

### **PRODUCT DATA SHEET**

Protein G - 30nm Au20/Ag80 Alloy Conjugate (0.5ml)

Catalog No. AAC-30-18-05 Size: 0.5ml

#### Description

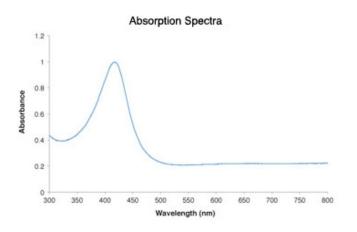
Protein G conjugated 30nm Au20/Ag80 Alloy nanoparticles. Suitable for use in immunoblotting, light microscopy, and electron microscopy applications.

Provides a permanent and sensitive label (100pg or less) when used in conjunction with Cytodiagnostics membrane and microscopy silver enhancer kits, see related product below.

**Concentration:** 0.15 mg/ml (OD=3)

**Conjugated Protein:** Protein G from *Streptococcus sp.* (expressed in *E. coli*)

**Storage Buffer:** 10mM PBS (pH 7.4), 20% glycerol (v/v), 1% BSA **Working Dilution:** 1:10 – 1:100



#### Storage

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

#### **Product Safety and Handling**

This product is for R&D use only, not for drug, household, or other uses. Please review the material safety datasheet (MSDS) 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

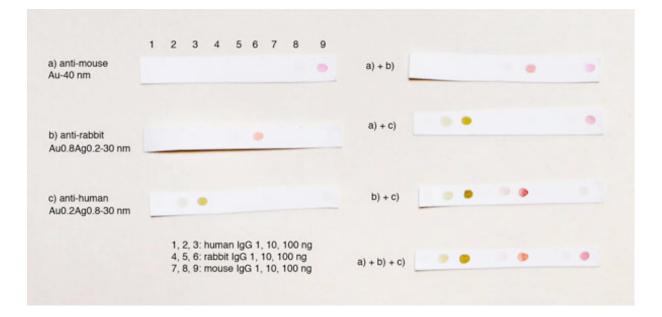
## Standard Immunogold Dot-Blot Protocol

(Adapted from Moeremans et al. [1])

- Spot one microlitre drops of a serial dilution of your protein (100-0.1ng) in PBS supplemented with 50ug/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.
- Incubate for 2 hours (or longer for increased sensitivity) with secondary alloy conjugate diluted 1:10 (OD=0.3) times with blocking solution (0.2% Blocking Solution). See Tech Note #102 for preparation of an alloy conjugate.
- 7. Wash 3x5 minutes as above.
- 8. Dry membrane and record data.
- 9. (OPTIONAL) Proceed with <u>silver</u> <u>enhancement</u> to improve sensitivity.

919 Fraser Drive Unit 11, Burlington, ON, L7L 4X8, Canada Tel: 866-344-3954 Fax: 289-288-0122 www.cytodiagnostics.com

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**Figure 1.** Dot-blot assay using three different types of noble metal nanoparticle conjugates, I.e. anti-mouse IgG 40nm gold conjugate (top left), anti-rabbit IgG Au20/Au80 30nm alloy conjugate (middle left) and anti-human IgG Au80/Au20 30nm alloy conjugate (bottom left). Simultaneous multiplex detection of two or more of the antigens are shown in the right column.

#### 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 at www.cytodiagnostics.com

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