Efficacy of Chlorhexidine, Polihexanide and Tissue-Tolerable Plasma against *Pseudomonas aeruginosa* Biofilms Grown on Polystyrene and Silicone Materials

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**Introduction**

The formation of biofilms is crucial in the pathogenesis of many acute and subacute microbial infections, including chronic wounds and foreign-body-related infections [1, 2]. Bacteria organized in biofilms are distinctly less susceptible to host defences and antimicrobial therapy compared with their planktonic counterparts [3–5]. Topical antimicrobial therapy with antiseptics or with tissue-tolerable plasma (TTP) could be an interesting option for biofilm elimination, given that these measures are effective in the inactivation and removal of microbial biofilms, but data available on this topic are limited.

We therefore compared the efficacy of chlorhexidine digluconate (CHX) with that of polihexanide (polhexamethylene biguanide, PHMB) and TTP against *Pseudomonas aeruginosa* biofilm grown in microtitre plates (polystyrene) and on silicone materials in an artificial wound fluid.

**Key Words**

Chlorhexidine · Polihexanide · *Pseudomonas aeruginosa* · Biofilm efficacy · Tissue-tolerable plasma

**Abstract**

**Background:** The formation of biofilms is crucial in the pathogenesis of many acute and subacute microbial infections, including chronic wounds and foreign-body-related infections. Topical antimicrobial therapy with chemical antiseptics or physical treatment with tissue-tolerable plasma (TTP) may be promising to control bacterial infection. **Methods:** We assessed the efficacy of 0.1% chlorhexidine digluconate (CHX), 0.02 and 0.04% polihexanide (polhexamethylene biguanide, PHMB) and of TTP against *Pseudomonas aeruginosa* SG81 biofilm grown in microtitre plates (polystyrene) and on silicone materials in an artificial wound fluid. **Results:** Overall, PHMB was as effective as CHX in reducing the total amount of biofilm (gentian violet assay) and in reducing the bacterial metabolism in biofilms (XTT assay). TTP also led to a significant reduction in colony-forming units. **Conclusion:** The antimicrobial activity of PHMB in biofilms is comparable to that of CHX. TTP could become an interesting physical alternative to chemical antisepsis in the future.
minal chlorobenzene substituents, it was to be compared with chlorhexidine in its effect against biofilms. A further reason for the choice of polihexanide is its growing importance as an antiseptic substance and its related increasing use, particularly for the treatment of infected wounds and for decolonization of skin [Hübner and Kramer, this supplement issue; 12–14], as well as for the decolonization of mucous membranes of patients colonized with methicillin-resistant *Staphylococcus aureus* [Eberlein and Assadian, this supplement issue].

The use of TTP as a physical alternative to chemical antisepsis appears promising, because plasma, too, depending on the dose, is microbicidally effective [15, 16] and in the body temperature range can also be used on the surface of the body [17]. Unlike chemical antisepsis, energy can be supplied to the wound tissue through the use of plasma, thus possibly contributing to promoting the healing process. The possibility of the use of TTP worldwide is therefore being studied in centres specializing in this.

**Materials and Methods**

**Bacterial Strains and Growth Conditions**

The bacterium used in this study was a mucoid environmental *P. aeruginosa* strain SG81 which was isolated from a biofilm in a technical water system and was provided by the biofilm center Duisburg, Germany. This strain is a well-characterized and stable alginate-producing strain that forms highly mucoid colonies on standard media agar. The content of the alginate was subclassified in mannose and glucose residues in which the mannose residues outweigh with 67% [18, 19]. The bacteria were grown overnight on Columbia blood agar at 37°C for 24 h. A single colony was then transferred to trypticase soy agar plates (Oxoid, Cambridge, UK) and was incubated for 24 h at 37°C. The grown cells were suspended in phosphate-buffered saline (PBS), harvested by centrifugation for 15 min at 3,000 g, washed twice in PBS and resuspended in artificial wound fluid (minimal essential medium + 10% fetal bovine serum) (GIBCO-Invitrogen, Karlsruhe, Germany) [11] to a final concentration of approximately 1 × 10⁹ colony-forming units (CFU)/ml.

**Culture of the Biofilms and Preparation of the Samples**

Biofilms were cultured on test objects made of silicone with a diameter of 0.6 cm (Thomasil-60, Reichelt Chemietechnik, Heidelberg, Germany) and in the wells of cell culture 24-well microtitre plates (Techno Plastic Products AG, Switzerland).

The final bacteria medium consisted of the artificial wound and the washed bacteria at a final concentration of 10⁶ CFU/ml. The sterile test objects were positioned in 24-well microtitre plates, covered with 0.7 ml microorganism suspension, and incubated aerobically on the agitator (Polymax, Heidolph, Germany) at 160 rpm for 4 h at 37°C. For the direct culture, on 96-well microtitre plates, 50 μl microorganism suspension per well was applied by pipette and likewise incubated. After 4 h, the suspension was drawn off and replaced by sterile medium; after renewed incubation, the medium was changed every 8 h. After 44 h, the medium was drawn off and the test object/well was washed with PBS.

