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Control of mucoid *Pseudomonas aeruginosa* biofilms in a highly porous skin model (*ex vivo/in vitro* model) with carboxylate dendrimer-endolysin complexes

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The alarming rise in antibiotic resistance in *Pseudomonas aeruginosa*, resulting from its ability to undergo enzymatic modifications, mutate, or form complex structures such as biofilms, has rendered many conventional therapies ineffective, creating an urgent need to search for new antimicrobial agents. Ligand-modified nanoparticles are a promising alternative to conventional agents. For example, silver-containing dendrimers (metallo-dendrimers), which permeabilize the bacterial outer membrane, enhance the production of reactive oxygen species, and, in combination with peptidoglycan-degrading proteins—CHAP-domain endolysins—can cross the barrier of Gram-negative bacteria. However, their activity against mature biofilms and potential cytotoxicity to mammalian tissues remain poorly understood. In this study, we used an *ex vivo* pig ear skin model to evaluate the ability of three carboxylate dendrimers and their combinations with endolysin to inhibit biofilm formation with *P. aeruginosa* PAO1. The irregular skin texture and richer nutrient resources in the *ex vivo* model may limit the effectiveness of nanomaterials compared to the biophilic *P. aeruginosa* in the *in vitro* model on a smooth plastic surface. Early application of dendrimers with endolysin significantly reduces bacterial survival, while late application causes excessive damage to the epidermis, making it difficult for dendrimers to reach the bacteria and hindering the assessment of the effectiveness of antibacterial drugs. These results clearly suggest that studies on the antibacterial activity of various nanomaterials on smooth surfaces differ significantly from the highly porous structure of the skin model, which has an important impact on their limited application in the treatment of skin wounds.

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1. Introduction

In 2028, we will celebrate the 100th anniversary of the discovery of the first antibiotic, penicillin, by Alexander Fleming, which changed the outlook for fighting infections. However, it was not expected that within 100 years, bacteria would acquire such resistance to these types of compounds that it would be almost

impossible to fight them. Current medicines face a serious challenge due to the resistance of bacteria, which causes the deaths of millions each year.^{1,2} The problem is amplified by a decrease in the production of new and effective antibiotics, which limits the ability to successfully treat infections. Current therapies usually rely on modifications of existing drugs, but these strategies provide only a temporary solution.^{3,4} With conventional antibiotics losing their effectiveness, it has become necessary to search for alternative treatments that are capable of combating MDR (multi-drug resistant) pathogens.

One of the most common and “critical” pathogens responsible for hospital-acquired infections is *Pseudomonas aeruginosa*.^{5,6} It is a rod-shaped, opportunistic, enveloped and highly adaptive Gram-negative bacterium that is widely distributed in aquatic and soil environments and is capable of growth under both aerobic and oxygen-limited conditions (facultatively anaerobic). Its remarkable metabolic flexibility allows it to survive in a variety of conditions.^{7,8} *P. aeruginosa* is particularly dangerous

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for immunosuppressed patients, in whom it can lead to chronic and nosocomial infections, hard-to-treat wounds, and urinary tract infections (UTI); it is also associated with pneumonia and bloodstream and surgical site infections.^{9,10}

Mechanisms of bacterial resistance include limited cell membrane permeability, active antibiotic removal systems (efflux pumps), various mutations that reduce drug sensitivity, and horizontal gene transfer that allows rapid transmission of resistance traits.^{10–13} One of the most effective ways in which *Pseudomonas aeruginosa* avoids antibiotics is its ability to form a biofilm.

A bacterial biofilm is a highly organized three-dimensional community of microorganisms embedded in the extracellular polymeric substances (EPS), consisting mainly of exopolysaccharides, such as alginate, proteins and nucleic acids.^{14,15} Bacteria adhere tightly to each other and/or to non-biological solid surfaces, such as catheters or endoprostheses, biological surfaces, such as wounds, burn injuries, ulcers, or cystic fibrosis.^{11,16} Bacteria possess many features that help them attach to a surface, such as hydrophobicity of the cell wall, surface potential, size and shape, the presence of quorum sensing, pili and curli, and the ability to produce extracellular adhesins. However, it is important to note that bacterial adhesion to a surface has different levels, depending on several surface factors, such as hydrophobicity, heterogeneity, patterns, roughness, stiffness, surface potential and charge. In addition, environmental conditions are crucial for bacterial attachment, e.g., pH, temperature and ionic strength.^{17–19} Free-living/planktonic bacteria, once in contact with the surface of a medical device or injured tissue, due to the action of van der Waals forces, the presence of adhesins, type IV pili, and lipopolysaccharide (LPS), start producing extracellular polymers that organize into a spatial matrix, enabling firm adhesion and biofilm development.^{20,21} The process of biofilm formation consists of several steps: (1) the adhesion of individual bacterial cells to the surface, which can take place through various surface structures, such as fimbriae, pili, or adhesion substances; (2) bacteria enter the microcolony phase, and their cell divisions lead to an increase in the number of cells within the structure; (3) biofilm maturation, during which the bacteria produce an extracellular matrix (EPS) that provides stability and protection against external agents.^{22,23}

In the mature biofilm, intensive cell–cell interactions take place, and the bacteria diversify in terms of metabolic functions, which promotes their survival under changing conditions.²⁴ In time, a dispersal phase occurs, in which some cells detach from the biofilm and can colonize new environments. The structure of the biofilm makes bacteria embedded in it up to hundreds or thousands of times more resistant to antibiotics than their planktonic analogs.^{25,26} This is due to several factors, including the limited penetration of drugs deep into the biofilm, the presence of spore forms, the ability of bacteria to enter a metabolic dormant state, the secretion of protective substances, and many others, as mentioned above, which reduce their vulnerability to drugs. The biofilm also provides protection from the immune system's response as a result of dynamic changes in gene expression, resulting in the

differentiation of surface antigens and leading to the chronic nature of the infection.²¹ Because of these properties, biofilms are a major cause of difficult-to-treat infections. The increase in antibiotic resistance and the ability of bacteria to form biofilms make classic treatments less and less effective.^{27,28} Consequently, the search is on for new therapeutic strategies that could inhibit biofilm formation or effectively eliminate it. Research into new methods of biofilm control is crucial for the future of treating bacterial infections and reducing the spread of multidrug-resistant strains.

Nanomaterials, such as dendrimers, also known as ligand-modified nanoparticles, are widely known for their antimicrobial properties and are a promising alternative to the currently studied non-functioning antibiotics.²⁹ The improvement of their structure and therapeutic potential is the subject of ongoing research. It is important to note that dendrimers can differ significantly from one another, and these differences in structure and surface modifications result in different mechanisms of antimicrobial action. Additional modification with metal ions, such as silver, zinc, copper or gold, as well as functional groups, including imidazolium or pyrimidine derivatives, has been shown to significantly increase their antimicrobial and biological efficacy.^{30,31}

There is increasing number of evidence that demonstrates the potential of combining dendrimers with bacteriophages or antimicrobial proteins (AMPs), such as lysins, to enhance antibacterial activity.^{32,33} Endolysins, for instance, are capable of specifically targeting and degrading bacterial peptidoglycan by cleaving bonds within peptide cross-links, such as tetrapeptides and penta-glycine bridges (endolysin CHAP domain). In Gram-positive bacteria, endolysins can directly access and degrade the exposed peptidoglycan.^{34–36} In contrast, the presence of an outer membrane (OM) in Gram-negative bacteria provides a significant barrier, which requires the use of membrane-permeabilizing agents, such as dendrimers, to enable the effective action of endolysins.³⁷ Based on numerous scientific reports, the use of dendrimers containing silver ions, particularly in combination with antimicrobial proteins, such as endolysin, represents a promising strategy for the effective elimination of planktonic bacterial cells. These compounds exhibit antibacterial effects through multiple mechanisms, including permeabilization of the outer membrane, degradation of the peptidoglycan layer, and increased reactive oxygen species (ROS) production.^{38–40}

Dendrimers are widely used as drug and gene carriers in nanomedicine and targeted pharmacology.⁴¹ Appropriately designed and modified, they can efficiently transport active substances without adversely affecting target cells or tissues.⁴² Nevertheless, in addition to their antimicrobial properties, dendrimers can also exhibit harmful effects on eukaryotic cells, manifested as cytotoxicity or genotoxicity. According to scientific reports, the intensity of these effects can be strongly dependent on the structure of the dendrimer, the type of surface ligands and the concentration used.⁴³ Since it is known that dendrimers containing silver ions can enhance the production of reactive oxygen species (ROS) and cationic dendrimers permeabilize the

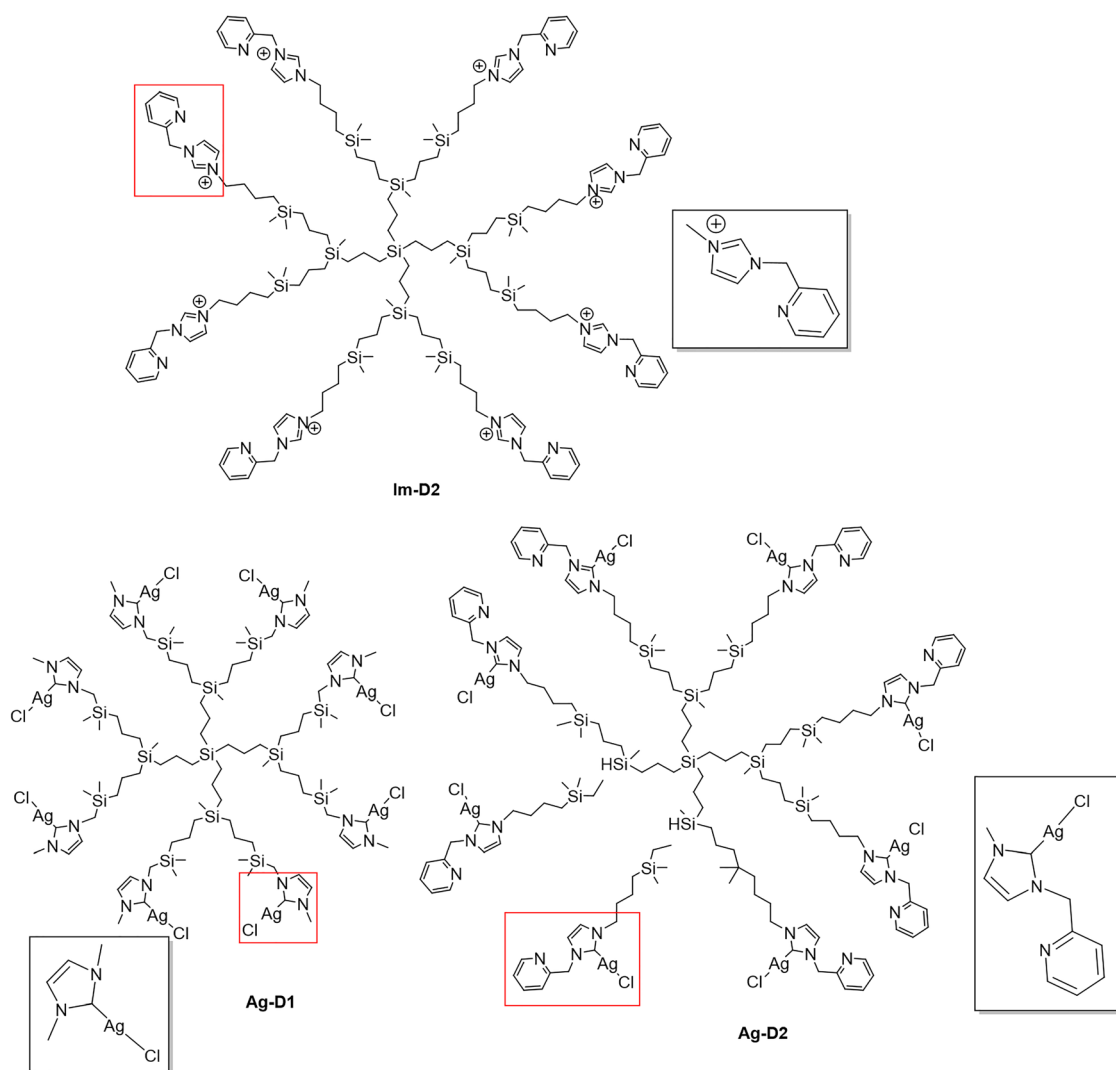


outer membrane in bacteria, there is a high probability that a similar effect will also occur in eukaryotic cells. This can lead to increased cytotoxicity, resulting in membrane destabilization and oxidative damage to lipids, proteins, and genetic material (DNA and RNA). As a consequence, dysfunction of cellular organelles, including mitochondria, can occur, and long-term accumulation of free radicals can trigger pathways leading to cell death (caspase-3 enzyme or p53 gene after genotoxic stress initiates apoptosis).⁴⁴ The use of antimicrobial proteins such as endolysins is considered safe in terms of cytotoxicity because eukaryotic cells do not contain peptidoglycan, which is the main substrate of these enzymes. Consequently, endolysins have no activity against mammalian cells, making them promising candidates for therapeutic applications.

