



User's Manual

# Mouse B. pertussis (FHA) IgG ELISA kit

**REF** DEIASL129

**Σ** 96T

**RUO**

This product is for research use only and is not intended for diagnostic use.

For illustrative purposes only. To perform the assay the instructions for use provided with the kit have to be used.

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## PRODUCT INFORMATION

### Intended Use

The Mouse Anti-B. Pertussis FHA IgG ELISA Kit detects and quantifies B. pertussis FHA-specific IgG in mouse serum or plasma of vaccinated or immunized animals. This immunoassay is suitable for:

- o Determining immune status relative to non-immune controls;
- o Assessing efficacy of vaccines, including dosage, adjuvantcy, route of immunization and timing;
- o Qualifying and/or standardizing vaccine batches and protocols.

The kit contains no active toxin or inactivated bacteria. The antigen used is highly purified pertussis toxoid (non-toxic). For in vitro research use only (RUO), not for therapeutic or diagnostic use.

### General Description

Pertussis, also known as Whooping Cough, is a highly contagious disease caused by *Bordetella pertussis* bacteria. Vaccines for pertussis, available in combination with vaccines for tetanus, diphtheria, H. influenza b, hepatitis & polio, use acellular components, primarily the inactivated pertussis toxin. The toxin, a protein exotoxin, produced only by B. pertussis, is central to pertussis pathogenesis; vaccination with the toxoid elicits high levels of protection from the disease. Also included are two other highly immunogenic pertussis proteins: pertactin, an outer membrane protein that promotes adhesion to host cells, and filamentous hemagglutinin (FHA).

### Principles of Testing

The Mouse Anti-FHA IgG ELISA kit is based on the binding of mouse anti-FHA IgG in samples to FHA immobilized on the microwells, and anti-FHA IgG antibody is detected by antimouse IgG specific antibody conjugated to HRP (horseradish peroxidase) enzyme. After a washing step, chromogenic substrate (TMB) is added and color is developed by the enzymatic reaction of HRP on the substrate, which is directly proportional to the amount of anti-FHA IgG present in the sample. Stopping Solution is added to terminate the reaction, and absorbance at 450nm is then measured using an ELISA microwell reader. The activity of mouse IgG antibody in samples is calculated relative to anti-FHA calibrators.

### Reagents And Materials Provided

1. FHA Antigen Microwell Strip Plate: 8-well strips (12)
2. Wash Solution Concentrate (100x): 10ml
3. Sample Diluent Concentrate (20x): 10ml
4. Anti-Mouse IgG - HRP Conjugate Concentrate (100x): 0.15ml
5. Anti-FHA IgG Calibrators: 1 U/ml, 2.5 U/ml, 5 U/ml, 10 U/ml, 0.65 ml for each.
6. Anti-FHA IgG Positive Control: 0.65 ml
7. Low NSB Sample Diluent: 30 ml, use as is for sample dilution.
8. TMB Substrate: 12 ml

9. Stop Solution: 12 ml

## Materials Required But Not Supplied

- Pipettors and pipettes that deliver 100ul and 1-10ml. A multichannel pipettor is recommended.
- Disposable glass or plastic 5-15ml tubes for diluting samples and Anti-Mouse IgG HRP Concentrate.
- Graduated cylinder to dilute Wash Concentrate; 0.2-1L.
- Stock bottle to store diluted Wash Solution; 200ml to 1L.
- Distilled or deionized water to dilute reagent concentrates.
- Microwell plate reader at 450 nm wavelength.

## Storage

The microtiter well plate and all other reagents, if unopened, are stable at 2-8°C until the expiration date printed on the box label.

## Specimen Collection And Preparation

Culture medium, serum and other biological fluids may be used as samples with proper dilution to avoid solution matrix interference. For serum, collect blood by venipuncture, allow clotting, and separate the serum by centrifugation at room temperature. For other samples, including tissue culture media, clarify the sample by centrifugation and/or filtration prior to dilution in Sample Diluent. If samples will not be assayed immediately, store refrigerated for up to a few weeks, or frozen for long-term storage.

## Plate Preparation

Bring all reagents to room temperature (18-30o C) equilibration (at least 30 minutes).

- Determine the number of wells for the assay run. Duplicates are recommended, including 8 Calibrator wells and 2 wells for each sample and control to be assayed.
- Remove the appropriate number of microwell strips from the pouch and return unused strips to the pouch. Reseal the pouch and store refrigerated.
- Add 200-300ul Working Wash Solution to each well and let stand for about 5 minutes. Aspirate or dump the liquid and pat dry on a paper towel before sample addition.