**Test Substances and Antiseptic Treatment**

CHX was used as a 0.1% aqueous solution (production by the pharmacy of Greifswald University Hospital). For production of the PHMB solutions, Cosmocil, a 20% PHMB solution (Arch Bio- cides, UK) was used.

The test objects were transferred into new, sterile microtitre plates, covered with 0.9 ml of the antiseptic, and incubated. After 30 min, the antiseptic was drawn off, and the antiseptic effect was halted by adding 1 ml inactivator (see below). In the case of the 96-well microtitre plates, 90 μl antiseptic was added, and the effect halted with 100 μl inactivator. PBS was used as the control.

Both antiseptics were inactivated using 40 g/l Tween 80, 30 g/l saponin, 4 g/l lecithin, 10 g/l SDS and 1 g/l sodium thioglycolate. The inactivation of the antiseptics by the inactivator was proved in the quantitative suspension test according to DIN 1040 (data not shown).

**Treatment with TTP**

As the plasma source, an HF plasma pen (frequency 1.82 MHz, input power 3 W; KINPen 09, INP Greifswald; fig. 1) was used with argon as the carrier gas [20, 21]. During treatment, the plasma generated in the jet (spatial afterglow plasma, effluent) is directed to the treated surface at an argon flow rate of 5 standard l/min, the temperature at the plasma tip being 42°C. At this temperature, a mean heat output of about 150 mW is generated on the surface.

For plasma use, the plasma source was attached to a computer-controlled x/y/z-table and the test object or microtitre plate was positioned below it (fig. 1). The distance from the test object or bottom of the microtitre plate was 7 mm. The entire surface of the...
test object was treated meander-like at a speed of 10 mm/s. The whole time of treatment was 60 s. The samples were also treated with inactivation solution to destroy possible existing radicals.

**Determination of the CFU**

After 5 min of inactivation, biofilms were detached in 5 ml PBS using sonication for 20 min at 130 W. CFU were determined by serial dilutions, which were plated on trypticase soy agar and incubated for 24 h at 37°C.

**Determination of the Biofilm Mass Using Gentian Violet Assay**

The evaluation was carried out according to published methods [22–24]. After 5 min of inactivation, the inactivator was drawn off, the objects were rinsed with fresh PBS and dried for at least 60 min at 37°C. The samples were then covered with 0.1% gentian violet (GV), incubated for 15 min, washed three times with distilled water, and then decoloured with 96% v/v ethanol HCl (Merck, Darmstadt, Germany). The supernatant was pipetted off and the extinction detected in the photometer at 590 nm (reference value 620 nm).

**Confocal Laser Scanning Microscopy**

Two different dyes, Live/Dead (BacLight, Invitrogen, Darmstadt, Germany, SKU L-7012) and acridine orange (AppliChem-BioChemica, Darmstadt, Germany), were used to visualize the components of the biofilm. The samples were transferred into a 24-well microtitre plate and incubated immediately according to the manufacturer’s instructions. After incubation and fixation, samples were rinsed 5 times with PBS to remove dye residues. Samples were then placed onto glass object slides covered with 10 μl Mowiol 4-88 (Polysciences Inc., Eppelheim, Germany) and a coverslip placed on top. Samples were observed using a Zeiss CLSM510 Exciter confocal laser scanning microscope with a 488-nm argon laser. Microscope work was carried out using the oil immersion Plan-Neofluar 63×/1.4 Oil DIC objective.

**Biostatistics**

All tests were carried out sixfold each for the CFU, biofilm mass determinations, and in duplicate for the staining. In addition, for each test, 6 samples were used as controls. The Friedman test was used for testing for global statistical differences in micro-organism count, biofilm mass and vitality, and if the result was significant, further testing was carried out with the Wilcoxon test. The level of significance was set at 0.05.
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**Results**

**GV Assay**
For the microtitre plates, the results of the extinction measurement of the samples stained with GV showed no significant differences between the treatments (Friedman test, p = 0.702) (table 1). On the silicone test objects, the treatment with antiseptics likewise led to no significant effect (table 2). Negative values are explained by the subtraction of the control values. In contrast, plasma and gas led to a significant increase in the extinction compared with control and CHX (table 2).

**XTT Assay**
The converted amount of XTT stain as a measure of the metabolic activity of the microorganisms in the biofilm was significantly reduced by the antiseptics in the microtitre plate, while plasma and gas did not lead to a significant reduction (table 1). On the silicone test objects, only 0.04% PHMB led to a significant reduction compared to the control, while, as in the microtitre plate, plasma and gas led to an increase in the extinction that differed significantly from CHX and PBS (table 2).

**Colony-Forming Units**
The antimicrobial effect of the antiseptics and of the plasma was seen most clearly in the reduction of the CFU. Both in the microtitre plates and on the silicone test objects the treatments (with the exception of 0.02% PHMB in the microtitre plate) led to a significant reduction in the CFU compared with the control, although there were no significant differences between the treatments. Only the plasma control gas on silicone and 0.02% PHMB in the microtitre plate were poorer than CHX (tables 1 and 2).