While dendrimers and antimicrobial proteins hold significant therapeutic potential, especially as targeted nanocarriers, their broader biomedical application requires a thorough understanding of their toxicity and biological interactions.

Although the initial results are promising, there is a persistent lack of data regarding their effects on complex bacterial structures, such as biofilms, and eukaryotic cells or tissues. Therefore, further *in vitro* and *ex vivo/in vivo* studies are essential to optimize therapeutic efficacy while minimizing adverse effects.

This study aimed to evaluate the inhibition of biofilm formation by planktonic *Pseudomonas aeruginosa* cells using both an *in vitro* laboratory model and an *ex vivo* approach utilizing pork ear skin. It is crucial to understand not only how bacterial biofilms respond to treatment under controlled laboratory conditions, but also how they might behave during therapeutic interventions targeting human or animal infections. Additionally, the influence of a complex of dendrimers and endolysin^{38,39} on normal, uninfected skin was examined to allow a comparative analysis between healthy and *P. aeruginosa*-infected tissue. Prior to testing the dendrimers and endolysin on *ex vivo* skin models, it was necessary to assess the cytotoxic effects of the compounds by evaluating the viability of VH10



Scheme 1 Structures of CBS dendrimers **Im-D2** (chloride anions omitted for clarity), **Ag-D1** and **Ag-D2**. The functional groups are highlighted in red squares.



fibroblast cells exposed to various concentrations of each tested agent.

2. Materials and methods

2.1. Carbosilane dendrimers

The silver and imidazolium metal carbosilane dendrimers **Ag-D1**,⁴⁵ **Ag-D2** and **Im-D2**³⁸ (Scheme 1) were prepared according to the procedures described previously.

2.2. Recombinant endolysin

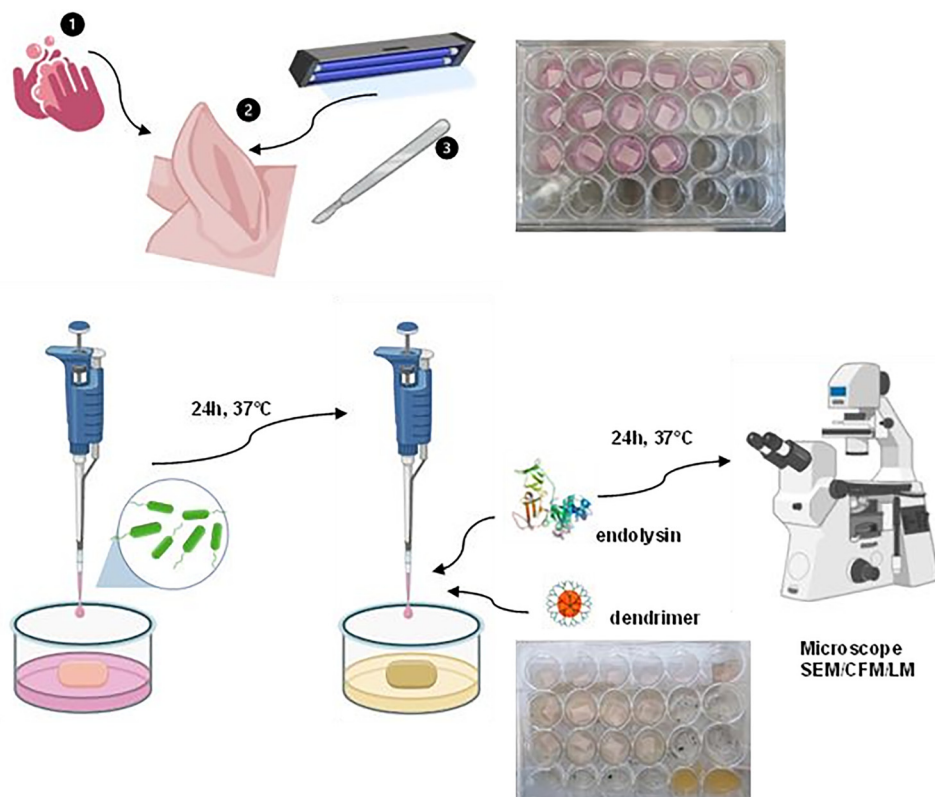
The recombinant CHAP endolysin fragment of the phage Φ 812 catalytic domain was purchased from Protean, Czech Republic.

2.3. Bacterial biofilm formation assay

2.3.1. Crystal violet assay. The inhibition of *P. aeruginosa* PAO1 biofilms was assessed using a crystal violet (CV) assay. This method quantifies the total biofilm mass based on the binding ability of the CV dye to extracellular polymeric substances (EPS) and both live and dead bacterial cells. Overnight-cultured bacteria were diluted in 10 mL sterile TSB and spotted onto a 96-well plate, supplemented with fresh TSB, and allowed to form a biofilm under the conditions of 37 °C & 5% CO₂ for 24 hours. The mature biofilms were gently washed twice with phosphate-buffered saline (PBS, pH 7.4) to remove unbound bacteria. PBS was removed, and the biofilms were left to dry. Next, the biofilms were stained for 15 min with 0.1% crystal

violet solution and then washed three times with PBS to remove the excess dye. Next, CV was dissolved in 99.9% ethanol. After 15 min of incubation, the solution was transferred to a new 96-well plate, and the absorbance at 570 nm was measured using a TECAN SPARK Magellan V2.2 STD spectrometer (Tecan Group Ltd., Switzerland). The absorbance of the de-staining solution represents the biofilm mass. Experiments were performed in triplicate.

2.3.2. FilmTracer™ LIVE/DEAD™ biofilm viability kit – *in vitro*. Labeling of live and dead bacterial cells in the formed biofilm was performed using the dedicated FilmTracer™ LIVE/DEAD™ Biofilm Viability Kit (Invitrogen, USA). The dye solution was prepared according to the manufacturer's instructions, *i.e.*, 3 μ L of SYTO-9 dye (480/500 nm) and 3 μ L of propidium iodide (490/635 nm) were added to 1 mL of distilled water (MilliQ, MERCK). Overnight-cultured bacteria were diluted in 10 mL sterile TSB and spotted onto a 24-well plate (Wuxi NEST Biotechnology Co., Ltd), supplemented with fresh TSB, and allowed to form a biofilm under the conditions of 37 °C & 5% CO₂ for 24 hours. Appropriate volumes of dendrimers and endolysin were added to the formed biofilm so that the concentrations were 8 and 32 μ g mL⁻¹ and 2 μ g mL⁻¹, respectively, and incubated in a 37 °C & 5% CO₂ for 24 hours. After this time, the old medium was gently pulled off and washed twice with *d*H₂O. The biofilm was stained with the prepared solution for 20 minutes in the dark, the dye was rinsed off, and, using a metal scratcher, was transferred to a primary slide,



Scheme 2 Method of preparing the skin for the experiments described in the text.



covered with a coverslip and observed under a confocal microscope (NIKON Eclipse Ti/A1, Japan), no longer than 30 min after staining.

2.3.3. Preparation of pork external ear skin and the *Pseudomonas aeruginosa* PAO1 bacterial biofilm testing model. The fresh pork ear came from a local slaughterhouse. The skin of the pork's external ear was cleaned by washing thoroughly with hot water and gray soap, and the hair was removed with a razor blade. The skin was then pre-sterilized by washing several times with 70% ethanol and irradiating with UV light for 20 minutes. After the sterilization process, under a laminar chamber, the skin was separated from the cartilage using a scalpel. The specimens were then divided into smaller pieces of about 1 × 1 cm and 1–3 mm thick. All further experimental procedures were carried out on the day of sterilization; the samples were not stored or frozen.

To analyze the formation of the *P. aeruginosa* PAO1 bacterial biofilm (Scheme 2), the culture medium was prepared using DMEM (Dulbecco's Modified Eagle Medium) containing L-glutamine and phenol red and further enriched with 10% fetal bovine serum (FBS). For the experiment, 24-well culture plates were used, in which previously prepared and UV-sterilized filter paper discs, matched in size to the diameter of the wells, were placed. Here, 250 μL of DMEM medium was added to each well, followed by the placement of sterile skin sections (1 × 1 cm). The bacteria were prepared as described above. Next, 10 μL of diluted bacteria were spotted in the center of a piece of skin and allowed to form a biofilm under the conditions of 37 °C & 5% CO₂ for 24 hours. Appropriate volumes of dendrimers (8 and 32 μg mL⁻¹) and endolysin (2 μg mL⁻¹) and fresh DMEM medium (appropriate volume, in total 400 μL) were added to the formed biofilm on skin and incubated at 37 °C with 5% CO₂ for 24 hours. After this time, the old medium was gently removed, followed by washing twice with dH₂O. The biofilm formed on the skin was stained by spotting 30 μL of dye and left for 20 min in the dark. The dye was then rinsed three times with distilled water and the skin was gently transferred with tweezers to a coverslip. The prepared samples were observed under a confocal microscope, no longer than 30 min after staining.

2.3.4. Scanning electron microscopy. Skin preparation for scanning microscope observation was the same as described above, with a smaller skin area (about 5 mm × 5 mm). After culture and incubation, skin samples were fixed in a 3% glutaraldehyde solution in 0.2 M cacodyl buffer for 3 hours, with the fixative solution replaced with fresh solution after 30 minutes. The samples were then rinsed twice in 0.1 M cacodyl buffer. In a further step, osmication was carried out in 2.5% osmium tetroxide (OsO₄) in 0.1 M cacodyl buffer for 2 hours to further fix and contrast cell structures. After that, the samples were again washed twice in a cacodyl buffer. Tissue dehydration was then carried out in increasing concentrations of ethanol: 40%, 50%, 60%, 70%, 80%, 90% and 100%, with each incubation lasting 20 minutes to effectively remove water from the tissues. After complete dehydration, the samples were subjected to screening in mixtures of ethanol and propylene

oxide in proportions as follows: 3:1, 1:1, and 1:3, and then they were suspended in pure propylene oxide. The samples were placed on a glass slide and left to dry completely, then placed on carbon discs, covered with gold using a JEOL JFC 110E Fine Coat Ion Sputter, and analyzed by SEM (JEOL JSM-7100F, JEOL Ltd, Japan) at an accelerating voltage of 15 kV.

