Antimicrobial Activity of Biosynthesized Silver Nanoparticles Compared to Standard Antibiotics Used in ORL Infections

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In recent years, bacterial infections in hospitals have grown particularly due to the development of antibiotic resistance. Recent research targets the discovery of new antibiotics that exhibit broad spectrum of action without adverse effects or minimizing adverse effects. In this study, the activity of biosynthesized silver nanoparticles against three bacteria commonly found in infectious diseases in the ORL sphere was evaluated. The recorded data revealed an activity comparable to that of the standard antibiotics used in these types of infections, with the observation that the activity of the nanoparticles could also be observed in the particular cases of antibiotic resistance.

Keywords: green silver nanoparticles, antibiotics, resistance, bacterial strains

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Experimental part

Materials and methods

The reagents used in the present study were of analytical grade, acquired from Sigma Aldrich, Germany and utilized as received without any other supplementary purification. Green silver nanoparticles were obtained as previously described by Andor et al.\textsuperscript{[8]}. Briefly, to an aqueous rosemary leaves extract (RAE) - recognized both as a reducing and stabilizing agent - was added an aqueous solution of AgNO\textsubscript{3} (1 mM) and the reaction mixture was inserted in an orbital shaker (away from light, for two days, 250 rpm, at room temperature), then purified and utilized in the in vitro experiments.

Preclinical testing. Green silver nanoparticles obtained with an aqueous extract of rosemary leaves were tested at different concentrations (0.1-10 \textmu g/mL) on three bacterial strains. Reference strains are presented in table 1.

\begin{table}[h]
\centering
\begin{tabular}{|l|l|l|}
\hline
Bacterial species & Code & Producer\\
\hline
\textbf{Pseudomonas aeruginosa} & ATCC\textsuperscript{TM} 27853\textsuperscript{TM} & Thermo Fischer Scientific\\
\textbf{Streptococcus pneumoniae} & ATCC\textsuperscript{TM} 27336\textsuperscript{TM} & Thermo Fischer Scientific\\
\textbf{Haemophilus influenzae} & ATCC\textsuperscript{TM} 39141\textsuperscript{TM} & Thermo Fischer Scientific\\
\hline
\end{tabular}
\caption{REFERENCE BACTERIAL STRAINS UTILIZED IN THE PRESENT STUDY}
\end{table}

\[\text{Table 1} \]

Multiplication of infections in hospitals due to treatment-resistant bacteria in which the usual antibiotics are no longer effective has become an alarming problem and a real danger to human health. The complex mechanism involved in the development of antibiotic resistance, even in the case of new generations, has not been fully elucidated. This requires additional studies to optimize diagnosis, find new therapies without side-effects or with minimal side effects and / or reduce abusive drug use [1, 2]. Biofilms are microbial formations wrapped in a polymeric matrix, with the major components exopolysaccharides. Microbes grow and develop in this matrix due to nutritional resources and protection against environmental factors. Biofilms are sources of pathogenic organisms that trigger disease. They are directly involved in the occurrence of ORL disorders, such as otitis media and otolaryngeal infections [3]. \textit{Pseudomonas aeruginosa}, is a gram negative bacteria, a pathogen responsible for the occurrence of infections (sepsis, endocarditis, etc.) that has become resistant to frequently used antibiotic treatment, responsible for high rates of morbidity and mortality. \textit{Haemophilus influenzae} is a gram-negative bacterium that triggers respiratory tract complications, and is known as a respiratory and invasive pathogen. The main diseases caused by \textit{H. influenzae} in children are pneumonia, meningitis and bacteremia, and in adults are pneumonia, acute otitis media, acute sinusitis and acute exacerbation of chronic bronchitis[4]. \textit{Streptococcus pneumoniae} is a gram-positive bacteria that colonizes the mucosal surfaces of the human upper respiratory tract and is the main cause of otitis media, community pneumonia, sepsis and meningitis [5].

The chemical synthesis methods applied to the metal nanoparticles most often require the use of toxic reagents, solvents, chemical coating agents, and special synthetic conditions that lead to increased production costs and environmental pollution. An inexpensive and safe alternative is that of nanoparticle biosynthesis [6]. Silver nanoparticles have captured the researchers' attention due to the remarkable properties of silver ions which in metallic form are inert, but in ionized form they become very reactive [7]. This study aimed testing the antimicrobial activity of biosynthesized silver nanoparticles compared to antibiotics used for infections produced by \textit{Pseudomonas aeruginosa}, \textit{Haemophilus influenzae} and \textit{Streptococcus pneumoniae} by different methods, namely: a) preclinical testing-on bacterial strains by disk diffusion method and anti-biofilm activity and b) clinical testing-classic antibiotics on biological samples originated from current clinical activity.

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Disk Diffusion Method

The antimicrobial activity of the silver nanoparticles was assessed by the agar disk diffusion method, as specified in the literature [9]. The experiments were realized in triplicate. The positive controls utilized in the current study, classic antibiotics (Fig. 1), were: penicillin (CT0042B 1.5 units), cefepime (CT0771B conc. 30 µg), ceftriaxone (CT0417B conc. 30 µg), erythromycin (CT0021B conc. 30 µg), piperacillin (CT0261B conc. 75 µg) and ticarcillin (CT0167B conc. 75 µg) disks (Thermo-Fischer Scientific) and the negative control, was represented by the blank paper disk treated with the solvent employed in the experiments.

