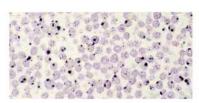
GoldEnhanceTM

Go beyond silver with GOLD enhancement!



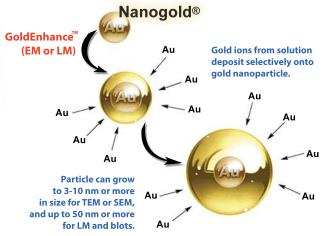


with GoldEnhance™
In situ hybridization detection of HPV-16 DNA with CARD-Cy3 and Nanogold®-Streptavidin.

Control with biotin-tyramide omitted

Get deep penetration with our ultra-small gold labels...

Develop with GoldEnhance for incredible imaging!



GoldEnhance™ EM Plus 8mL kit

Our new favorite for EM!

- Slow enhancement for easy size control
- Better for SEM: Much better backscatter signal than silver
- Low background
 No autonucleation for 40 minutes
- Neutral pH for best ultrastructural preservation
- Safe to use before osmium tetroxide etching Silver would dissolve

GoldEnhance™ LM

60mL kit

for light microscopy and blots

- Minimal autonucleation even after 1-2 hours
 Convenient for multiple samples
 and automated processing
- Observe with brightfield optics! simpler and less expensive than fluorescence
- Much faster than chemiluminescence
- Permanent staining: does not fade
- Low viscosity, Mix-and-use

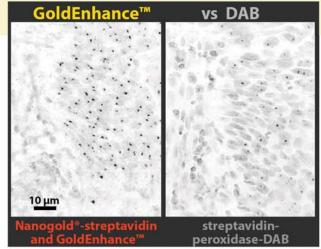
Why Gold?

lanogol@

GoldEnhance

Better than silver for many applications!

- · Lower backgrounds than silver in many cases
- Light insensitive! Permanent staining Does not fade
- Better for SEM -much better backscatter signal than silver
- Compatible with physiological buffers like PBS, halides silver would precipitate
- · Reaction is less pH sensitive than silver
- Near neutral pH for best structural preservation many silver enhancers have a pH of 3-4
- Can be used for specimens on metal surfaces (e.g. cell culture substrates)
- · Excellent shelf life, easy mixing with low viscosity



Formalin-fixed serial sections of cervical carcinoma, in situ hybridized for HPV-16/18 using a biotinylated probe. Courtesy G. W. Hacker, University of Salzburg.



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Questions? Ask our scientists: tech@nanoprobes.com