2.3.5. Histomorphology of uninfected and *Pseudomonas aeruginosa*-infected pork skin after treatment with dendrimers and their complexes with endolysin. Preparation for the histomorphological observation of non-infected and infected pork skin after treatment with tested concentrations of dendrimers and endolysin was carried out as described above for confocal microscopy, along with fixation with 10% formalin for 24 h. The next day, the tissues were washed twice in PBS, embedded in OCT and frozen at -20 °C. The cryosections were then cut

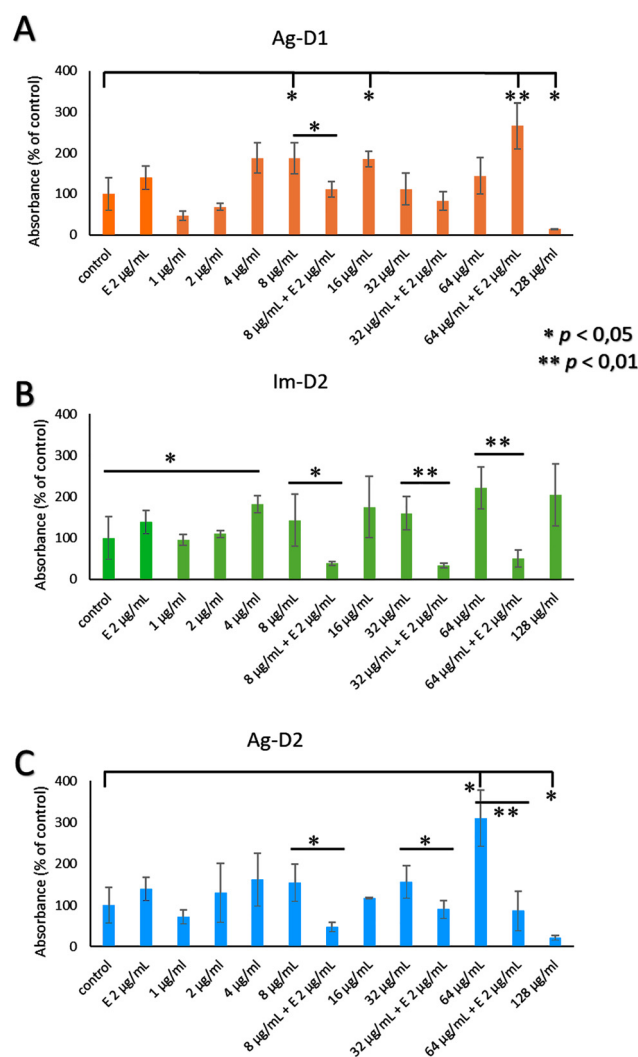


Fig. 1 Inhibition of bacterial biofilm formation quantified by measuring crystal violet absorbance in the presence of increasing concentrations of the dendrimers (A) **Ag-D1**, (B) **Im-D2**, (C) **Ag-D2** and selected concentrations of the metallodendrimer-endolysin (endolysin - E 2 μg mL⁻¹) complexes after 24 h incubation. Experiments were performed in triplicate, and the results are presented as percentages of the control.



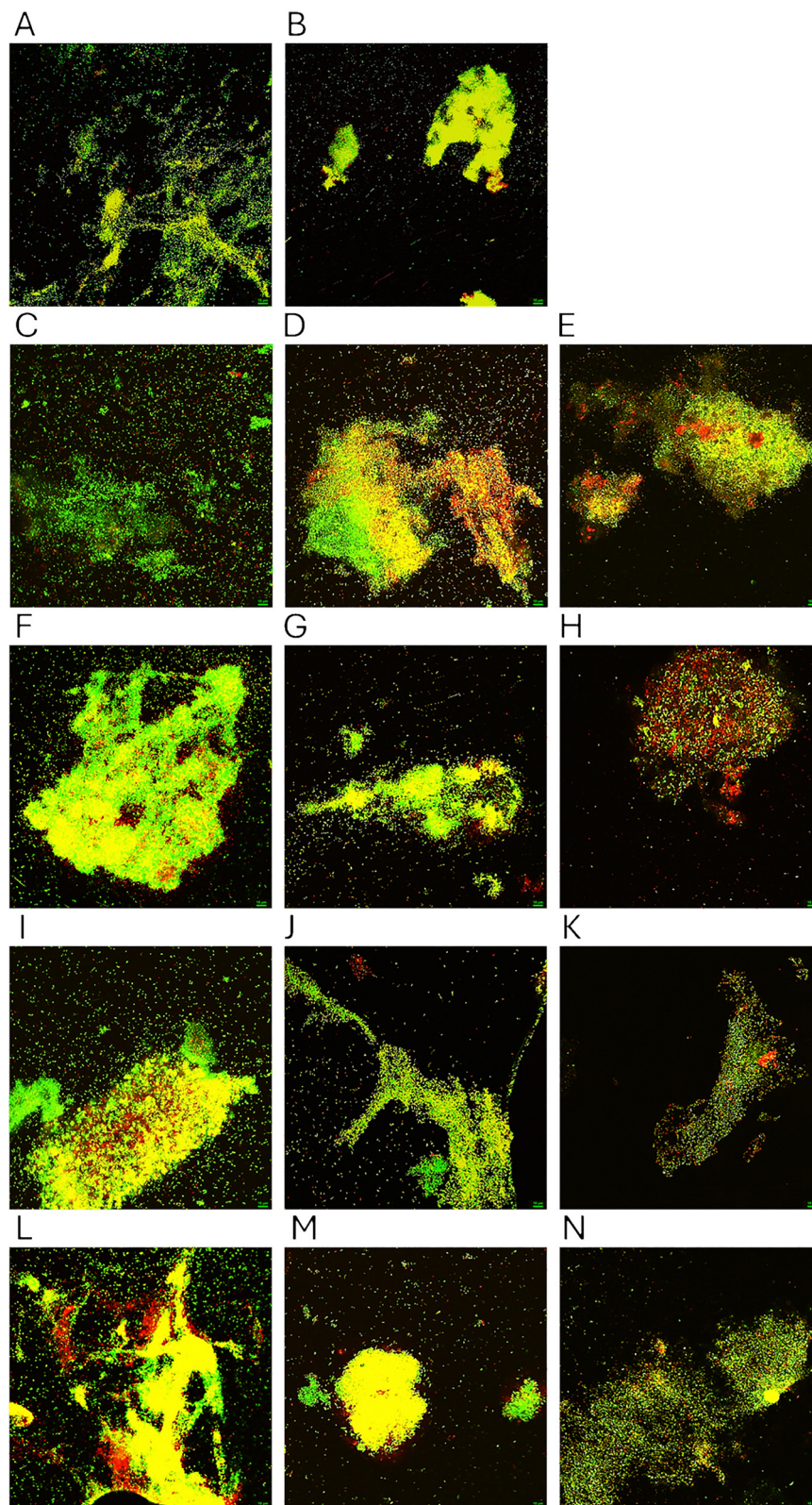


Fig. 2 FilmTracer™ LIVE/DEAD™ Biofilm Viability staining results of the biofilm formed *in vitro* by *Pseudomonas aeruginosa* PAO1. (A) Untreated control group. Twenty four hours after treatment (B) with endolysin, (C) with $8 \mu\text{g mL}^{-1}$ Ag-D1, (D) with $8 \mu\text{g mL}^{-1}$ Im-D2, (E) with $8 \mu\text{g mL}^{-1}$ Ag-D2, (F) with $8 \mu\text{g mL}^{-1}$ Ag-D1 + $2 \mu\text{g mL}^{-1}$ endolysin, (G) with $8 \mu\text{g mL}^{-1}$ Im-D2 + $2 \mu\text{g mL}^{-1}$ endolysin, (H) with $8 \mu\text{g mL}^{-1}$ Ag-D2 + $2 \mu\text{g mL}^{-1}$ endolysin, (I) with $32 \mu\text{g mL}^{-1}$ Ag-D1, (J) with $32 \mu\text{g mL}^{-1}$ Im-D2, (K) with $32 \mu\text{g mL}^{-1}$ Ag-D2, (L) with $32 \mu\text{g mL}^{-1}$ Ag-D1 + $2 \mu\text{g mL}^{-1}$ endolysin, (M) with $32 \mu\text{g mL}^{-1}$ Im-D2 + $2 \mu\text{g mL}^{-1}$ endolysin and (N) with $32 \mu\text{g mL}^{-1}$ Ag-D2 + $2 \mu\text{g mL}^{-1}$ endolysin. Green fluorescence was interpreted as live bacterial cells, whereas bacteria with disrupted membranes show red fluorescence. The scale bar is $20 \mu\text{m}$.



using a Leica CM1100 cryotome, transferred to SuperFrost™ glass slides, stained with 1% toluidine blue for 5 seconds, mounted with a glycerin/PBS mixture under a coverslip, and observed under a Nikon Eclipse 50i light microscope (Japan) with a DS-Fi1 digital camera.

2.4. Cytotoxic effect on fibroblasts

The VH10 human fibroblast cell line (ATCC, Manassas, VA, USA; catalogue no. PCS-201-012) was maintained and cultured in Fibroblast Growth medium (Cat no. 116-500, Cell Application, Inc.) at 37 °C in a humidified atmosphere with 5% CO₂. The culture medium was changed every 2 days. The viability of fibroblast cells treated with dendrimers (concentrations of 1, 2, 4, 8, 16, 32, 64, 128 μg mL⁻¹) and their complexes with endolysin (at concentrations of 1 and 2 μg mL⁻¹) was determined using an MTS assay. The fibroblasts were seeded in 96-well plates. After reaching the appropriate confluence, the cells were treated with dendrimers, endolysin, and dendrimer + endolysin for 24 h. The MTS Cell Proliferation Assay Kit (Colorimetric) (Abcam) was used according to the manufacturer's recommendations, and absorbance at 490 nm was measured using a TECAN SPARK Magellan V2.2 STD spectrometer (Tecan Group Ltd, Switzerland).

3. Results and discussion

3.1. Bacterial biofilm formation in the presence of dendrimers and complexes with endolysin – *in vitro* studies

Fig. 1 shows the effect of the tested dendrimers on mature *P. aeruginosa* PAO1 bacterial biofilm after 24 hours of incubation at various concentrations. In each of the three systems analyzed, a relationship was observed between the concentration of the dendrimer and the degree of inhibition of bacterial growth. For the metallodendrimer **Ag-D1** alone (Fig. 1A), none of the concentrations used caused a reduction in biofilm, and an increase in its amount was even observed. Only the highest concentration, 128 μg mL⁻¹, completely degraded the biofilm and bacteria (~14%). Similarly, the addition of endolysin to selected concentrations, 8, 32, and 64 μg mL⁻¹ did not cause significant degradation of the biofilm. When the imidazolium **Im-D2** dendrimer alone was used (Fig. 1B), as described above, no reduction in biofilm quantity was observed, while an increase in its mass was noted at all concentrations tested, reaching an average of ~161% of the control value. However, the addition of endolysin to the **Im-D2** dendrimer at concentrations of 8, 32, and 64 μg mL⁻¹ resulted in a significant decrease in biofilm quantity, to an average value of ~40% compared to the control. A similar trend was observed for the metallodendrimer **Ag-D2** dendrimer (Fig. 1C). The use of the dendrimer alone did not inhibit biofilm formation; an upward trend was observed for all concentrations tested, reaching an average of ~157%. In contrast, in the presence of endolysin, a significant reduction in biofilm quantity was observed, particularly at a concentration of 8 μg mL⁻¹, where the value dropped to approximately 48%, and when using the highest concentration

of the dendrimer alone (128 μg mL⁻¹), which resulted in a further reduction in biofilm quantity to approximately 21% of the control value.

