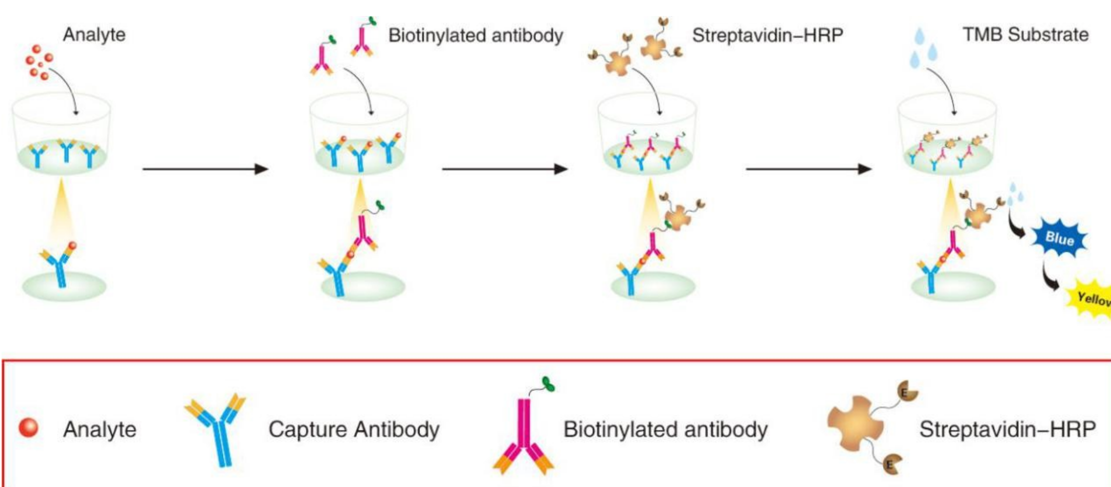
BACKGROUND

Immunoglobulin E (IgE) is a kind of antibody (or immunoglobulin (Ig) "isotype") that has only been found in mammals. IgE's main function is immunity to parasites such as helminths like Schistosoma mansoni, Trichinella spiralis, and Fasciola hepatica. IgE is utilized during immune defense against certain protozoan parasites such as Plasmodium falciparum. IgE also has an essential role in type I hypersensitivity, which manifests various allergic diseases, such as allergic asthma, most types of sinusitis, allergic rhinitis, food allergies, and specific types of chronic urticaria and atopic dermatitis. IgE also plays a pivotal role in responses to allergens, such as: anaphylactic drugs, bee stings, and antigen preparations used in desensitization immunotherapy.

PRINCIPLE OF THE ASSAY

This assay employs the quantitative sandwich enzyme immunoassay technique. A monoclonal antibody specific for IgE has been pre-coated onto a microplate. Standards and samples are pipetted into the wells and any IgE present is captured by the coated antibody after incubation. Following extensive washing, a biotin-conjugate antibody specific for IgE is added to detect the captured IgE protein in sample. For signal development, horseradish peroxidase (HRP)-conjugated Streptavidin is added, followed by Tetramethyl-benzidine (TMB) reagent. Following a wash to remove any unbound combination, and enzyme conjugate is added to the wells. Solution containing sulfuric acid is used to stop color development and the color intensity which is proportional to the quantity of bound protein is measurable at 450nm.

Schematic diagram：



TECHNICAL HINTS AND LIMITATIONS

1. This ELISA should not be used beyond the expiration data on the kit label.
2. To avoid cross-contamination, use a fresh reagent reservoir and pipette tips for each step.
3. To ensure accurate results, some details, such as technique, plasticware and water sources should be emphasized.
4. A thorough and consistent wash technique is essential for proper assay performance.
5. A standard curve should be generated for each set of samples assayed.
6. It is recommended that all standards and samples be assayed in duplicate.
7. Avoid microbial contamination of reagents and buffers. Buffers containing protein should be made under aseptic conditions and be prepared fresh daily.
8. In order to ensure the accuracy of the results, the standard curve should be made every time.

PRECAUTIONS

The Stop Solution suggested for use with this kit is an acid solution. Wear protective gloves, clothing, eye, and face protection. Wash hands thoroughly after handling.

