



# ELISA for Mouse IgG

Product Code: 3825-1AD-6

#### **CONTENTS:**

## Vial 1 (yellow top)

Anti-IgG antibody (150 µl) Concentration: 0.5 mg/ml

### Vial 2 (green top)

ALP-conjugated anti-IgG antibody (80 µl)

#### Vial 3

Lyophilised mouse IgG standard

To ensure total recovery of stated quantity, vials have been overfilled.

#### **STORAGE:**

Shipped at ambient temperature. On arrival box 1 should be stored refrigerated at 4-8°C and box 2 should be stored frozen at -20°C.

# General

Intended use: For quantitative determination of native mouse IgG in serum and plasma.

**Reagents:** Anti-IgG antibody is supplied in sterile-filtered (0.2  $\mu$ m) PBS with sodium azide (0.02%). ALP-conjugated anti-IgG antibody is supplied in 0.1 M Tris-buffer with sodium azide (0.02%).

 $\textbf{Recommended standard dilution:}\ 0.1\text{-}500\ ng/ml$ 

## Guidelines for Mouse IgG ELISA

- Day 1 1. Coat a high protein binding ELISA plate with anti-IgG antibody, diluted to 1 μg/ml in PBS, pH 7.4, by adding 100 μl/well. Incubate overnight at 4-8°C.
- **Day 2** 2. Wash twice with PBS (200 μl/well).
  - 3. Block plate by adding 200 µl/well of PBS with 0.05% Tween 20 (PBS-Tween) containing 0.1% BSA (incubation buffer\*). Incubate for 1 hour at room temperature.
  - 4. Wash five times with PBS-Tween.
  - 5. Prepare mouse IgG standard by reconstituting contents of vial 3 in 500 µl PBS to a make up a stock solution of 50 µg/ml. The stock solution should be used immediately or stored in aliquots at -20°C for future use. We recommend the aliquots not be refrozen after initial use. For the test, prepare dilutions of the stock using the standard range as a guideline.
  - 6. Add 100 µl/well of samples or standards diluted in incubation buffer and incubate for 2 hours at room temperature.
  - 7. Wash as in step 4.
  - 8. Add 100 µl/well of anti-IgG-ALP diluted 1:1000 in incubation buffer. Incubate for 1 hour at room temperature.
  - 9. Wash as in step 4.
  - 10. Add 100 µl/well of appropriate substrate solution e.g. p-nitrophenyl-phosphate (pNPP).
  - 11. Measure the optical density (405 nm for pNPP) in an ELISA reader after suitable developing time.
    - \* The same buffer is used for blocking and for dilution.

NOTE; for research use only.









