Exploiting the interaction between live *E. coli* and citrate Au nanoparticles to form a label-free diagnostic test for bacterial contamination

Camilla Gazzana, Stella Valenzuela, Andrew McDonagh, Michael Cortie

15 Broadway, Ultimo, 2007 School of Mathematical and Physical Sciences, University of Technology Sydney Sydney, NSW, Australia

camilla.gazzana@uts.edu.au, michael.cortie@gmail.com

There is a need for sensitive and robust label-free detection of bacteria in food and environmental samples. Here we show how colloidal gold nanoparticles can be combined with the spontaneous multiplication of live *Escherichia coli* bacteria in a growth medium to produce an enhanced optical signal that is sensitive down to ca. 10 CFU/mL of live bacteria. The assay exploits the insight that an *in situ* suspension of nanoparticles in a growth solution amplifies the optical changes caused by multiplication of the bacteria. The method relies upon the change in optical spectrum as gold nanoparticles aggregate onto the cell walls of a proliferating population of planktonic bacteria. The sensitivity is comparable to that of many state-of-theart techniques used in the field. The method does not detect dead or inactive bacteria. Since there is no need for an expensive and heat-sensitive antibody, the technique that we describe may offer some advantages for testing in the field.

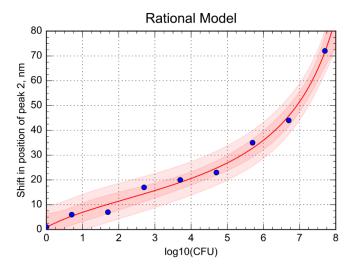


Figure 1: Rational model ($R^2 = 0.9927$) of the proposed AuNP sensor whereby the dark pink regions show the 95% confidence interval for the average log10(CFU/mL) of the bacterial test solution and the light pink regions show the 95% prediction interval for any individual future calculation as a function of the shift in the position of the second plasmon peak in nm.

References:

1. Pissuwan, D.; Gazzana, C.; Mongkolsuk, S.; Cortie, M. B., Single and multiple detections of foodborne pathogens by gold nanoparticle assays. *Wiley Interdisciplinary Reviews: Nanomedicine and Nanobiotechnology* **2020**, *12* (1), e1584.