To confirm the reduced viability of bacteria in formed biofilm due to CBS dendrimers and endolysin, a FilmTracer™ LIVE/DEAD™ Biofilm Viability was performed. Fig. 2A represents the untreated bacterial biofilm used as a control; among a large number of living normal bacteria and stable biofilm, a few red spots appeared. A similar effect was observed for biofilm in the presence of endolysin (Fig. 2B). Higher numbers of red spots were observed near the biofilm exhibiting green fluorescence, but with reduced density after 8 μg mL⁻¹ **Ag-D1** treatment, compared to the control (Fig. 2C). Applying 8 μg mL⁻¹ **Im-D2** (Fig. 2D) caused an increase in the number of dead cells and a decrease in living cells was observed among the dense biofilm; orange and red signals dominated, with a small amount of green. A similar effect was observed for 8 μg mL⁻¹ **Ag-D2** (Fig. 2E), but with a lower intensity of dead cells. Treatment of bacterial biofilm with the additional presence of endolysin did not significantly reduce the biofilm structure (Fig. 2F – 8 μg mL⁻¹ **Ag-D1** + 2 μg mL⁻¹ endolysin), but several red signals were observed. Fig. 2G presents bacterial biofilm after 8 μg mL⁻¹ **Im-D2** + 2 μg mL⁻¹ endolysin, where a reduction in biomass was observed, while small amounts of red spots were present. In the case of 8 μg mL⁻¹ **Ag-D2** + 2 μg mL⁻¹ endolysin (Fig. 2H), the red signal dominated the green one, and the biofilm biomass was reduced.

The use of a higher dose of dendrimers (32 μg mL⁻¹) **Ag-D1** (Fig. 2I), **Im-D2** (Fig. 2J) and **Ag-D2** (Fig. 2K) did not significantly increase the number of dead bacterial cells or decrease the biofilm biomass compared to the lower dose used. Moreover, the additional presence of endolysin with higher doses of dendrimers resulted in an increase in the number of red spots and a reduction in biomass after **Ag-D1** + 2 μg mL⁻¹ endolysin (Fig. 2L), while after **Im-D2** + 2 μg mL⁻¹ endolysin (Fig. 2M), the biofilm was more condensed, and in the case of **Ag-D2** + 2 μg mL⁻¹ endolysin (Fig. 2N), it was diluted and thin.

To explain the mechanism of biofilm inhibition by the tested compounds, both the activity of silver ions and the

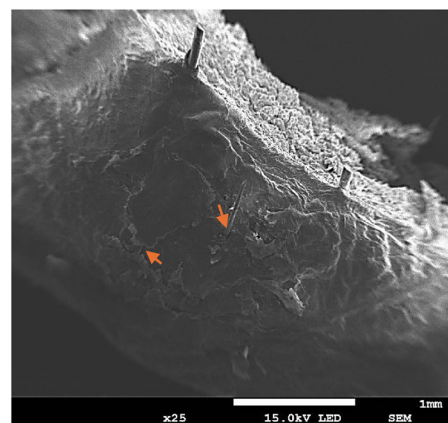


Fig. 3 SEM image of the skin surface.



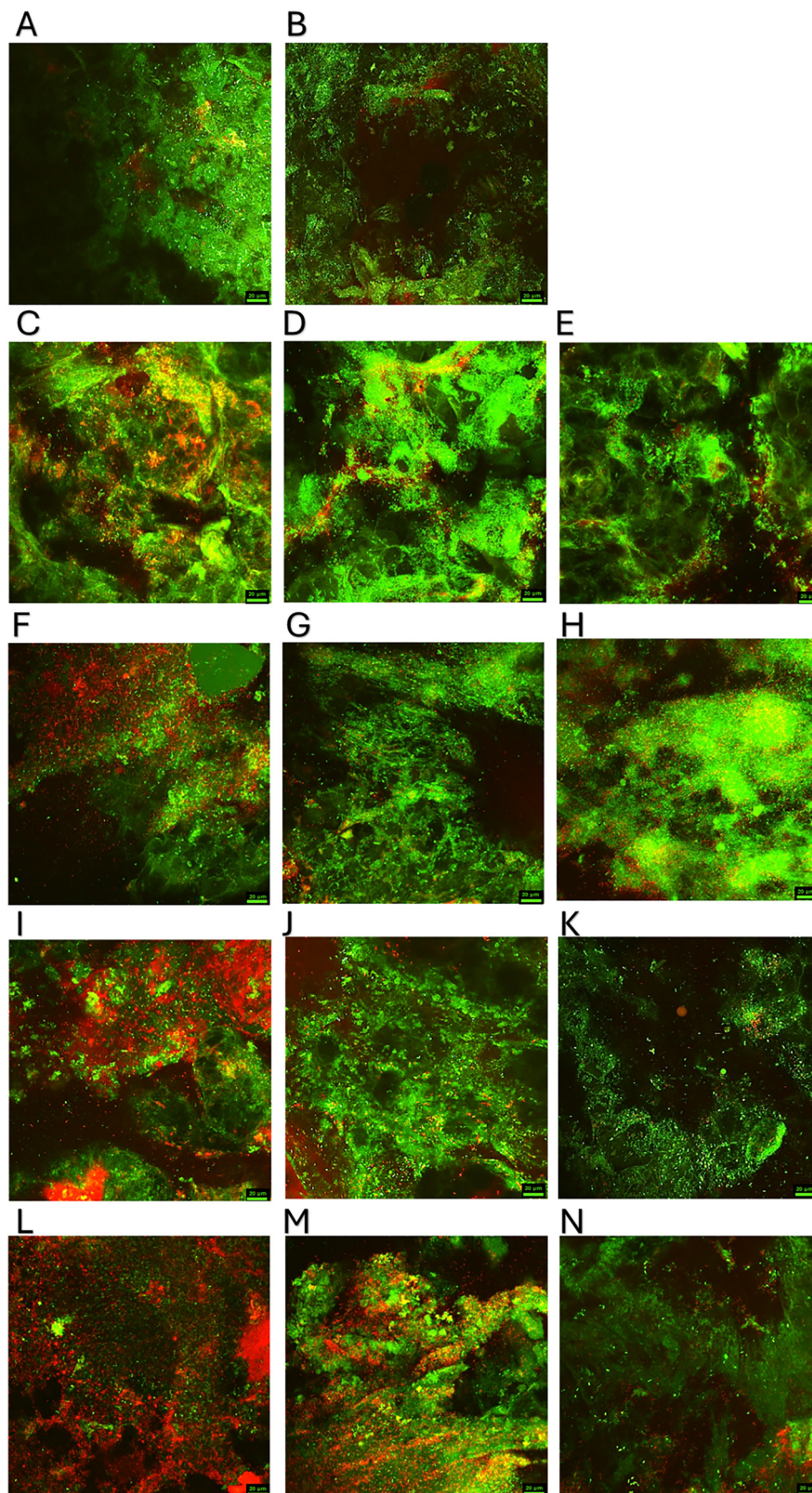


Fig. 4 FilmTracer™ LIVE/DEAD™ Biofilm Viability staining results of biofilm formed *ex vivo* by *Pseudomonas aeruginosa* PAO1 on pork skin. (A) Untreated control group. Twenty-four hours after treatment (B) with endolysin, (C) with $8 \mu\text{g mL}^{-1}$ Ag-D1, (D) with $8 \mu\text{g mL}^{-1}$ Im-D2, (E) with $8 \mu\text{g mL}^{-1}$ Ag-D2, (F) with $8 \mu\text{g mL}^{-1}$ Ag-D1 + $2 \mu\text{g mL}^{-1}$ endolysin, (G) with $8 \mu\text{g mL}^{-1}$ Im-D2 + $2 \mu\text{g mL}^{-1}$ endolysin, (H) with $8 \mu\text{g mL}^{-1}$ Ag-D2 + $2 \mu\text{g mL}^{-1}$ endolysin, (I) with $32 \mu\text{g mL}^{-1}$ Ag-D1, (J) with $32 \mu\text{g mL}^{-1}$ Im-D2, (K) with $32 \mu\text{g mL}^{-1}$ Ag-D2, (L) with $32 \mu\text{g mL}^{-1}$ Ag-D1 + $2 \mu\text{g mL}^{-1}$ endolysin, (M) with $32 \mu\text{g mL}^{-1}$ Im-D2 + $2 \mu\text{g mL}^{-1}$ endolysin and (N) with $32 \mu\text{g mL}^{-1}$ AgD2 + $2 \mu\text{g mL}^{-1}$ endolysin. Green fluorescence was interpreted as live bacterial cells, whereas bacteria with disrupted membranes show red fluorescence. The scale bar is $20 \mu\text{m}$.



specific properties of dendrimers must be taken into account. Ag^+ ions, present in dendrimers **Ag-D1** and **Ag-D2**, are characterized by high affinity for thiol groups, especially cysteine, which is present in numerous enzymes responsible for the metabolism of exopolysaccharides. The breaking of S–H bonds

leads to a weakening of bacterial cell adhesion to the substrate and disruption of the quorum-sensing system. Besides, dendrimers with Ag^+ ions increase the permeability of the bacterial OM and cause an increase in ROS production, leading to planktonic cell disintegration and death,^{39,46} which can occur

