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# Datasheet for 109-1102 Human IgG (H&L) Antibody

#### **Overview**

| Description:         | Goat Anti-Human IgG (H&L) Antibody - 109-1102 |
|----------------------|---|
| Item No.:            | 109-1102                                      |
| Size:                | 2 mL  |
| Applications:        | ELISA, Microarray                             |
| Reactivity:          | Human   |
| <b>Host Species:</b> | Goat  |

## **Product Details**

Background: Anti-Human IgG (H&L) generated in goat detects human Immunoglobulin G (IgG), both heavy

and light chains of the antibody molecule are present. It is a protein complex composed of four peptide chains — two identical heavy chains and two identical light chains arranged in a Y-shape typical of antibody monomers. Each IgG has two antigen binding sites. Representing approximately 75% of serum immunoglobulins in humans, IgG is the most abundant antibody isotype found in the circulation. IgG molecules are synthesized and secreted by plasma B cells. Secondary Antibodies are available in a variety of formats and conjugate types. When choosing a secondary antibody product, consideration must be given to species and immunoglobulin

specificity, conjugate type, fragment and chain specificity, level of cross-reactivity, and host-species source and fragment composition.

Synonyms: goat anti-Human IgG Antibody, goat anti Human IgG

Host Species: Goat

Specificity: IgG (H&L)

Clonality: Polyclonal

Format: Antiserum

## **Target Details**

Reactivity: Human

Immunogen: Human IgG whole molecule

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**Purity/Specificity:** This product was prepared from monospecific antiserum by a delipidation and defibrination.

Assay by immunoelectrophoresis resulted in a single precipitin arc against anti-goat serum,

Human IgG and Human Serum.

## **Application Details**

| Suggested Applications: | ELISA, Microarray (Based on references)   |
|-------------------------|---|
| Application Note:       | Secondary antibody reagents are ideal for ELISA, western blotting, Immunohistochemistry, Fluorescence Microscopy, Flow Cytometry as well as other antibody detection methods. |
| Assay Dilutions:        | All assays should be optimized by the user. Recommended dilutions (if any) may be listed below.   |
| ELISA:                  | 1:20,000 - 1:100,000  |
| IHC:                    | 1:1,000 - 1:5,000   |
| WB:                     | 1:2,000 - 1:10,000  |

## **Formulation**

| Physical State:        | Lyophilized   |
|------------------------|---|
| Concentration:         | 90 mg/mL by Refractometry                               |
| Buffer:                | 0.01 M Sodium Phosphate, 0.15 M Sodium Chloride, pH 7.2 |
| Preservative:          | None  |
| Stabilizer:            | None  |
| Reconstitution Volume: | 2.0 mL  |
| Reconstitution Buffer: | Restore with deionized water (or equivalent)            |

## **Shipping & Handling**

| Shipping Condition: | Ambient   |
|---------------------|---|
| Storage Condition:  | Store vial at 4° C prior to restoration. For extended storage aliquot contents and freeze at -20° C or below. Avoid cycles of freezing and thawing. Centrifuge product if not completely clear after standing at room temperature. This product is stable for several weeks at 4° C as an undiluted liquid. Dilute only prior to immediate use. |
| Expiration:         | Expiration date is one (1) year from date of receipt.   |

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#### References

- Serrano-Coll, H et al. Social and environmental conditions related to Mycobacterium leprae infection in children and adolescents from three leprosy endemic regions of Colombia. *Bmc Infectious Diseases* (2019)
- Metz I, Beißbarth T, Ellenberger D, et al. Serum peptide reactivities may distinguish neuromyelitis optica subgroups and multiple sclerosis. *Neurol Neuroimmunol Neuroinflamm*. (2016)
- Bucukovski, J et al. A Multiplex Label-Free Approach to Avian Influenza Surveillance and Serology. PloS One (2015)

## **Disclaimer**

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