

# FITC-labeled Goat Anti-Mouse IgG (H+L) FLUORESCENT CONJUGATE

**Catalog Number:** FL-07

**Quantity:** 500 micrograms

Format: PBS (0.14 M Sodium Chloride; 0.003 M Potassium Chloride; 0.002 M Potassium

Phosphate; 0.01 M Sodium Phosphate; pH 7.4), no preservative.

**Host:** Goat

## **Background:**

FITC-labeled goat anti-mouse IgG can be used to verify specific binding of mouse IgG to its receptor. By first incubating cells with the primary mouse antibody, and then binding the FITC-labeled goat anti-mouse IgG to the primary antibody, a fluorescent marker is formed that can demonstrate expression of a receptor or affinity of an antibody for its receptor. FITC is excited by 488 nm wavelength light, and emits at 525 nm.

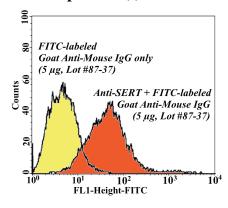
# **Specificity and Preparation:**

This fluorescent conjugate was prepared using goat anti-mouse IgG (H+L) and the fluorescent compound, fluorescein isothiocyanate (FITC). The antibody binds to mouse IgG, and is affinity-purified to decrease background and non-specific binding. This antibody exhibits maximal binding to mouse IgG antibodies, and minimal cross-reactivity with other molecules. This product is routinely tested by flow cytometry.

## **Usage and Storage:**

Applications include flow cytometry (ATS in-house;  $1 \mu g/10^6$  cells per  $200 \mu l$ ). Store at 4°C. DO NOT STORE FROZEN. The material may display diminished activity as a result of repeated freezing and thawing. Gently spin down material before use; 5-10 seconds in a microfuge should be adequate.

#### To view protocol(s) for this and other products please visit: www.ATSbio.com/protocols



A clone of RBL-2H3 cells, rat basophilic leukemia cells, were incubated with a protein-A purified mouse monoclonal primary antibody for one hour. The cells were then incubated for 30 minutes with FITC-labeled goat anti-mouse IgG secondary. Sample was read on a FACScan flow cytometer and data was analyzed with Cell Quest software.