

XIX CONGRESO DE LA SOCIEDAD ARGENTINA DE MICROBIOLOGÍA GENERAL

22 al 25 de octubre del 2024

Centro cultural y Pabellón Argentina de la Universidad Nacional de Córdoba, Córdoba, ARGENTINA.



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GOLD@N-BUTIL POLYCYANOACRYLATE NANOSPHERE AS A COATING WITH ANTIBACTERIAL AND ANTI-BIOFILM PROPERTIES

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Bacterial adhesion and biofilm formation on biomaterials represent serious economic and health problems. Biofilms, whether mono-microbial or poly-microbial, exhibit significant resistance to antimicrobials and host defenses. Although antibiotics are a common treatment and prevention strategy, their effectiveness decreases due to the resistance induced by biofilms and the increase in multi-resistant strains. As an alternative, new materials and coatings are being developed. Nano-structured coatings, such as gold nanoparticles (Au-NPs), have intrinsic antimicrobial properties. Additionally, alkyl polycyanoacrylates (PCA) are biocompatible polymers with antimicrobial activity against Gram-positive and some Gram-negative bacteria. This study synthesizes and evaluates a coating based on gold@polycyanoacrylate n-butyl nanoparticles (Au-PCAB-Nps) to combat biofilms on biomaterials. Au-PCAB-Nps were synthesized by nanoprecipitation using a non-ionic surfactant as a stabilizer. They were characterized by Transmission Electron Microscopy, with hydrodynamic radius determined by Dynamic Light Scattering and Z-potential measured by Electrophoretic Light Scattering. The viability of *E. coli* 144, *P. mirabilis* 2921, *S. aureus* ATCC 6538, *S. epidermidis* M20200221, *A. baumannii* ATCC 19606, and *P. aeruginosa* ATCC 902 against the nanoparticles (Nps) was assessed using resazurin assays. The ability of the Nps to inhibit and eradicate biofilms was evaluated in a static assay on 96-well plates, using concentrations of 1x, 2x, and 4x. The obtained dispersion revealed two nanoparticle populations with hydrodynamic diameters of 79 ± 1 nm and 167 ± 2 nm, and a polydispersity index of 0.169. The observed Z-potential was -18.15 ± 0.02 mV. Au-PCAB-Nps exhibited minimum inhibitory concentration and minimum bactericidal concentration values above 4x for all tested strains. The viability assay showed

that Au-PCAB-Nps had no cytotoxic effect on *E. coli*, *P. mirabilis*, *S. aureus*, *S. epidermidis*, and *A. baumannii* under the tested conditions (24 h, concentrations of 0.002x to 4x), but did show cytotoxic effects on *P. aeruginosa* at 24 h and concentrations of 0.5x to 4x. Compared to Au-Nps, Au-PCAB-Nps had a similar effect, while PBCA-Nps showed no cytotoxic effects on any of the evaluated strains. Au-PCAB-Nps significantly reduced biofilm formation of *E. coli*, *P. mirabilis*, *S. aureus*, *S. epidermidis*, and *A. baumannii* compared to the control. PBCA-Nps showed a similar effect, while Au-Nps had no impact under any of the studied conditions. However, Au-PCAB-Nps did not reduce the biomass of mature biofilms of the studied bacteria, a behavior also observed with Au-Nps. PBCA-Nps reduced the biomass of mature biofilms only in *P. mirabilis* and *S. aureus*. These results suggest that Au-PCAB-Nps have promising potential as a coating for biomaterials to prevent bacterial biofilm-related infections. However, their efficacy does not surpass that of PBCA-Nps and is comparable to Au-Nps.

Palabras clave: Biofilms - Nanoparticles - Gold - n-Butyl Polycyanoacrylate - Coating Materials