



GOAT ANTI-HUMAN IGE, EPSILON CHAIN SPECIFIC

SKU: FI-3040-.5



DESCRIPTION

Fluorescein Goat Anti-Human IgE, epsilon chain specific, can be used for immunofluorescence and 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
- Optimally labeled with fluorescein to provide the brightest label for fluorescence microscopy
- Supplied in solution
- Excitation: 495 nm
- Emission: 515 nm
- Color: Green
- This chain-specific antibody is produced specifically to distinguish between chains or classes of target immunoglobulins. This chain specific antibody has virtually no cross-reactivity with other immunoglobulin classes or other heavy or light chains.

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



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	0.5 mg
Storage Instructions	2-8 °C
Usage Summary	The recommended concentration range for use is 5-20 µg/ml. If this fluorescein-labeled 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, Blotting Applications, Flow Cytometry/Cell Separation
Target Species	Human
Concentration	1.0 mg active conjugate/ml
Conjugate	Fluorescein
Reactive Species	Goat
Source Species	Human
Host Species	Goat

TECHNICAL INFORMATION

The goat anti-human 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 labeled antibody then undergoes a further series of quality control assays, including immunohistochemical analysis.

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CITATIONS



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DOCUMENTS

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