Anti-biofilm activity

The colorimetric test is often utilized in order to evaluate the activity of different compounds on biofilm. The steps required to apply the technique are as follows: (i) $10^6$/mL bacterial strain are cultured in 96-well plates, (ii) various concentrations of RAE_AgNPs are added after 24 h, (iii) the plates are incubated with test samples for 4 h at 37°C, (iv) the wells are washed, dried and treated with crystal violet aqueous solution, and (v) final washings and absorbance reading at 595 nm using an ELISA reader. The percentage inhibition of biofilm activity was calculated by the formula presented in the literature [10].

Clinical testing

Biological samples originated from current clinical activity, were harvested according to the internal protocol prior to antimicrobial administration. Bacterial isolation was performed on Columbia agar with blood and Chocolat (Sanimed, Romania) agar and the plates were incubated at 37°C. Api NH (bioMerieux, France) kits were used to identify Haemophilus influenzae strains while for the rest of the isolated species the Vitek 2 Compact automatic analyser (bioMerieux, France) was employed. Sensitivity testing for anti-infectious chemotherapies was performed by diffusion and E-tests (AB Biodisk, Sweden) according to the CLSI standard.

Results and discussions

The antibacterial activity of silver nanoparticles compared to classic antibiotics used as positive control was evaluated on three bacterial strains. The *Pseudomonas aeruginosa* and *Streptococcus pneumoniae* were significantly influenced by silver nanoparticles. The zone of inhibition diameters are noted in Table 2.

RAE_AgNPs were utilized in order to inhibit the activity of biofilms. In this research, the assessment of nanoparticles on biofilms activity in the human pathogens *P. aeruginosa*, *S. pneumoniae* and *H. influenzae* was evaluated by in vitro methods. The test strains were treated with various concentrations of RAE_AgNPs (0.1 to 10 µg/mL) and data showed that, green silver nanoparticles prevented the activity of biofilms, significant even from a concentration of 1 µg/mL, as can be seen in the figure 2.

Treatment of *Pseudomonas aeruginosa* for 24 h with 1 µg/mL of RAE_AgNPs decreased biofilm activity by almost

| Positive control/Test compounds | *Pseudomonas aeruginosa* | *Streptococcus pneumoniae* | *Haemophilus influenzae* |
|---------------------------------|--------------------------|---------------------------|-------------------------|
| Penicillin                      | 16 mm                    | 6 mm                      | 12 mm                   |
| Cefepime                        | 14 mm                    | 15 mm                     | 15 mm                   |
| Ceftriaxone                     | 12 mm                    | 16 mm                     | 15 mm                   |
| Erythromycin                    | 10 mm                    | 11 mm                     | 8 mm                    |
| Piperacillin                    | 8 mm                     | 15 mm                     | 9 mm                    |
| Ticarcillin                     | 7 mm                     | 15 mm                     | 7 mm                    |
| RAE_AgNPs                       | 19 mm                    | 16 mm                     | 14 mm                   |

Table 2

ZONE OF INHIBITION (mm) OF POSITIVE CONTROLS AND TESTED SAMPLES AGAINST THREE BACTERIAL STRAINS
40% while in case of a higher concentration, 5 µg/mL the percentage of inhibition is almost 80%. In case of evaluation of the *Streptococcus pneumoniae* and *Haemophilus influenzae* biofilm activity, inhibition values were close, around 60% at a concentration of 5 µg/mL, respectively about 80% at a concentration of 10 µg/mL.

In the case of clinical trials, only positive controls were used. 20 bacterial strains were identified, including: 13 - *Streptococcus pneumoniae*, 4 - *Haemophilus influenzae* and 3 - *Pseudomonas aeruginosa*. Each of the three identified batteries showed resistance to the antibiotics tested as follows: 4 *Streptococcus pneumoniae* strains showed resistance to penicillin, but they kept their sensitivity to ceftriaxone and cefepime; 7 *Streptococcus pneumoniae* strains were resistant to erythromycin and one *Pseudomonas aeruginosa* strain was resistant to piperacillin and ticarcillin.

Gurunathan *et al.* tested silver nanoparticles in combination with antibiotics on both gram positive bacteria and gram negative bacteria. They noticed a more pronounced increase in the association of nanoparticles with ampicillin against Gram-negative negative bacteria *Pseudomonas aeruginosa* and *Shigella flexneri*; and vancomycin association for Gram-positive bacteria *Staphylococcus aureus* and *Streptococcus pneumoniae* [11]. In another recent study, Kamble and Shinde tested the biofilm inhibition activity of green silver nanoparticles obtained with Curcuma longa and observed promising activity against biofilm formation [12]. The interest in finding new active compounds against resistant bacteria, obtained by safe and cheap methods without the use of harmful chemical reagents, remains a topical one.

**Conclusions**

The antibacterial activity of silver nanoparticles obtained by biosynthesis with an aqueous extract of rosemary leaves has been tested against strains of *P. aeruginosa*, *H. influenzae* and *S. pneumoniae* known to be resistant to several antibiotics. An increased antibacterial activity of RAE AgNPs with higher efficacy against *P. aeruginosa* has been observed. Also, RAE AgNPs have been shown to be effective inhibitors of biofilm formation even at low concentrations.

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