

## Assay Performance Characteristics:

Standard range: 500-0.98ng/mL  
 Limit of Detection: 1.95ng/mL  
 Background: OD<0.08 at 450nm  
 Coefficient of Determination: R-squared>0.98

## Plate Template:

	1	2	3	4	5	6	7	8	9	10	11	12
A												
B												
C												
D												
E												
F												
G												
H												

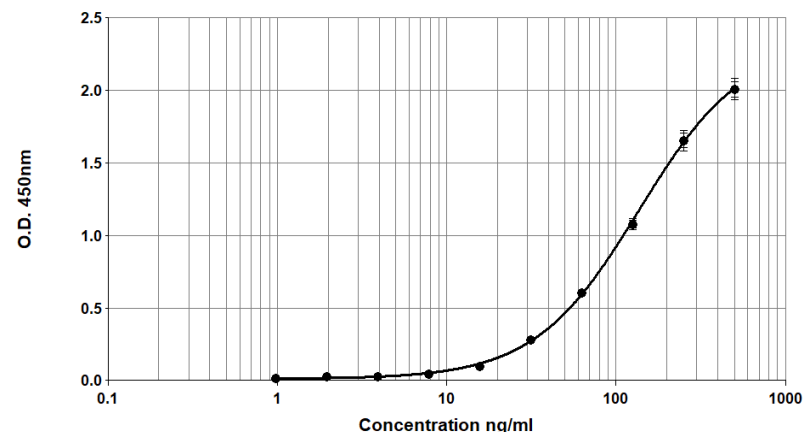
## References:



## Gal d 1 ELISA 2.0 Pre-coated Plate Kit

Product Code: EPC-GD1-X  
 Lot Number: xxxxx

### Sample curve:



### Contents:

Microtiter plate coated with anti-Gal d 1 monoclonal antibody 5G11

Gal d 1 allergen standard (white cap)  
 Concentration: 5,000ng/mL

Biotinylated monoclonal antibody 2B2 (brown cap)

Streptavidin-peroxidase (blue cap)

Wash buffer (10x concentrate)  
 Assay buffer (10x concentrate)  
 TMB developing substrate  
 Stop solution (0.5N sulfuric acid)

Store kit at 2-8°C  
 Expiry: xxxxx xx, xxxx

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

**An InBio® product**



Indoor Biotechnologies, Inc.  
 700 Harris Street,  
 Charlottesville, VA 22903  
 United States

Indoor Biotechnologies Ltd.  
 Vision Court  
 Caxton Place  
 Cardiff, Wales CF23 8HA  
 United Kingdom

Indoor Biotechnologies,  
 India Private Limited  
 Bangalore Bioinnovation Centre,  
 BioTech Park, Electronic City Phase 1  
 Bangalore-560100, India

Tel: (434) 984-2304  
 Fax: (434) 984-2709  
 Email: [mail@inbio.com](mailto:mail@inbio.com)

Tel: +44 (0) 29 2167 4640  
 Email: [Info@indoorbiotech.co.uk](mailto:Info@indoorbiotech.co.uk)

Tel: +91-9901722009  
 Email: [info@inbioindia.com](mailto:info@inbioindia.com)

## Certificate of Analysis

<b>Pre-coated Plate:</b>	96-well polystyrene microtiter plate coated with monoclonal antibody 5G11 and treated with stabilizing agent. Sealed in foil pouch with desiccant.
Monoclonal Antibody:	5G11
Immunogen:	Gal d 1
Isotype:	Mouse IgG1/kappa
Specificity:	Binds to species specific epitope present on <i>Gallus domesticus</i> allergen, Gal d 1.
Purification:	Produced in cell culture and purified by affinity chromatography using Protein G. Single heavy and light chain bands on SDS-PAGE.
Lot Number:	xxxxx

<b>Detection Antibody:</b>	2B2
Immunogen:	Gal d 1
Isotype:	Mouse IgG1/IgG2b/kappa
Specificity:	Binds to species specific epitope present on <i>Gallus domesticus</i> allergen, Gal d 1.
Purification:	Produced in cell culture and purified by affinity chromatography using Protein G. Single heavy and light chain bands on SDS-PAGE.
Biotinylation:	Biotinylated and titrated for use in ELISA at 1/1000 dilution. Prepared in 1% BSA/50% glycerol/PBS, pH 7.4, 0.22µm filtered, preservative free.
Lot Number:	xxxxx

<b>Allergen Standard:</b>	Purified natural Gal d 1 prepared in 1% BSA/30% glycerol/PBS, pH 7.4.
Concentration:	5,000ng/mL (based on amino acid analysis)
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
- 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°C for up to 1 week  
\*Use of an automated plate washer will require a larger volume of wash buffer than provided. Most PBS-based wash buffers should be compatible with the assay. Visit the support page at [www.inbio.com](http://www.inbio.com) for recommended wash buffer formulary.
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 Gal d 1 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 can be prepared directly on the pre-coated plate: add 20µL sample to 180µL assay buffer. 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 dilutions 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 biotinylated 2B2 and prepare a 1:1,000 detection antibody/conjugate mix by adding 10µL biotinylated 2B2 and 10µL streptavidin-peroxidase 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 solution 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. Gently tap the plate and monitor the reaction as the blue color develops. Once OD450 reaches 0.08-0.09 for Standard 1, 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.

*A list of frequently asked questions and troubleshooting guide can be found under the 'Support' tab on our web site: [www.inbio.com](http://www.inbio.com).*