### Assay Design

Review Calculation of Results and Limits of the Assay before proceeding:

- Select the proper sample dilutions accounting for expected potency of positives and minimizing non-specific binding (NSB) and other matrix effects; for example, net signal for nonimmune samples should be <0.5 OD. This is usually 1:100 or greater dilution for mouse sera with normal levels of IgG and IgM. Note: normal mouse sera may contain anti-pertussis FHA activity from prior exposure to the organisms.

- Run a Sample Diluent Blank. This signal is an indicator of proper assay performance, especially of washing efficacy, and is used for net OD calculations, if required (See Method A,B). Blank OD should be <0.3.
- Run a set of Calibrators. Calibrators validate that the assay was performed to specifications; results can be used to normalize between-assay variation for enhanced precision. Reading values off a Calibrator curve, Method C, has limitations. See Limits of the Assay.
- Run the Anti-FHA IgG Positive Control; value range is on label.
- Run a range of sample dilutions for expected higher positives that allows calculation of antibody Titer (when specific titer is at least 4-fold higher than non-immune). See Method D.
- Run samples in duplicate if used for quantitation; non-immunes that are significantly lower than immunes may be run in singlicate. The Calibrators that are used for quantitation, e.g., for between-assay normalization, should be run in duplicate. When determining titer from a dilution curve, singlicates can be run if more than two dilution points are used for titer calculations.

## Reagent Preparation

### Working Wash Solution

Dilute the entire volume 10ml + 990ml with distilled or deionized water into a clean stock bottle. Label as Working Wash Solution and store at ambient temperature until kit is used entirely.

### Working Sample Diluent

Dilute the entire volume, 10ml + 190ml with distilled or deionized water into a clean stock bottle. Label as Working Sample Diluent and store at 2-8°C until the kit lot expires or is used up.

### Working HRP Conjugate

Dilute fresh as needed; 10ul of concentrate to 1ml of Working Sample Diluent is sufficient for 1 8-well strip. Use within the working day and discard. Return 100X to 2-8°C storage.

## Assay Procedure

ALL STEPS ARE PERFORMED AT ROOM TEMPERATURE. After each reagent addition, gently tap the plate to mix the well contents prior to beginning incubation.

### 1. 1 st Incubation

- o Add 100ul of calibrators, samples and controls each to pre-determined wells.
- o Tap the plate gently to mix reagents and incubate for 60 minutes.
- o Wash wells 4 times and pat dry on fresh paper towels. As an alternative, an automatic plate washer may be used. Improper washes may lead to falsely elevated signals and poor reproducibility.

### 2. 2 nd Incubation

- o Add 100ul of diluted Anti-Mouse IgG HRP to each well.
- o Incubate for 30 minutes. o Wash wells 5 times as in step 1.

### 3. Substrate Incubation

- o Add 100ul TMB Substrate to each well. The liquid in the wells will begin to turn blue.
- o Incubate for 15 minutes in the dark, e.g., place in a drawer or closet.

Note: If your microplate reader does not register optical density (OD) above 2.0, incubate for less time, or read OD at 405-410 nm (results are valid).

#### 4. Stop Step

- o Add 100ul of Stop Solution to each well.
- o Tap gently to mix. The enzyme reaction will stop; liquid in the wells will turn yellow.

#### 5. Absorbance Reading

- o Use any commercially available microplate reader capable of reading at 450nm wavelength. Use a program suitable for obtaining OD readings, and data calculations if available.
- o Read absorbance of the entire plate at 450nm using a single wavelength within 30 minutes after Stop Solution addition. If available, program to subtract OD at 630nm to normalize well background.

## Calculation

Consider several data reduction methods to best represent the relationships among experimental and control groups, to determine Positive Immune and Negative Non-immune, and to Quantitate positive antibody levels.

#### Method A. Antibody Activity [ELISA Signal & Sample Dilution]

Represent data as net OD units (A450 signal; blank subtracted) ÷ dilution = Total Activity Units.

A Calibrator value in the mid-OD range (e.g., 5 U/ml) can be used to normalize inter-assay values.

#### Method B. Positive Index

Experimental sample values may be expressed relative to the values of Control or Non-immune samples, by calculation of a Positive Index. One typical method is as follows:

1. Calculate the net OD mean + 2 SD of the Control/Non-immune samples = Positive Index.
2. Divide each sample net OD by the Positive Index. Values above 1.0 are a measure of Positive Antibody Activity; below 1.0 are Negative for antibody.

