

## PRODUCT DATA SHEET

### Anti-Mouse IgG (H+L), Rat Serum Adsorbed - Gold Conjugate



**Anti-Mouse**

#### Description

Affinity isolated rat serum adsorbed anti-mouse IgG antibody produced in goat and coupled to gold nanoparticles. Suitable for use in applications such as lateral flow, immunoblotting, light microscopy, and electron microscopy applications procedures for secondary detection of mouse antibody labeled samples.

Provides a permanent and sensitive label when used separately or in conjunction with Cytodiagnostics membrane and microscopy silver enhancer kits, see related product below.

**Concentration:** 0.15 mg/ml (@ OD=3), 0.5mg/ml (@ OD=10)

**Conjugated Antibody:** Goat affinity purified anti-mouse IgG (H+L), rat serum adsorbed

**Clonality:** Polyclonal

**Storage Buffer:** 20mM Tris (pH 8.0), 150mM NaCl, 20% glycerol (v/v), 1% BSA

**Working Dilution:** 1:10 – 1:100 (application dependent, optimization might be required)

#### Storage

Store undiluted in storage buffer at 2-8° C. Stable for at least 4 months if stored as specified.

**DO NOT FREEZE.**

#### Product Safety and Handling

This product is for R&D use only, not for drug, household, or other uses. Please review the safety datasheet (SDS) available online for proper safety and handling procedures.

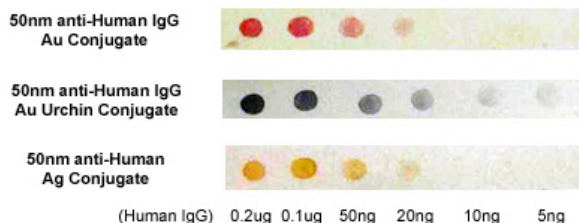
#### Related Products

Silver Enhancer Kit for Membranes Cat No. SR-01-02  
Silver Enhancer Kit for Microscopy Cat No. SR-01-01

#### Standard Immunogold Dot-Blot Protocol

*(Adapted from Moeremans et al. [1])*

1. Spot one microlitre drops of a serial dilution of your protein (1ug-1ng) in PBS supplemented with 0.5 ug/ml of BSA on nitrocellulose or PVDF membrane.
2. Let protein drops dry into the membrane.
3. Block Membrane for 30 minutes using 1% (w/v) dry milk in 1X PBS at room temperature.
4. Incubate with primary antibody for 2 hours at room temperature.
5. Wash membrane 3x5 minutes with blocking solution prepared as above.
6. Incubate for 2 hours (or longer for increased sensitivity) with secondary gold conjugate diluted 1:10 (OD=0.3) times with blocking solution (0.2% Blocking Solution).
7. Wash 3x5 minutes as above.
8. Dry membrane and record data.
9. (OPTIONAL) Proceed with silver enhancement to improve sensitivity.



**Figure 1.** Example dot-blot assay for Cytodiagnostics streptavidin gold conjugate (top left) and our streptavidin silver conjugate (top right) before and after enhancement using Cytodiagnostics silver enhancement kit for membranes. Bottom picture illustrates and highlights the difference in appearance (color) of 50nm anti-human IgG noble metal nanoparticle conjugates prepared using NHS-activated gold nanoparticles, NHS-activated gold nanourchins, and NHS-activated silver nanoparticles, respectively.

## References

1. M. Moeremans, et al., Journal of Immunological Methods, 1984, 74, 353

## Ordering Information

For ordering call 866-344-3954 or visit us online.