KIT COMPONENTS& STORAGE CONDITIONS

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

PART	SIZE	STORAGE OF OPENED/ RECONSTITUTED MATERIAL
Microwell Plate - antibody coated 96-well Microplate (8 wells ×12 strips)	1 plate	Return unused wells to the foil pouch containing the desiccant pack. Reseal along entire edge of the zip-seal. May be stored for up to 1 month at 2 – 8℃**
Standard-lyophilized,40ng/ml upon reconstitution	2 vials	Aliquot and Store at -20°C** for six months
Concentrated Biotin-Conjugated antibody(100X) - 120 ul/vial	1 vial	Store at 2-8°C **for six months
Concentrated Streptavidin-HRP solution(100X) - 120 ul/vial	1 vial	Store at 2-8°C** for six months
Standard /Sample Diluent - 16 ml/vial	1 bottle	Store at 2-8°C** for six months
Biotin-Conjugate antibody Diluent - 16 ml/vial	1 bottle	Store at 2-8°C** for six months
Streptavidin-HRP Diluent - 16 ml/vial	1 bottle	Store at 2-8°C** for six months
Wash Buffer Concentrate (20x) - 30 ml/vial	1 bottle	Store at 2-8°C** for six months
Substrate Solution - 12 ml/vial	1 bottle	Store at 2-8°C** for six months
Stop Solution - 12 ml/vial	1 bottle	Store at 2-8°C** for six months
Plate Cover Seals	4 pieces	

OTHER SUPPLIES REQUIRED BUT NOT SUPPLIED

1. Microplate reader capable of measuring absorbance at 450 nm.
2. Pipettes and pipette tips.

3. Deionized or distilled water.
4. Squirt bottle, manifold dispenser, or automated microplate washer.
5. 500 mL graduated cylinder.

SPECIMEN COLLECTION & STORAGE

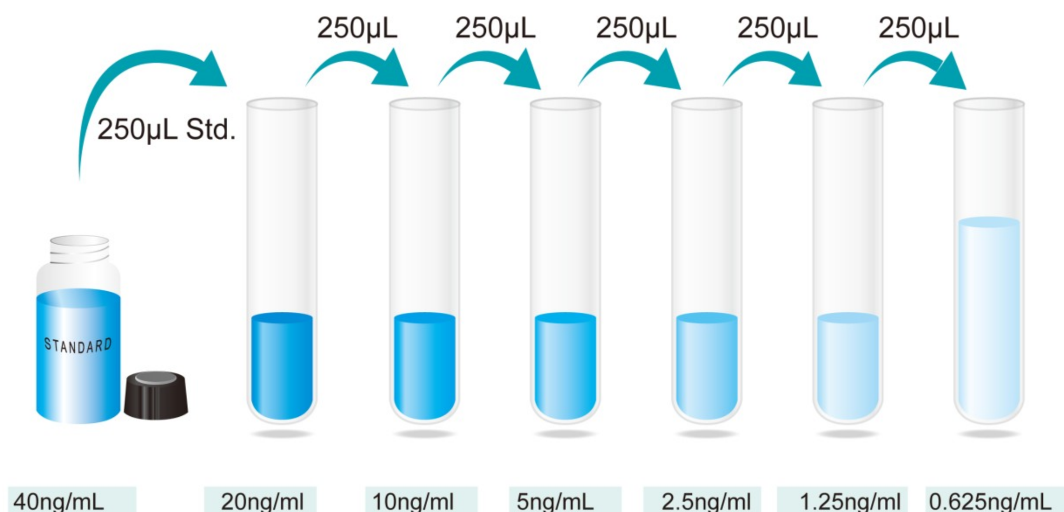
Cell Culture Supernates - Centrifuge cell culture media at 1000×g to remove debris. Assay immediately or aliquot and store samples at ≤ -20 °C. Avoid repeated freeze-thaw cycles.
Serum - Use a serum separator tube (SST) and allow samples to clot for 2 hours at room temperature or overnight at 2-8℃. Centrifuge approximately for 15 minutes at 1000×g. Assay immediately or aliquot and store samples at ≤ -20 °C. Avoid repeated freeze-thaw cycles.
Plasma - Collect plasma using EDTA, heparin, or citrate as an anticoagulant. Centrifuge for 15 minutes at 1000×g within 30 minutes of collection. Assay immediately or aliquot and store samples at ≤ -20 °C. Avoid repeated freeze-thaw cycles.
Note: It is recommended to conduct a pre-test before the formal experiment to determine the dilution ratio.