A sample value would be Positive if significantly above the value of the pre-immune serum sample or a suitably determined non-immune panel or pool of samples, tested at the same sample dilution. This calculation quantifies the positive Antibody Activity level.

For Example:

Sample	Assay Net OD		Calculation using Positive Index	
	Control	Exptl	Control	Exptl
1	0.243	2.358	0.49	<b>4.79</b>
2	0.351	0.597	0.71	<b>1.21</b>
3	0.286	1.421	0.58	<b>2.89</b>
4	0.357	1.268	0.73	<b>2.58</b>
5	0.512	0.857	<b>1.04</b>	<b>1.74</b>
6	0.342	1.296	0.70	<b>2.63</b>
7	0.298	0.608	0.61	<b>1.24</b>
8	0.285	0.369	0.58	0.75
9	0.157	0.864	0.32	<b>1.76</b>
10	0.187	0.543	0.38	<b>1.10</b>
Mean	0.302			
SD	0.095			
Mean +2 SD	<b>0.492</b>		<b>= Positive Index</b>	

Method C. Use of a Calibrator Curve

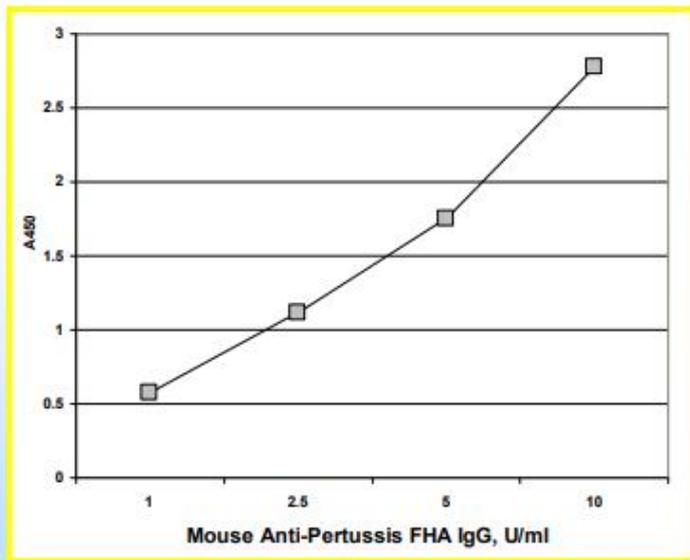
When the dilution curves of samples are parallel to the Calibrator curve (see Limits of the Assay), the Anti-FHA activity units may be determined by interpolation from the Calibrator curve. The results may be calculated using any immunoassay software package. If software is not available, Anti-FHA activity concentrations may be determined as follows:

1. Calculate the mean OD of duplicate samples.
2. On graph paper plot the mean OD of the calibrators (yaxis) against the concentration (U/ml) of Anti-FHA (x-axis). Draw the best fit curve through these points to construct the calibrator curve. A point-to-point construction is most common and reliable.
3. The Anti-FHA activity concentrations in unknown samples and controls can be determined by interpolation from the calibrator curve.
4. Multiply the values obtained for the samples by the dilution factor of each sample.
5. Samples producing signals higher than the 10 U/ml calibrator should be further diluted and re-assayed.

**Typical Results:**

Wells	Calibrators & Samples		A450 nm
A1,2	Negative Diluent Blank		0.12
B1,2	1 U/ml	Calibrator	0.58
C1,2	2.5 U/ml	Calibrator	1.11
D1,2	5 U/ml	Calibrator	1.75
E1,2	10 U/ml	Calibrator	2.78
F1,2	Sample	1:100	1.36

Sample Result: 3.3 U/ml x 100 dilution = 330 U/ml



**Method D. Titers from Sample Dilution Curves**

The titer of antibody activity calculated from a dilution curve of each sample is recommended as the most accurate quantitative method. Best precision can be obtained using the following guidelines:

1. Use an OD value Index in the mid-range of the assay (2.0 – 0.5 OD); this provides the best sensitivity and reproducibility for comparing experimental groups and replicates. An arbitrary 0 OD is commonly used.
2. Prepare serial dilutions of each sample to provide a series that will produce signals higher and lower than the selected index. With accurate diluting, duplicates may not be required if at least 4 dilutions are run per sample.
3. A 5-fold dilution scheme is useful to efficiently cover a wide range which produces ODs both above and below 1.0 OD. The dilution scheme can be tightened to 3-fold or 2-fold for more precise comparative data.
4. A Calibrator value in the mid-OD range (e.g., 5 U/ml) can be used to normalize inter-assay values.

