Assay Performance Characteristics:

Standard range: 100-0.19ng/mL
Limit of Detection: 0.78ng/mL
Background: OD<0.1 at 450nm
Coefficient of Determination: R-squared>0.98

References:

1. Pollart SM, Smith TF, Morris E, Gelber LE, Platts-Mills TAE, Chapman MD. Environmental exposure to cockroach allergens: analysis with monoclonal antibody based enzyme immunoassays. J Allergy Clin Immunol 87:505-510, 1991.

2. Gelber LE, Seltzer LH, Bouzoukis JK, Pollart SM, Chapman MD, Platts-Mills TAE. Sensitization ad exposure to indoor allergens as risk factors for asthma among patients presenting to hospital. Amer Rev Respir Dis 147:573-578, 1993.

A list of frequently asked questions and troubleshooting guide can be found under the 'Support' tab on our web site: www.inbio.com.

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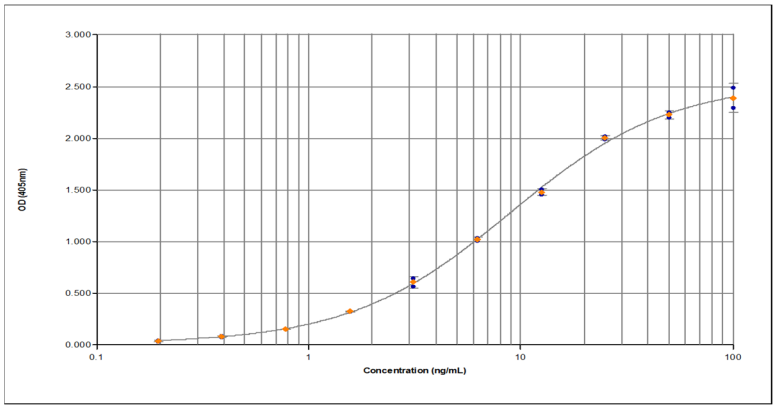
Bla g 2 ELISA 2.0
Pre-coated Plate Kit

Product Code: EPC-BG2-x
Lot Number: xxxxx

Sample curve:



Contents:

Microtiter plate coated with anti-Bla g 2 monoclonal antibody 7C11
Bla g 2 allergen standard (white cap)
Rabbit anti-Bla g 2 polyclonal antiserum (brown cap)
Peroxidase-conjugated goat anti-rabbit IgG (blue cap)
Wash buffer (10x concentrate)
Assay buffer (10x concentrate)
TMB developing substrate
Stop solution (0.5N sulfuric acid)

Store kit at 2-8^{O}C
Expiry:

For research and commercial use *in vitro*:
not for human *in vivo* or therapeutic use.

An InBio® product. Made in the USA.

Certificate of Analysis

Pre-coated Plate:	96-well polystyrene microtiter plate coated with monoclonal antibody 7C11 and treated with stabilizing agent. Sealed in foil pouch with desiccant.
Monoclonal Antibody:	7C11 (clone 7C11 C2 A6)
Immunogen:	Bla g 2
Isotype:	Mouse IgG1
Specificity:	Binds to an epitope on cockroach *Blattella germanica* allergen, Bla g 2.
Purification:	Produced in ascites and purified by ammonium sulphate precipitation and affinity chromatography using Protein A. Single heavy and light chain bands on SDS-PAGE.
Lot Number:	XXXXX

Detection Antibody:	Rabbit polyclonal antiserum
Immunogen:	Recombinant Bla g 2
Isotype:	Multiple
Specificity:	The antiserum contains IgG antibodies to cockroach *Blattella germanica* allergen, Bla g 2 and does not react with Bla g 1.
Activity:	Titrated for use in ELISA at 1/1000 dilution. Prepared in 1% BSA/50% glycerol/PBS, pH 7.4, 0.22 µm filtered, preservative free.
Lot Number:	XXXXX

Allergen Standard:	Purified natural Bla g 2 prepared in 1% BSA/50% glycerol/PBS, pH 7.4.
Concentration:	1,000ng/mL (based on amino acid analysis)
Lot Number:	XXXXX

Peroxidase-conjugated goat anti-rabbit:
Lot Number: XXXXX

Materials required, but not provided:

- Type I ultrapure water or 18.2MΩ de-ionized water
- Volumetric measuring equipment (e.g. serological pipettes, graduated cylinders)
- Clean containers for buffer and reagent preparation
- Reagent reservoirs
- Calibrated single and multi-channel micropipettes and tips
- Vortex mixer
- Plate reader capable of reading absorbance at 450nm
- Analysis software (recommended, but not required)

Protocol

Please read the entire protocol before starting the assay
Bring all reagents to room temperature before use

1. Prepare 1x working dilutions of the 10x wash and assay buffers in clean containers using 18.2MΩ de-ionized water or Type I ultrapure water. For one plate:
   Wash buffer: add 15mL concentrate to 135mL water
   Assay buffer: add 2.5mL concentrate to 22.5mL water
   Adjust volumes accordingly for multi-plate assays.
   *Diluted buffers may be stored at 4^{O}C for up to 1 week
2. Remove the plate from the foil pouch and wash by adding 150µL wash buffer to each well. Empty the wells by inverting the plate and then tap on absorbent paper to remove residual buffer. Repeat the wash cycle two more times.
3. Add standards, samples, and blanks to the plate (final volume in all wells is 100µL).

   Standards: add 180µL assay buffer into wells A1 and B1, and 100µL into remaining wells of rows A and B. Vortex the Bla g 2 standard and add 20µL to wells A1 and B1. Mix well by pipetting up and down 7-10 times and then transfer 100µL into wells A2 and B2. Mix well and continue the serial doubling dilution scheme across the plate to column 10.
   The assay buffer in wells A11, B11 and A12, B12 will serve as **Blanks.**
   Samples: Dust extracts are routinely tested starting at 1/10 dilution and may be prepared on the pre-coated plate: add 20µL sample to 180µL assay buffer in column A, mix, then transfer 100µL into 100µL assay buffer in the next well. Continue across the plate for the desired number of dilutions. A minimum of three dilutions per sample should be tested; 6-12 are recommended. Air filter extracts, allergen extracts, and other types of samples may require a different dilution scheme.
   *Sample dilutions may also be prepared in tubes or on a 96-well dilution plate and transferred to the pre-coated plate.
4. Incubate the plate at room temperature (away from direct sunlight) for 1 hour.
5. Wash the plate 3x with 150µL wash buffer per well. Vortex the polyclonal antibody and prepare a 1:1,000 detection antibody/conjugate mix by adding 10µL polyclonal antibody and 10µL peroxidase-conjugated goat ant-rabbit IgG to 10mL assay buffer.
   Mix thoroughly and add 100µL to each well.
6. Incubate the plate at room temperature (away from direct sunlight) for 1 hour.
7. Pour the TMB substrate and stop solutions into separate basins so they are ready to use in the next step. Wash the plate 3x with 150µL wash buffer per well.
8. Use a **multi-channel** pipette to add 100µL TMB to each well and monitor the reaction as the blue color develops. After 10-15 minutes, use a **multi-channel** pipette to add 50µL stop solution to each well (the color will change to yellow).
9. Read the plate at 450nm. The OD for Standard 1 should be between 1.2 and 3.5, with an ideal range of 2.0 - 2.5.