REAGENTS PREPARATION

1. **Temperature returning** - Bring all kit components and specimen to room temperature(20-25℃) before use.

2. **Wash Buffer** - Dilute 30mL of Wash Buffer Concentrate with 570mL of deionized or distilled water to prepare 600mL of Wash Buffer. If crystals have formed in the concentrate Wash Buffer, warm to room temperature and mix gently until the crystals have completely dissolved.

3. **Standard/Specimen** - Reconstitute the Standard with 0.5mL of Standard/Sample Diluent. This reconstitution produces a stock solution of 40ng/mL. Allow the standard to sit for a minimum of 15 minutes with gentle agitation prior to making dilutions. Pipette 250μL of Standard/ Sample Diluent into 20ng/ml tube and the remaining tubes. Use the stock solution of 40ng/mL to produce a 2-fold dilution series (below). Mix each tube thoroughly and change pipette tips between each transfer. The 40ng/mL standard serves as the high standard. The Standard/ Sample Diluent serves as the zero standard (0 pg/mL).



***If you do not run out of re-melting standard, store it at -20°C. Diluted standard shall not be reused.**

4. **Working solution of Biotin-Conjugate anti-mouse IgE antibody:** Make a 1:100 dilution of the concentrated Biotin-Conjugate solution with the Biotin-Conjugate antibody Diluent in a clean plastic tube.

   ***The working solution should be used within one day after dilution.**

5. **Working solution of Streptavidin-HRP**: Make a 1:100 dilution of the concentrated Streptavidin-HRP solution with the Streptavidin-HRP Diluent in a clean plastic tube.

   ***The working solution should be used within one day after dilution.**

ASSAY PROCEDURE

Prepare all reagents and standards as directed.

⇩

Add 100l standard or samples to each well, incubate 60 minutes,room temperature(25±2℃).

Add 100l working solution of Biotin-Conjugate anti-mouse IgE antibody to each well, incubate 60 minutes,room temperature(25±2℃).

⇩ Aspirate and wash 4 times

Add 100l working solution of Streptavidin-HRP to each well, incubate 30 minutes, room temperature(25±2℃).Avoid placing the plate in direct light.

⇩ Aspirate and wash 5 times

Add 100l Substrate solution to each well, incubate 5-20 (depending on signal) minutes,room temperature(25±2°C).Protect from light.

⇩

Add 50l Stop solution to each well. Read at 450nm within 30 minutes.

CALCULATION OF RESULTS

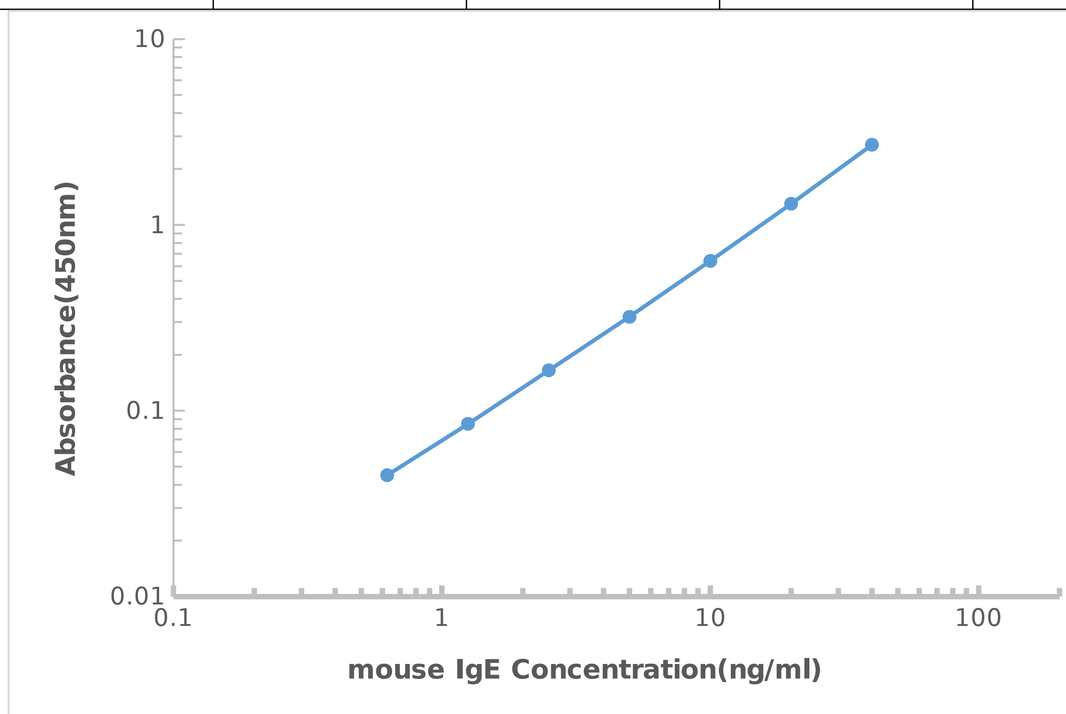
1. The standard curve is used to determine the amount of specimens.
2. First, average the duplicate readings for each standard, control, and sample. All O.D. values are subtracted by the mean value of blank control before result interpretation.
3. Construct a standard curve by reducing the data using 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.
4. The data may be linearized by plotting the log of the IgE 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.