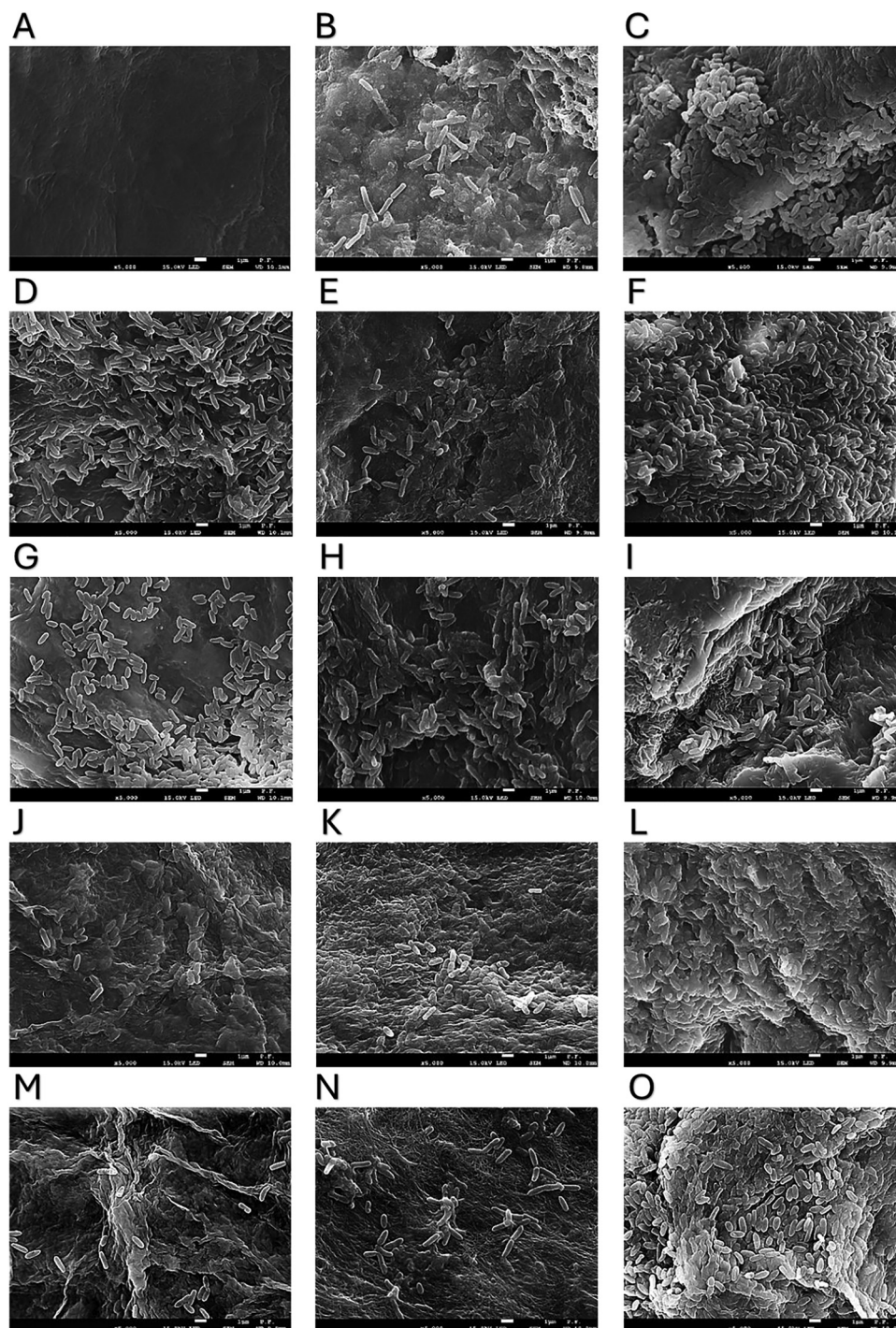


Fig. 5 SEM images present the morphology of the bacterial biofilm formed by *Pseudomonas aeruginosa* PAO1 on pork skin. (A) Uninfected skin. (B) Untreated bacterial biofilm. Twenty-four hours after treatment (C) with endolysin, (D) with $8 \mu\text{g mL}^{-1}$ **Ag-D1**, (E) with $8 \mu\text{g mL}^{-1}$ **Im-D2**, (F) with $8 \mu\text{g mL}^{-1}$ **Ag-D2**, (G) with $8 \mu\text{g mL}^{-1}$ **Ag-D1** + $2 \mu\text{g mL}^{-1}$ endolysin, (H) with $8 \mu\text{g mL}^{-1}$ **Im-D2** + $2 \mu\text{g mL}^{-1}$ endolysin, (I) with $8 \mu\text{g mL}^{-1}$ **Ag-D2** + $2 \mu\text{g mL}^{-1}$ endolysin, (J) with $32 \mu\text{g mL}^{-1}$ **Ag-D1**, (K) with $32 \mu\text{g mL}^{-1}$ **Im-D2**, (L) with $32 \mu\text{g mL}^{-1}$ **Ag-D2**, (M) with $32 \mu\text{g mL}^{-1}$ **Ag-D1** + $2 \mu\text{g mL}^{-1}$ endolysin, (N) with $32 \mu\text{g mL}^{-1}$ **Im-D2** + $2 \mu\text{g mL}^{-1}$ endolysin, and (O) with $32 \mu\text{g mL}^{-1}$ **Ag-D2** + $2 \mu\text{g mL}^{-1}$ endolysin. The scale bar is $1 \mu\text{m}$.



in biofilm. Additionally, a small dose of the tested compound may increase EPS and protein matrix production, which would change the biofilm structure, and the metabolic activity of bacterial cells would decrease due to the disruption of intracellular processes.⁴⁷ At the same time, the electrical charge of the EPS matrix, which is mainly negative, must be taken into account. Positively charged dendrimers can be strongly attracted to the biofilm by electrostatic forces, which may facilitate the delivery of silver ions and increase matrix penetration. Moreover, it may promote troublesome aggregation of dendrimers or bacterial cells, leading to increased biofilm formation or changing physical properties, such as density and adhesion capacity to the surface.⁴⁸

3.2. Bacterial biofilm formation on skin tissue in the presence of dendrimers and complexes with endolysin – *ex vivo* studies

The skin tissue consists of three main layers: the epidermis with the stratum corneum (the outermost part of the epidermis), the dermis, and the subcutaneous tissue. It acts as a basic protective barrier against pathogens, thanks to several key mechanisms: acidic pH (around 5.5), production of antimicrobial proteins (lysozyme, cathelicidin, β -defensins, lactoferrin, and others), and the presence of salts. The stratum corneum, composed of keratinized cells, prevents the permanent adhesion of microorganisms and promotes their removal through continuous exfoliation. Although it contains few nutrients, it is rich in keratin, which some bacteria can use as a food source.⁴⁹

The surface of the skin, unlike the smooth surface of a 24- or 96-well laboratory plate, is characterized by numerous depressions, protrusions, and pores that drain sebaceous and sweat gland secretions, as well as hair deposits (Fig. 3). These microscopic structures provide excellent anchorage points for bacteria, which can colonize them and develop biofilms under natural conditions or infection. The resulting complex bacterial matrix is then highly resistant to mechanical and chemical removal methods, which significantly hinders the effective treatment of skin infections. Furthermore, due to the presence of cholesterol and other lipids in the stratum corneum, the surface charge of the skin is mostly negative, which is important for bacterial adhesion.^{50,51}

To assess the effects of the CBS imidazolium dendrimer and metal dendrimers and endolysin on the formed biofilm of *Pseudomonas aeruginosa* PAO1, an *ex vivo* model was used on pork skin pieces, which, after infection and treatment with dendrimers, were subjected to staining to differentiate between live and dead bacterial cells in the biofilm to be analyzed by confocal fluorescence microscopy. The results are shown in Fig. 4. Fig. 4A shows a dense, unchanged biofilm of the *P. aeruginosa* PAO1 (control group). After treatment with endolysin alone ($2 \mu\text{g mL}^{-1}$, Fig. 4B), no significant changes were observed – the biofilm structure was characterized by high bacterial viability, comparable to the control. Treatment of the biofilm with **Ag-D1** at a concentration of $8 \mu\text{g mL}^{-1}$ (Fig. 4C) led to partial thinning of the matrix, with gaps appearing in the lower layers and red signals indicating dead cells. Similar changes, with a predominance of intense green fluorescence of surface bacteria and periodic red foci, were observed after the application of **Im-D2** ($8 \mu\text{g mL}^{-1}$, Fig. 4D) and the **Ag-D2** complex ($8 \mu\text{g mL}^{-1}$, Fig. 4E). The addition of endolysin increased the number of dead cells in the biofilm in combination with **Ag-D1** (Fig. 4F) and **Ag-D2** (Fig. 4H), and a clear increase in red signal was observed, while in combination with **Im-D2** (Fig. 4G), this effect was weaker. Increasing the concentration of dendrimers to $32 \mu\text{g mL}^{-1}$ enhanced the bactericidal effect. **AgD1** (Fig. 4I) and **Im-D2** (Fig. 4J) caused a significant increase in the red signal, while **Ag-D2** (Fig. 4K) slightly increased the proportion of dead cells. The combination of $32 \mu\text{g mL}^{-1}$ **Ag-D1** with endolysin (Fig. 4L) showed the highest efficacy, as the biofilm then contained mostly dead bacteria with a minimal proportion of live ones. Under similar conditions, the **Im-D2** + endolysin complex (Fig. 4M) showed a slightly weaker effect, while **Ag-D2** + endolysin (Fig. 4N) showed a moderate but noticeable improvement in efficacy compared to the dendrimer alone.

Scanning electron microscopy (SEM) was used to evaluate bacterial biofilm morphology after treatment with dendrimers and their endolysin complexes (Fig. 5), and the results are summarized in Table 1. The negative control group (Fig. 5A) shows a sterile, uninfected skin surface. The positive control group (Fig. 5B) shows a properly formed biofilm and intact

Table 1 Bacterial biofilm changes after treatment with dendrimers and their complexes with endolysin

Group	EPS	Bacterial visibility	Appearance of bacteria
A. Control	None	None	—
B. PAO1	Present, dense	High	Undamaged, normal
C. $2 \mu\text{g mL}^{-1}$ E	Present, dense, undulating	Medium	Normal
D. $8 \mu\text{g mL}^{-1}$ Ag-D1	Present, less compact	Very high	Shortened, irregular shape
E. $8 \mu\text{g mL}^{-1}$ Im-D2	Weakened, flat	Medium	Normal
F. $8 \mu\text{g mL}^{-1}$ Ag-D2	Present, dense, undulating	Low (EPS coated)	
G. $8 \mu\text{g mL}^{-1}$ Ag-D1 + $2 \mu\text{g mL}^{-1}$ E	Diluted, cracked, delaminated	Low (EPS coated)	
H. $8 \mu\text{g mL}^{-1}$ Im-D2 + $2 \mu\text{g mL}^{-1}$ E	Present, dense, compact	Low (EPS coated)	Irregular shapes
I. $8 \mu\text{g mL}^{-1}$ Ag-D2 + $2 \mu\text{g mL}^{-1}$ E	Diluted	High	Irregular shapes
J. $32 \mu\text{g mL}^{-1}$ Ag-D1	Dense, stratified	Low (EPS coated)	
K. $32 \mu\text{g mL}^{-1}$ Im-D2	Severely fibrosis	Low (EPS coated)	
L. $32 \mu\text{g mL}^{-1}$ Ag-D2	Diluted	High	Irregular shapes
M. $32 \mu\text{g mL}^{-1}$ Ag-D1 + $2 \mu\text{g mL}^{-1}$ E	Undulating, cracked, dense	Medium	Damaged, irregular shapes
N. $32 \mu\text{g mL}^{-1}$ Im-D2 + $2 \mu\text{g mL}^{-1}$ E	Present, dense	Low (EPS coated)	Numerous cell wall damages
O. $32 \mu\text{g mL}^{-1}$ 2Ag-D2 + $2 \mu\text{g mL}^{-1}$ E	Diluted	High	



bacterial cell structure. The administration of endolysin at a concentration of $2 \mu\text{g mL}^{-1}$ (Fig. 5C) did not significantly reduce EPS continuity; the matrix remained dense and wavy and was associated with moderate visibility of bacteria with an intact appearance. In contrast, treatment of the biofilm with $8 \mu\text{g mL}^{-1}$ **Ag-D1** (Fig. 5D) resulted in a reduction in the compact EPS structure, which became looser, and the bacteria, despite their very high numbers, exhibited shortened, irregular shapes. The use of $8 \mu\text{g mL}^{-1}$ **Im-D2** (Fig. 5E) caused a weakened and flattened EPS matrix, restoring the average level of bacteria and

their normal appearance. A lower dose of **Ag-D2** ($8 \mu\text{g mL}^{-1}$, Fig. 5F) resulted in dense, wavy EPS and poorly visible bacteria. The combination of **Ag-D1** with endolysin ($8 \mu\text{g mL}^{-1}$ **Ag-D1** + $2 \mu\text{g mL}^{-1}$ endolysin, Fig. 5G) resulted in dilution and cracking of the EPS. **Im-D2** at a concentration of $8 \mu\text{g mL}^{-1}$ and $2 \mu\text{g mL}^{-1}$ endolysin (Fig. 5H) caused a compact and dense EPS layer in which the visibility of bacteria was low, and their shapes were irregular. The addition of endolysin to **Ag-D2** (Fig. 5I) diluted the EPS but increased the visibility of the bacteria, which retained their irregular shape. A high dose of **Ag-D1** (Fig. 5J) resulted in a

