



## DIAGNOSTIC AUTOMATION, INC.

23961 Craftsman Road, Suite E/F, Calabasas, CA 91302

Tel: (818) 591-3030 Fax: (818) 591-8383

[onestep@rapidtest.com](mailto:onestep@rapidtest.com)

[technicalsupport@rapidtest.com](mailto:technicalsupport@rapidtest.com)

[www.rapidtest.com](http://www.rapidtest.com)



See external label



2°C-8°C



$\Sigma=96$  tests



Cat # 1802-09

# Total Human IgA Assay

Cat # 1802-09

## Intended Use

To quantitate total human Immunoglobulin A (IgA)

## Principle of Procedure:

Solid phase capture sandwich ELISA assay using a microwell format.

## Shelf Life:

The expiration date for the package and each component is stated on the label(s). Store all components at 2°-8° degrees C. Store all components at 2-8°C with the exception of the standard, which should be stored frozen.

## Patient and Standard Dilutions:

Dilute each serum or plasma specimen to be tested 1:10,000 with the IgA specimen diluent provided. Prepare serial two fold dilutions of the human IgA standard: Neat, 1:2, 1:4, 1:8 etc. with the specimen diluent provided. Use the specimen diluent alone as the blank control well.

## Materials Supplied:

Anti-Human IgA coated microwell strips 12x8 with plastic frame  
HRP conjugated goat anti-human IgA -12mL  
IgA standard (pre-diluted 1:10,000)  
TMB/peroxide substrate color developer –12mL  
IgA specimen diluent (Specimen Diluent Green II) -60mL  
Sulfuric acid termination reagent (0.5N) –12mL  
15 X Wash buffer concentrate – 60mL

## Limitations of the Procedure:

No single assay should be used as the only basis for arriving at a diagnostic conclusion. For research use only.

## Dynamic Range:

0.031 µg/mL - 2µg/mL

## Reproducibility:

C.V. 6%-10% depending upon the region of the standard curve.

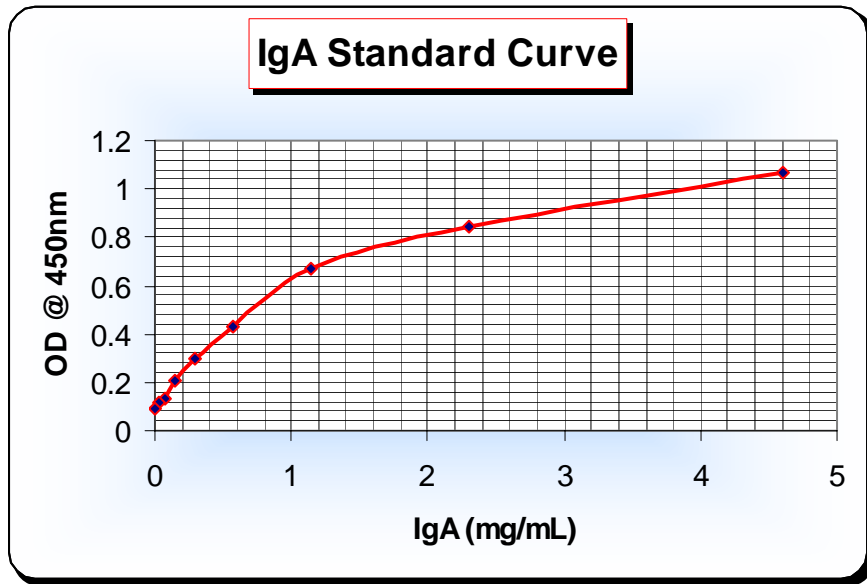
## Assay Procedure:

\*Allow each reagent to reach room temperature before use

1. Add 100uL of *diluted* specimen or standard to each microwell.
2. Incubate at room temperature for 60 minutes.
3. Decant and wash each microwell four times with wash buffer (dilute buffer 1:15 with reagent grade water).
4. Add 100uL of HRP conjugated goat anti-human IgA to each well.
5. Incubate at room temperature for 60 minutes.
6. Decant and wash as in step 3.
7. Add 100uL of TMB/peroxide substrate and incubate at room temperature for 30 minutes.
8. Terminate the reaction with 100uL of 0.5N sulfuric acid.
9. Zero the microwell reader at 450nm using the specimen diluent zero control well.
10. Determine the optical density (O.D.) of the remaining wells.
11. Construct a standard curve using the O.D. values obtained for each of the standards.
12. Interpolate the unknowns from the standard curve.

**Note: Constructed 4 parameter curve displayed below.**

Typical Standard Curve:



<b>Date Adopted</b>	<b>Reference No.</b>
2005-09-27	DA-Total Human IgA Assay-2008



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