**Calculations**

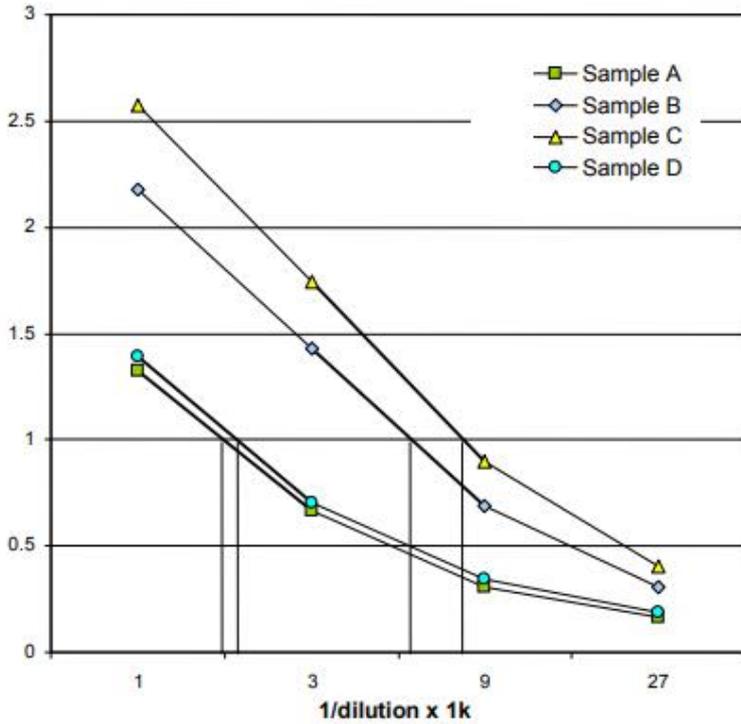
1. On a log scale of inverse of Sample Dilution as the x-axis, plot the OD values of the two dilutions of each

positive sample having ODs above and below the OD value of the Index (arbitrary or selected Calibrator).

- From a point-to-point line drawn between the two sample ODs, read the dilution value (x-axis) corresponding to the OD of the selected Index = IgG Antibody Activity Units

For example:

**II. A 1.0 OD Index was used to determine titer of 4 antibodies.**



Titer Values	
Sample A	= 1.72 kU
Sample B	= 5.70 kU
Sample C	= 1.85 kU
Sample D	= 7.90 kU

### Sensitivity

The FHA coating level, HRP conjugate concentration and Low NSB Sample Diluent are optimized to differentiate anti-FHA IgG from background (non-antibody) signal with mouse serum samples diluted 1:100.

### Specificity

Purified pertussis FHA (200 kDa) is used to coat the microwells; thus the assay is specific for antibodies directed to pertussis FHA. The anti-Mouse IgG HRP conjugate reacts with mouse IgG antibodies that bind to FHA on the plate. IgA, IgM and IgE class antibodies would not be measured above background signals.

### Precautions

Calibrators, Sample Diluent, and Antibody HRP contain bromonitrodioxane (BND: 0.05%, w/v). Stop Solution

contains dilute sulfuric acid. Follow good laboratory practices, and avoid ingestion or contact of any reagent with skin, eyes or mucous membranes. All reagents may be disposed of down a drain with copious amounts of water. MSDS for TMB, sulfuric acid and BND can be requested.

## Limitations

### Quantitation of Antibody in a Sample

The ELISA measures anti-FHA activity, a combination of antibody concentration and avidity for the FHA antigen. Antibodies with substantially different total IgG concentrations may display similar anti-FHA activities, due to differences in avidity. The quantitation or activity of the samples is, therefore, appropriately expressed in activity Units (titer), rather than mass units of IgG (e.g., ug/ml).

### Calibrator Curve Quantitation

To quantitate antibody activity from a calibrator curve (such as provided with the kit), the dilution curve of the samples must be parallel to the calibrator curve, to avoid different values being obtained from different regions of the curve. Antibodies that are not matched in anti-FHA avidity will often have non-parallel dilution curves. In these cases, antibody activity is best expressed as a titer relative to a reference positive such as the 5 U/ml Calibrator, or another Calibrator in the kit (see Calculation of Results).