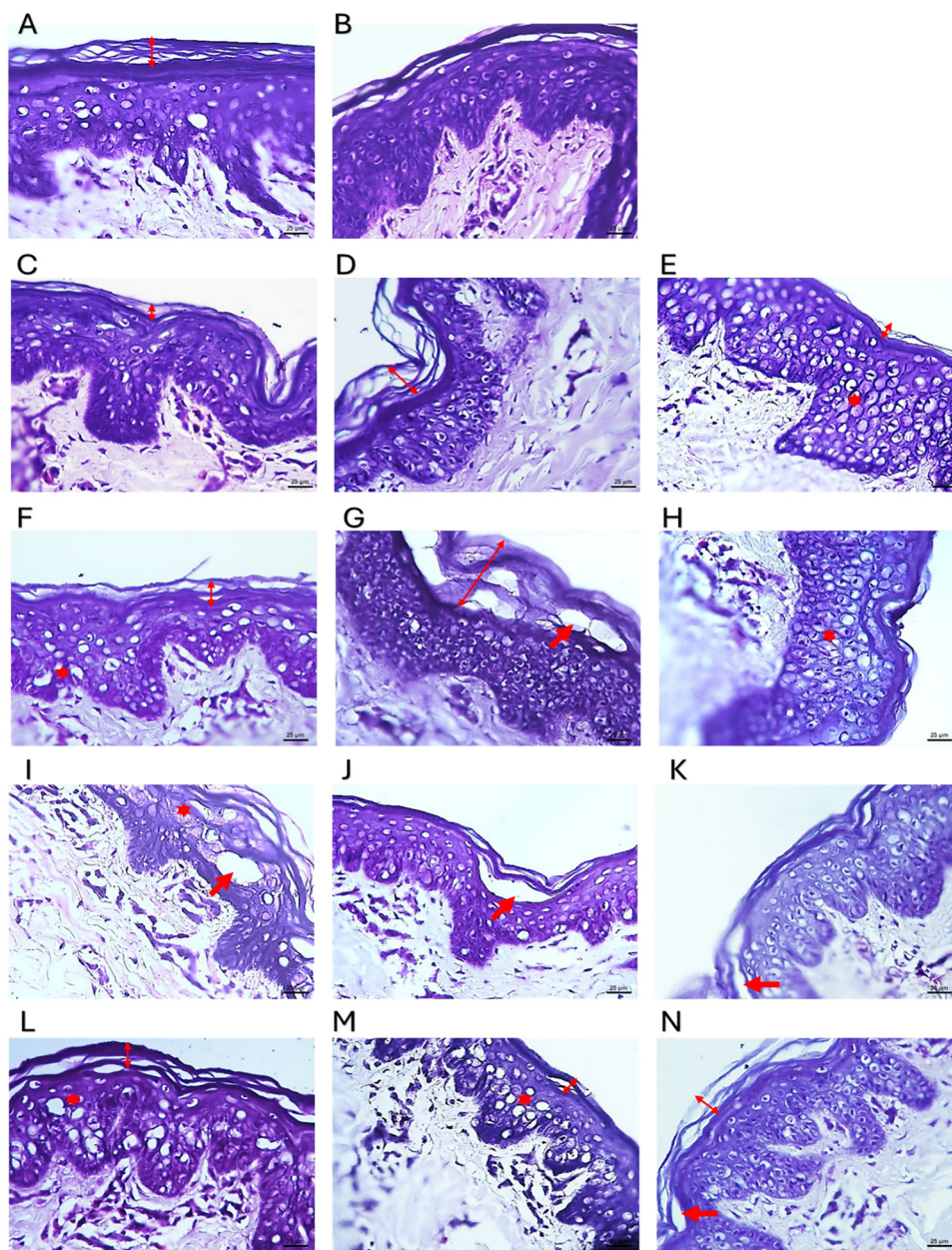


Fig. 6 Histomorphology of cryosections of uninfected pork skin. (A) Untreated control group. Twenty-four hours after treatment (B) with endolysin, (C) with $8 \mu\text{g mL}^{-1}$ **Ag-D1**, (D) with $8 \mu\text{g mL}^{-1}$ **Im-D2**, (E) with $8 \mu\text{g mL}^{-1}$ **Ag-D2**, (F) with $8 \mu\text{g mL}^{-1}$ **Ag-D1** + $2 \mu\text{g mL}^{-1}$ endolysin, (G) with $8 \mu\text{g mL}^{-1}$ **Im-D2** + $2 \mu\text{g mL}^{-1}$ endolysin, (H) with $8 \mu\text{g mL}^{-1}$ **Ag-D2** + $2 \mu\text{g mL}^{-1}$ endolysin, (I) with $32 \mu\text{g mL}^{-1}$ **Ag-D1**, (J) with $32 \mu\text{g mL}^{-1}$ **Im-D2**, (K) with $32 \mu\text{g mL}^{-1}$ **Ag-D2**, (L) with $32 \mu\text{g mL}^{-1}$ **Ag-D1** + $2 \mu\text{g mL}^{-1}$ endolysin, (M) with $32 \mu\text{g mL}^{-1}$ **Im-D2** + $2 \mu\text{g mL}^{-1}$ endolysin and (N) with $32 \mu\text{g mL}^{-1}$ **Ag-D2** + $2 \mu\text{g mL}^{-1}$ endolysin. The scale bar is $25 \mu\text{m}$ (cross-sections, magnification $400\times$, 1% toluidine blue staining).



dense, layered EPS and, at the same time, low visibility of irregularly shaped cells. A concentration of $32 \mu\text{g mL}^{-1}$ of **Im-D2** (Fig. 5K) led to severe matrix fibrosis, low bacterial visibility, and cell wall disruption. The use of **Ag-D2** at a concentration of $32 \mu\text{g mL}^{-1}$ (Fig. 5L) resulted in diluted EPS and high visibility of cells. For **Ag-D1** in combination with endolysin (Fig. 5M), the matrix became compact and wrinkled, and the visibility level dropped to average values. For **Im-D2** (Fig. 5N) in combination with endolysin, EPS remained dense, and bacteria were hardly visible due to abundant coverage. Finally, the **Ag-D2** complex with endolysin (Fig. 5O) led to a dilution of the matrix and an increase in the number of damaged cells, which exhibited numerous cell wall defects.

3.3. Histomorphology of uninfected and *Pseudomonas aeruginosa*-infected pork skin after treatment with dendrimers and their complexes with endolysin

The effects of individual therapies on *ex vivo* skin structures, indicating changes in the stratum corneum (SC), stratum lucidum (SL), stratum granulosum (SG), stratum spinosum (SS), stratum basale (SB) and dermis (D), are shown in Fig. 6. In the control group (Fig. 6A) and after the application of endolysin ($2 \mu\text{g mL}^{-1}$) (Fig. 6B), the entire skin structure remained intact. Treatment with the **Ag-D1** dendrimer ($8 \mu\text{g mL}^{-1}$) (Fig. 6C) caused local flattening (\downarrow) and abrasion of keratinocytes in the stratum corneum, without significant changes in the deeper layers of the epidermis; the dermis remained intact. The addition of endolysin **Ag-D1** ($8 \mu\text{g mL}^{-1} + 2 \mu\text{g mL}^{-1}$) (Fig. 6F) also caused local flattening of the stratum corneum, while hypertrophy (\star) of some cells was observed in the spinous layer. A higher concentration of **Ag-D1** ($32 \mu\text{g mL}^{-1}$) (Fig. 6I) caused dilatation of the stratum corneum, disruption of the continuity (\uparrow) of the cells of the stratum spinosum with the stratum basale of the epidermis, atrophy of the cells of the stratum spinosum, without changes in the dermis. The complex of $32 \mu\text{g mL}^{-1}$ **Ag-D1** + $2 \mu\text{g mL}^{-1}$ endolysin (Fig. 6L) caused thickening of the stratum corneum, hyperplasia and hypertrophy of keratinocytes in the stratum spinosum, while

maintaining the integrity of the dermis. Significant hyperplasia of the stratum corneum and stratum spinosum was observed, without significant changes in the dermis after using the pyrimidine imidazolium **Im-D2** - $8 \mu\text{g mL}^{-1}$ (Fig. 6D) and its complex with endolysin ($8 \mu\text{g mL}^{-1}$ **Im-D2** + $2 \mu\text{g mL}^{-1}$ endolysin) (Fig. 6G). At a higher concentration of $32 \mu\text{g mL}^{-1}$ **Im-D2** (Fig. 6J), a strong flattening of the stratum corneum was observed, with a break in continuity between the SC and SL, while the dermis remained unchanged. The complex with endolysin (Fig. 6M) caused incomplete atrophy of the stratum corneum, as well as hypertrophy of the cells of the stratum spinosum, with the dermis remaining intact. The introduction of pyrimidine **Ag-D2** ($8 \mu\text{g mL}^{-1}$) (Fig. 6E) led to significant SC atrophy and severe hyperplasia of the spinous layer, while the dermis remained unchanged. The complex of $8 \mu\text{g mL}^{-1}$ **Ag-D2** + $2 \mu\text{g mL}^{-1}$ endolysin (Fig. 6H) caused the same changes as in the case of the dendrimer itself. The use of a higher concentration of **Ag-D2** ($32 \mu\text{g mL}^{-1}$) (Fig. 6K) caused local reduction and disruption between the stratum corneum and stratum lucidum, without significant changes in the stratum spinosum and dermis, while the addition of endolysin ($32 \mu\text{g mL}^{-1} + 2 \mu\text{g mL}^{-1}$) (Fig. 6N) caused sporadic disruption of the continuity between the stratum corneum and the stratum lucidum, while maintaining the normal thickness of the stratum corneum and leaving the stratum spinosum and dermis unchanged. The reduction in the toxicity of dendrimers in the presence endolysin could be due to their complexation. Due to its charge, the protein is able to bind to the dendrimer, thus obscuring the active surface groups. Furthermore, local aggregation of dendrimers in the presence of endolysin can quite often occur, thus reducing their properties. All measurements of the thickness of the stratum corneum and the surface area of individual inner layers of the epidermis, *i.e.*, the granular layer, spinous layer, and basal layer, are summarized in Table 2. Overall, the tested compounds have concentration-dependent effects on the structure of the epidermis, while maintaining the integrity of the dermis, and the presence of endolysin tended to mitigate the extent of epidermal disruption induced by the

Table 2 Measurements of the thickness of the stratum corneum and the surface area of the inner layers of the epidermis (granular layer SG, spinous layer SS, and basal layer SB)

