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HORSE ANTI-MOUSE IGG ANTIBODY (H+L), FLUORESCEIN

SKU: FI-2000-1.5



DESCRIPTION

Fluorescein Horse Anti-Mouse IgG Antibody can be used for tissue staining or other applications. Optimal F/P ratios have been established for each conjugate to ensure maximum fluorescence with minimal background staining.

Features:

- Affinity-purified, ultrapure, high affinity antibody
- Thoroughly adsorbed against serum and immunoglobulins from potentially interfering species
- Recognizes both heavy and light chains (H+L)
- Optimally labeled with fluorescein to provide the brightest label for fluorescence microscopy
- Supplied in solution
- Excitation: 495 nm
- Emission: 515 nm

For research use only. Not intended for therapeutic or diagnostic use in animals or humans.

• Color: Green

SPECIFICATIONS

Color of Fluorescence	Green
Format	Concentrate
Formulation	10 mM HEPES, 0.15 M NaCl, pH 7.5, 0.08% sodium azide.
Maximum Emission	510-520 nm
Maximum Excitation	490-500 nm
Unit Size	1.5 mg
Storage Instructions	2-8 °C
Usage Summary	The recommended concentration range for use is 5-20 µg/ml. If this antibody is to be used in tissues which may contain cross-reacting endogenous immunoglobulins, dilution of this antibody may be made in buffers containing 2 % normal serum from the same species as the tissue.
Applications	Immunofluorescence, In situ hybridization, Blotting Applications, Flow Cytometry/Cell Separation
Target Species	Mouse
Concentration	1.5 mg active conjugate/ml
Conjugate	Fluorescein
Reactive Species	Horse
Source Species	Mouse
Host Species	Horse

TECHNICAL INFORMATION

The anti-mouse Ig antibodies are prepared by hyperimmunizing animals in a manner that produces high affinity antibodies. These are then purified by an affinity chromatography procedure designed to remove any low affinity antibodies which may be present. Cross-reactivities that are likely to interfere with specific labeling are removed by solid-phase adsorption techniques. The final product is then subjected to rigorous quality control assays including immunodiffusion, solid-phase enzyme immunoassays, gel electrophoresis and solid-phase binding assays. In preparing the labeled antibodies, great care is taken to ensure the maximum degree of labeling with no alteration in the specificity and affinity of the antibody. The

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labeled antibody then undergoes a further series of quality control assays, including immunohistochemical analysis.

CITATIONS

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DOCUMENTS

- Safety Data Sheet
- Download CoA
- Datasheet

GALLERY IMAGES



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