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

Typical data using the IgE ELISA

Standard(ng/ml)	OD.	OD.	Average	Corrected
0	0.056	0.052	0.054	-------
0.625	0.186	0.185	0.185	0.131
1.25	0.214	0.213	0.214	0.160
2.5	0.319	0.317	0.318	0.264
5.0	0.506	0.504	0.505	0.451
10.0	0.822	0.819	0.821	0.766
20.0	1.339	1.333	1.336	1.282
40.0	2.181	2.172	2.177	2.122



Representative standard curve for IgE ELISA.

Performance Characteristics

SENSITIVITY: The minimum detectable dose was0.12ng/mL.

SPECIFICITY: This assay recognizes both natural and recombinant mouse IgE. The factors listed below were prepared at 100ng/ml in Standard /sample Diluent and assayed for cross-reactivity and no significant cross-reactivity or interference was observed.

Factors assayed for cross-reactivity

Recombinant mouse	Natural proteins	Recombinant human
BMP1	IL-2	
BMP2	IL-4	
BMP4	IL-6	
IL-1β	IL-8	
IL-1ra		

REPEATABILITY: The coefficient of variation of both intra-assay and inter-assay were less than 10%.

RECOVERY： The recovery of IgE spiked to three different levels in four samples throughout the range of the assay in various matrices was evaluated.

Recovery of IgE in two matrices

Sample Type	Average % of Expected Range (%)	Range (%)

Citrate plasma	92	84-101
Cell culture supernatants	95	97-103

LINEARITY: To assess the linearity of the assay, three samples were spiked with high concentrations of IgE in various matrices and diluted with the appropriate Sample Diluent to produce samples with values within the dynamic range of the assay. (The plasma samples were initially diluted 1:1)

Dilution ratio	Recovery (%)	Citrate plasma	Cell culture supernatants
1:2	Average% of Expected	**102**	**101**
	Range (%)	**92－111**	**98－104**
1:4	Average% of Expected	**104**	**104**
	Range (%)	**94－113**	**103－105**
1:8	Average% of Expected	**97**	**105**
	Range (%)	**94－100**	**94－115**
1:16	Average% of Expected	**100**	**108**
	Range (%)	**96－103**	**99－116**

REFERENCE

1. Erb KJ (2007). Eur. J. Immunol. 37 (5): 1170–3.
2. Pfister K, et al. (1983). "IgE production in rat fascioliasis".
   Parasite Immunol. 5 (6): 587–93.
3. Gould HJ, et al. (2003). Annu. Rev. Immunol. 21: 579–628.
4. Winter WE, et al. (2000). Arch. Pathol. Lab. Med. 124 (9): 1382–5.
5. Karagiannis SN, et al. (2003). Eur. J. Immunol. 33 (4): 1030–40.
6. Elkayam O, et al. (1995). Allergy 50 (1): 94–6.