	Mean thickness of the stratum corneum (SC) [μm]	$p < 0.001$	Mean surface area of the stratum granulosum, stratum spinosum and stratum basale (SG + SS + SB) [μm^2]	$p < 0.001$
Control	26.68 ± 3.68		89008.5 ± 11367.57	$p < 0.001$
Endolysin $2 \mu\text{g mL}^{-1}$	16.01 ± 6.6		52648.05 ± 12989.59	
$8 \mu\text{g mL}^{-1}$ Ag-D1	14.04 ± 4.65		42352.25 ± 10198.6	
$8 \mu\text{g mL}^{-1}$ Im-D2	22.69 ± 6.59		34443.78 ± 7608.09	
$8 \mu\text{g mL}^{-1}$ Ag-D2	13.59 ± 3.49		60013.19 ± 5580.38	
$8 \mu\text{g mL}^{-1}$ Ag-D1 + $2 \mu\text{g mL}^{-1}$ E	10.02 ± 3.34		58781.8 ± 7462.56	
$8 \mu\text{g mL}^{-1}$ Im-D2 + $2 \mu\text{g mL}^{-1}$ E	14.51 ± 5.44		64693.54 ± 10175.75	
$8 \mu\text{g mL}^{-1}$ Ag-D2 + $2 \mu\text{g mL}^{-1}$ E	14.79 ± 4.94		60489.31 ± 6394.41	
$32 \mu\text{g mL}^{-1}$ Ag-D1	16.1 ± 4.47		70132.42 ± 13635.32	$p = 0.008$
$32 \mu\text{g mL}^{-1}$ Im-D2	12.63 ± 3.97		52961.31 ± 8276.02	$p < 0.001$
$32 \mu\text{g mL}^{-1}$ Ag-D2	12.69 ± 3.19		55602.58 ± 4384.04	
$32 \mu\text{g mL}^{-1}$ Ag-D1 + $2 \mu\text{g mL}^{-1}$ E	16.26 ± 5.95		51156.31 ± 7975.58	
$32 \mu\text{g mL}^{-1}$ Im-D2 + $2 \mu\text{g mL}^{-1}$ E	13.66 ± 3.89		52731.35 ± 6431.55	
$32 \mu\text{g mL}^{-1}$ Ag-D2 + $2 \mu\text{g mL}^{-1}$ E	21.28 ± 5.98		50038.99 ± 7576.00	



dendrimers, although the magnitude of this effect varied between dendrimer types and concentrations.

Structural changes in the biofilm of *Pseudomonas aeruginosa* PAO1 and layers of skin after 24 hours of treatment with dendrimers at concentrations of 8 and 32 $\mu\text{g mL}^{-1}$, their complexes with endolysin (2 $\mu\text{g mL}^{-1}$) are presented at Fig. 7. In the control group (Fig. 7A) with the stratum corneum (SC) detached in places and surrounded by dense biofilm, the

epidermis (E) was degraded, while the dermis (D) was partially degraded by bacteria, which manifested itself in places with a flat surface. The administration of endolysin (Fig. 7B) did not change this condition. Treatment with 8 $\mu\text{g mL}^{-1}$ **Ag-D1** (Fig. 7C) and 8 $\mu\text{g mL}^{-1}$ **Ag-D1** + 2 endolysin (Fig. 7F) did not reduce biofilm formation or SC or E reconstruction, and affected the D as described above. The use of a higher concentration of 32 $\mu\text{g mL}^{-1}$ **Ag-D1** (Fig. 7I) or its complex with

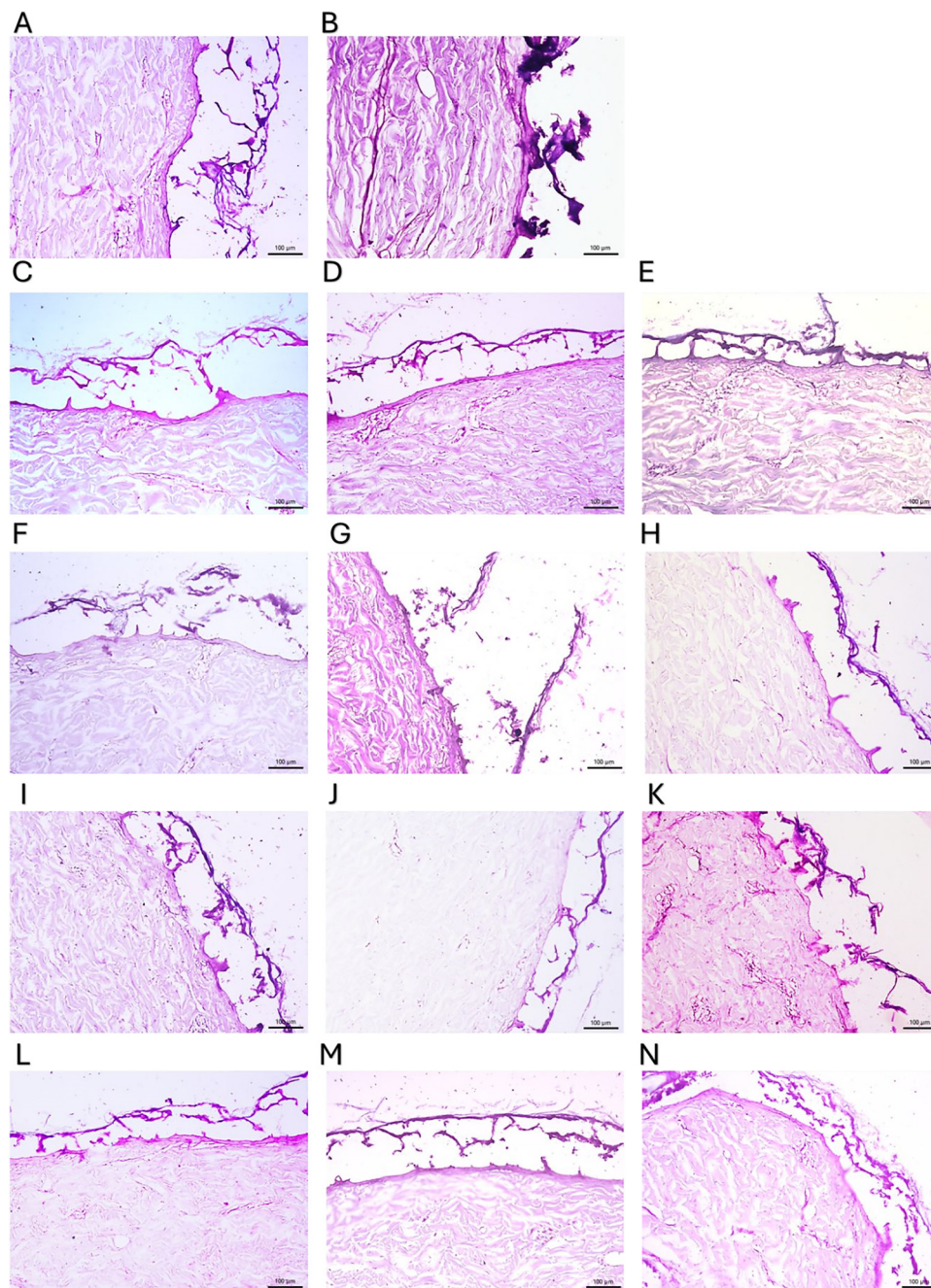


Fig. 7 Histomorphology of cryosections of infected pork skin with bacterial biofilm formed by *Pseudomonas aeruginosa* PAO1. (A) Untreated infected skin. Twenty-four hours after treatment (B) with endolysin, (C) with 8 $\mu\text{g mL}^{-1}$ **Ag-D1**, (D) with 8 $\mu\text{g mL}^{-1}$ **Im-D2**, (E) with 8 $\mu\text{g mL}^{-1}$ **Ag-D2**, (F) with 8 $\mu\text{g mL}^{-1}$ **Ag-D1** + 2 $\mu\text{g mL}^{-1}$ endolysin, (G) with 8 $\mu\text{g mL}^{-1}$ **Im-D2** + 2 $\mu\text{g mL}^{-1}$ endolysin, (H) with 8 $\mu\text{g mL}^{-1}$ **Ag-D2** + 2 $\mu\text{g mL}^{-1}$ endolysin, (I) with 32 $\mu\text{g mL}^{-1}$ **Ag-D1**, (J) with 32 $\mu\text{g mL}^{-1}$ **Im-D2**, (K) with 32 $\mu\text{g mL}^{-1}$ **Ag-D2**, (L) with 32 $\mu\text{g mL}^{-1}$ **Ag-D1** + 2 $\mu\text{g mL}^{-1}$ endolysin, (M) with 32 $\mu\text{g mL}^{-1}$ **Im-D2** + 2 $\mu\text{g mL}^{-1}$ endolysin and (N) with 32 $\mu\text{g mL}^{-1}$ **Ag-D2** + 2 $\mu\text{g mL}^{-1}$ endolysin. The scale bar is 100 μm (cross-sections, magnification 100 \times , 1% toluidine blue staining).



endolysin (Fig. 7L) did not prevent damage to the epidermis and dermis. Similarly, $8 \mu\text{g mL}^{-1}$ **Im-D2** (Fig. 7D) and **Im-D2** + 2 endolysin (Fig. 7H) did not restore the integrity of SC or E, while $32 \mu\text{g mL}^{-1}$ **Im-D2** (Fig. 7J) and **Im-D2** + 2 endolysin (Fig. 7M) caused their complete detachment, leaving SC intact. The introduction of chelated dendrimer **Ag-D2** at $8 \mu\text{g mL}^{-1}$ (Fig. 7E) and $8 \mu\text{g mL}^{-1}$ + 2 endolysin (Fig. 7G) also did not prevent SC detachment or E loss, similar to $32 \mu\text{g mL}^{-1}$ **Ag-D2** (Fig. 7K) and **Ag-D2** + 2 endolysin (Fig. 7N). In each of these cases, the biofilm adhered to the completely detached stratum corneum, no deeper layers of the epidermis were observed, and the dermis remained degraded, manifesting itself as a flattened surface. This means that none of the tested combinations prevented the mechanical destruction of the skin layers by the PAO1 biofilm (Fig. 8).

Pseudomonas aeruginosa forms clusters of cells that facilitate its attachment to the wound and secretes various enzymes and toxins, such as type A exotoxins, the endotoxin LPS, and elastase, which damage host tissues and immune system cells, leading to necrosis and intensifying the inflammatory process.⁵² Additional tests must be performed to check whether, after such an advanced bacterial infection, tissue regeneration would be possible in connection with the preservation of the structure of the dermis, despite the penetration of bacteria deep into the remaining tissue.

3.4. Cytotoxic effects on eukaryotic cells

Dendrimers used as wound dressings in skin infections exhibit antibacterial properties; however, their impact on the viability and function of skin cells cannot be overlooked. Therefore, an experiment was conducted to assess the viability and proliferative capacity of VH10 cells in the presence of increasing concentrations of dendrimers. As shown in Fig. 9A, low concentrations of the tested metallodendrimer dendrimer

(**Ag-D1**) – (Fig. 9A) ($1\text{--}4 \mu\text{g mL}^{-1}$) did not significantly affect cell viability, which remained at an average of $\sim 96\%$. A slightly higher concentration ($8 \mu\text{g mL}^{-1}$) led to a marked decrease in survival ($\sim 50\%$), and the highest concentrations ($16, 32, 64,$ and $128 \mu\text{g mL}^{-1}$) further reduced viability to approximately 28% . Based on the results, concentrations that exhibited low or no cytotoxicity were chosen, and endolysin was added at doses of 1 and $2 \mu\text{g mL}^{-1}$. As shown in graphs, endolysin alone, at both tested concentrations, did not exhibit cytotoxic effects. Furthermore, the addition of endolysin to **Ag-D1** at 4 and $8 \mu\text{g mL}^{-1}$ did not induce cytotoxicity (cell viability $\sim 91\%$). However, at $16 \mu\text{g mL}^{-1}$, survival decreased to $\sim 52\%$ and $\sim 42\%$, respectively, and at 32 and $64 \mu\text{g mL}^{-1}$, viability dropped to $\sim 30\%$ in the presence of endolysin, indicating that high dendrimer concentrations remain cytotoxic despite the addition of the protein.

