

Plasmonic gold biosensors for quantitative determination of aflatoxins in corn matrix

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We developed a new selective sensor for detection of aflatoxins. Aflatoxins are carcinogenic and harmful for gene and occur in nature. The aim of our work to detect this toxin with selective sensors. We use functionalised gold nanoparticles and gold nanofilms and the surface modification of gold surface was made with cyclodextrines which are modified by thiol groups.

Sodium borohydride reduced gold nanoparticles in presence of modified cyclodextrin by thiol groups (AuSH- β -CD-NaBH₄): Materials used for the preparation of Au nanoparticles were: HAuCl₄*3H₂O (Sigma-Aldrich), sodium borohydride (Sigma), MQ water and SH-modified cyclodextrin (Cyclolab Ltd.). Identification and characterization of sodium borohydride reduced gold nanoparticles were performed by their XRD diffractograms, UV-Vis spectra and TEM images. We could establish uniform conductive films on silica crystal and glass surfaces by spray-coating technology from the prepared gold nanodispersion if the SiO₂ surfaces were treated previous by PEI (positive charged layer). These specifically produced films are suitable for biosensors.

The attachment of aflatoxin molecules to bare gold surface and to those covered gold thin films was studied by SPR and QCM measurements. The adsorbed amount of gold surface (full coverage ~ 140 ng/cm² was achieved at 0,05 mg/ml SH- β -CD) SH- β -CD was attached. AFB₂ molecules could attach neither to gold surface nor AuSH- β -CD-NaBH₄ covered surface contrary to AFB₁ (~ 50 ng AFB₁ attach to 1 cm² pure gold surface, while if it is covered by AuSH- β -CD-NaBH₄. The adsorbed amount is ~ 530 ng AFB₁).

For comparison a quantitative analysis of aflatoxin molecules on surfaces of modified gold nanoparticles will be done by ELISA (Enzyme-Linked ImmunoSorbent Assay) method. Validation of thiol modified gold nanosensors on silica crystal will be performed by HPLC-MS method. To compared the ELISA results against the SPR measurement, we established that the Toxi watch ELISA kit for AFB₁ sensitivity limit is 30 ppt and the SPR sensitivity limit is (54 nm/ RIU) ~ 1 ppt.

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References

- [1] J. Turkevich, *Gold Bulletin*, 1985, 18, 3
- [2] F. Fernández, K. Hegnerová, M. Piliarik, F. Sanchez-Baeza, J. Homola, M.-P. Marco, *Biosensors and Bioelectronics*, 26, 4, pp. 1231-1238, 2010
- [3] E. Pál, V. Hornok, D. Sebők, A. Majzik, I. Dékány, *CollSurfB*, 79, pp. 276–283, 2010
- [4] J. H. Kim, K. S. Kim, K. M. Manesh et al., *CollSurfA*, 313-314, pp. 612-616, 2008