Imidazolium dendrimer **Im-D2** (Fig. 9B) exhibited lower cytotoxicity than **Ag-D1**; at concentrations of 8 and $16 \mu\text{g mL}^{-1}$, no significant reduction in cell survival was observed, with viability maintained at around 95% . A moderate decrease to $\sim 50\%$ was observed at $32 \mu\text{g mL}^{-1}$, while the highest concentrations (64 and $128 \mu\text{g mL}^{-1}$) reduced survival to $\sim 30\%$. In contrast, the addition of endolysin to **Im-D2** at lower concentrations ($4\text{--}8 \mu\text{g mL}^{-1}$) did not result in cytotoxicity, with average cell viability exceeding $\sim 100\%$. At an **Im-D2** concentration of 16 and $32 \mu\text{g mL}^{-1}$ combined with a low dose of endolysin, the cell survival rate was approximately 60% . The use of endolysin with dendrimer concentrations of $64 \mu\text{g mL}^{-1}$ and $128 \mu\text{g mL}^{-1}$ resulted in a fibroblast survival rate of $\sim 30\%$. The metallodendrimer **Ag-D2** (Fig. 9C) showed an intermediate cytotoxic profile.

Concentrations ranging from 1 to $8 \mu\text{g mL}^{-1}$ maintained the cell viability at $\sim 102\%$, while 16 and $32 \mu\text{g mL}^{-1}$ resulted in a survival rate of $\sim 77\%$. At dendrimer concentrations of 64 and $128 \mu\text{g mL}^{-1}$, the cell viability decreased to $\sim 30\%$. In contrast,

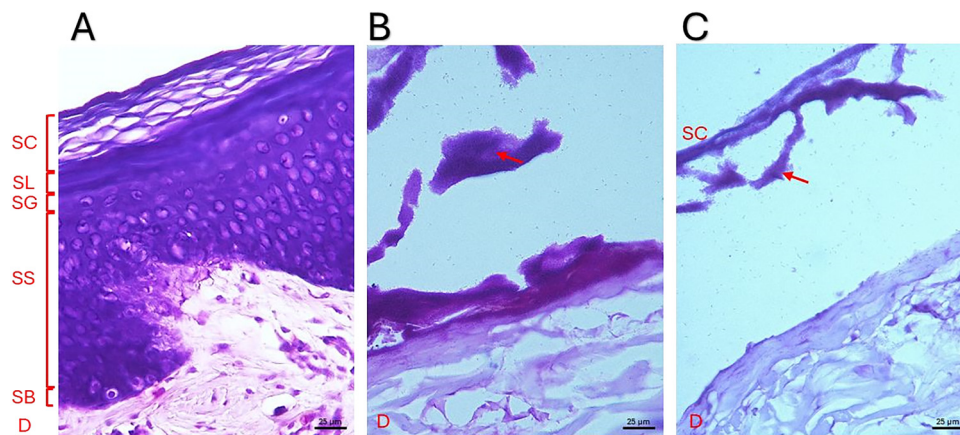


Fig. 8 Comparison of the histomorphology of cryosections of pork skin tissue from uninfected and *Pseudomonas aeruginosa*-infected groups. (A) Uninfected skin tissue with a clear division into layers of the epidermis: stratum corneum (SC), stratum lucidum (SL), stratum granulosum (SG), stratum spinosum (SS), stratum basale (SB) and dermis (D). (B, C) *Pseudomonas aeruginosa*-infected skin tissue with distinct clusters of bacteria – bacterial biofilm – in the presence of detached stratum corneum with a distinct absence of deeper layers of the epidermis and near the surface of the dermis. The scale bar is $25 \mu\text{m}$ (cross-sections, magnification $400\times$, 1% toluidine blue staining).



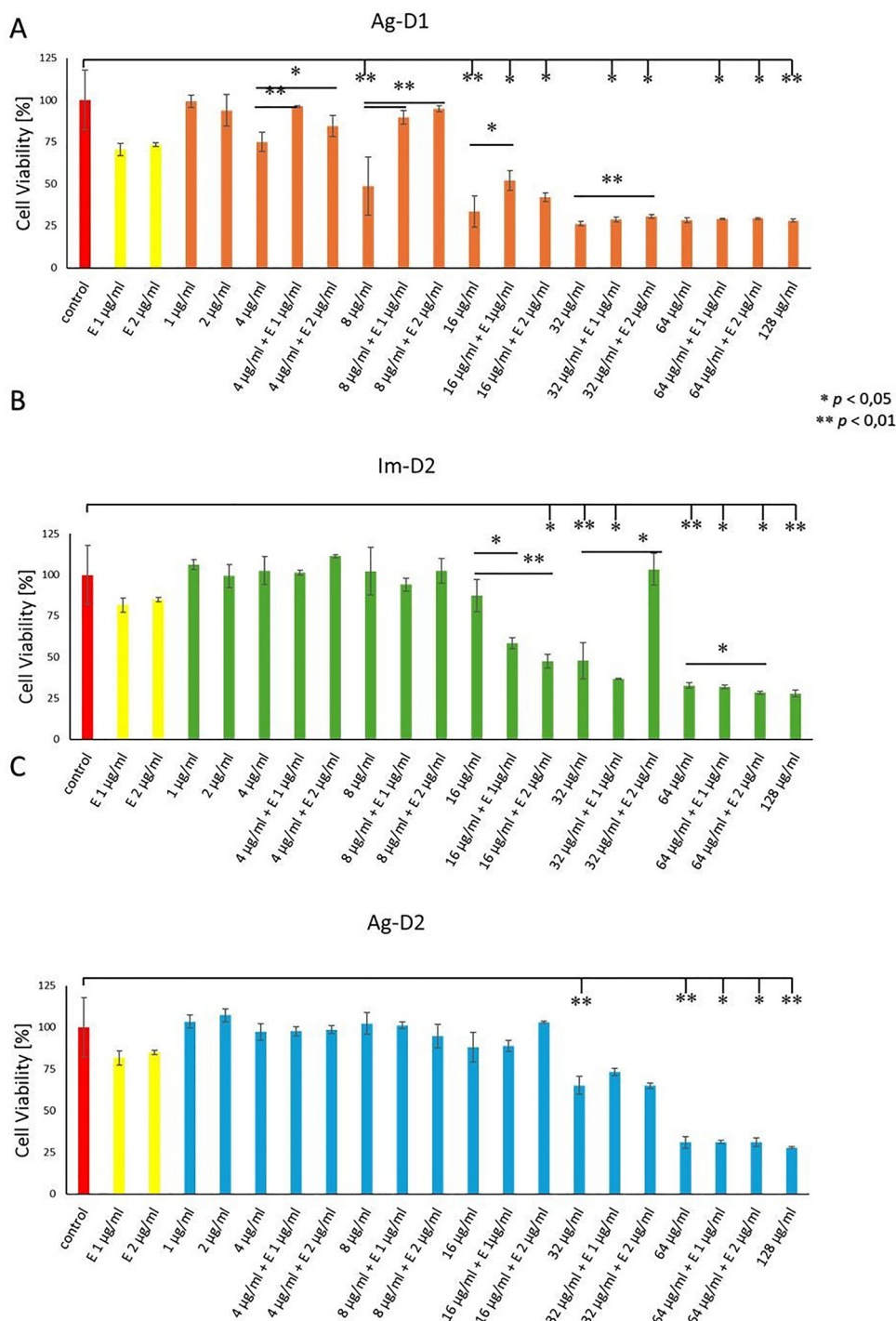


Fig. 9 The viability of VH10 cells treated with metallo dendrimers and complexes with endolysin (E $1 \mu\text{g mL}^{-1}$ and $2 \mu\text{g mL}^{-1}$) after 24 h incubation. (A) **Ag-D1**, (B) **Im-D2**, (C) **Ag-D2**. Untreated cells were used as the control. Experiments were performed in triplicate and results are presented as a percentage of the control.

Ag-D2 at $16 \mu\text{g mL}^{-1}$ in combination with endolysin showed no significant effect on cell viability. Moreover, the addition of 1 or $2 \mu\text{g mL}^{-1}$ of endolysin to 32 and $64 \mu\text{g mL}^{-1}$ of dendrimer did not reduce cytotoxicity, resulting in a comparable cell survival of $\sim 65\%$ and $\sim 31\%$, respectively.

4. Conclusions

In this study, we have demonstrated that carboxilane CBS dendrimers, imidazolium (**Im-D2**) and carboxilane metallo dendrimers-based silver nanocomposites (**Ag-D1** and **Ag-D2**), when combined



with a catalytic CHAP-domain endolysin, act as a modern combat tool against fully formed and highly resistant wild-type *Pseudomonas aeruginosa* PAO1 bacterial biofilm in *in vitro* and *ex vivo* imitations of animal skin infection. On comparing the effectiveness of dendrimer therapy combined with endolysin against *P. aeruginosa* biofilm in an *in vitro* model on a smooth plastic surface and *ex vivo* on a rough, irregular pork skin, a more pronounced reduction in bacterial viability and cellular damage was observed *in vitro*. This is due to both the higher toxicity of the tested compounds in a simple laboratory environment and the fact that the irregular texture of the skin and richer nutrient resources in the *ex vivo* model favored the survival of microorganisms. The absence of significant skin damage after the application of dendrimers and their complexes with endolysin indicates that early implementation of therapy could effectively eliminate bacteria without compromising tissue integrity. At the same time, histomorphological analysis revealed that even a 24-hour infection leads to excessive damage to the epidermis, and the bacteria became highly resistant to dendrimers, making it difficult to assess the effectiveness of late interventions. It is worth noting that a significant amount of time often passes between the onset of infection and the appearance of clear pathological changes, diagnosis, and obtaining medical assistance (including the implementation of targeted antibiotic therapy). For example, in an analysis of severe soft tissue infections, the average time from symptom onset to hospital admission was approximately 4.5 days. During this period, the infection may already reach an advanced stage, accompanied by tissue damage and biofilm maturation, which hinders effective pathogen elimination. Therefore, the 24-hour infection variant used in our study has significant translational and therapeutic value; it reflects the stage at which intervention is undertaken after a certain delay, and the effectiveness of treatment may be limited by developed biofilm and tissue changes.

Author contributions

Karolina Lach: writing – original draft, methodology, investigation, formal analysis, data curation. Samuel Takvor-Mena: methodology, investigation, writing – review & editing. Oscar Barrios-Gumiel: methodology, investigation. Javier Sanchez-Nieves: writing – review & editing, supervision, funding acquisition, conceptualization. Jacek Kuchinka: methodology, investigation. Piotr Furmańczyk: methodology, investigation. Małgorzata Łysek-Gładysińska: methodology, investigation. Karol Ciepluch: writing – review & editing, supervision, project administration, methodology, funding acquisition, conceptualization.

Conflicts of interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

Data availability

The data will be made available from the corresponding author upon reasonable request.

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