**Fluorochroming**
Acridine orange intensely fluoresces after intercalating with the DNA/RNA in living and dead bacteria. Compared to the untreated sample, the number of microorganisms in the preparation stained with acridine orange is clearly reduced (fig. 2). The clearest effects were seen for 0.1% CHX, TTP and 0.04% PHMB. The live/dead-stained preparation permits additional information about the vitality of the microorganisms in the biofilm (fig. 3). The dense, vital ‘massive microorganism growth’ in the control is killed by the antimicrobial treatments and reduced in its density. Live/dead-staining included the fluorochromes Syto9 and propidium iodide, both intercalating the DNA. Worth noting are the differences between gas and TTP on the one hand and between 0.02 and 0.04% PHMB on the other. Both with TTP and with 0.04% PHMB, hardly any vital microorganisms stain after treatment. This is in conformity with the preparations

Fig. 2. Effect of antiseptics and TTP against biofilms. a Control in PBS. b CHX 0.1%. c PHMB 0.02%. d PHMB 0.04%. e TTP. f Argon (gas control). Bacteria stained with acridine orange.
stained with acridine orange, after which TTP and the higher PHMB concentration even inactivate ‘islands’, i.e. areas of high microorganism density.

Discussion

Biofilms make treatment of microbial infections more difficult. This applies especially if abiotic foreign bodies are present in the wound and form a basis for biofilm formation that is difficult for the immune system to access [1]. For accessibility to microbial biofilms, topical treatment with antiseptic measures is superior to the chemotherapeutic use of antibiotics, and in view of the increasing microbial resistance to antibiotics, it is of particular significance. In the study described here, the effects of PHMB and TTP in vitro in the presence of artificial wound fluid were compared with that of chlorhexidine as a substance with a proven effect against biofilms and seen as the gold standard to eliminate dental plaque [28–31]. However, it must be remembered that, depending on the test system, chlorhexidine also proved to be ineffective against biofilms [9]. The effectiveness of PHMB against biofilms was also shown [2, 32–34]. First investigations on the antimicrobial abilities of physical plasma against bacteria in biofilms have been published [35–37]. In the present study, the biofilms were cultured not only on routinely used microtitre plates, but also on silicone as a material frequently used in medicine with organic burden.

The results show that PHMB is comparable to CHX in its effect against the P. aeruginosa biofilm, irrespective of the material. In addition to efficacy, local tolerability and systemic toxicity play a decisive role for clinical use. PHMB has proved to be well tolerable and tissue-friendly [38], and in these respects is clearly superior to CHX [see Hübner and Kramer, this supplement issue]. On account of the deficient data situation on the clinical efficacy of CHX for wound treatment [39] and its tissue toxicity [11], in Germany CHX-based wound antiseptics are no longer authorized. The results in the biofilm model substantiate the results of clinical studies in which the potential of PHMB and products containing PHMB for the treatment of infected wounds could be demonstrated [14, 40–42].

TTP is a physical alternative to chemical antiseptics. In recent years, plasma sources for wound treatment have been developed and tested at several centres [43–46]. Two
advantages of physical procedures are the fact that they can be standardized and the absence of entry of the antiseptic into the wound. In our tests, the microbicidal effect of plasma was comparable to that of CHX, whereas the simple gassing with argon proved to be significantly poorer than CHX. This confirms that the antimicrobial effect is not just a result of drying by the gas current, but truly of the plasma itself. These reductions through TTP were not only statistically significant, but with \( >3 \log_{10} \) correspond to the required efficacy for antiseptics [47]. This confirms the results of other authors reporting on the antimicrobial effects of TTP [36, 48].

However, the results regarding the determination of the biofilm mass could only be evaluated for the microtitre plates, as the drying caused by the flow of gas apparently altered the stainability on silicone.

To ensure the reproducibility of the results, standard methods of biofilm research were used. For the method of analysis of the solid substance, GV was used. This method is applied in many variations, which all follow the same scheme [22, 23]. In contrast, the XTT test was modified with regard to the combination with menadione and N-methylphenazinium methyl sulphate [25–27], because in the results of our previous studies in the case of \( P. \) aeruginosa the most intensive conversion of XTT was measured with this combination (data not shown). Nevertheless, the study has numerous limitations that must be taken into consideration in the evaluation. Evaluation of the metabolic activity by means of XTT must be made with caution for microorganism counts \( <10^5 \) CFU/ml on account of the S-shaped calibration curve for the stain. This particularly applies for the values for TTP and gas in the microtitre plate. To make the test conditions as similar as possible to the conditions in vivo, all tests were carried out in the presence of artificial wound fluid. However, this makes comparison with the results of other authors more difficult, as in the absence of such a load greater reductions tend to be achieved. With microtitre plates (polystyrene) and silicone, the choice of samples involved two materials that are frequently used in biofilm research. The results show that the results cannot be extrapolated uncritically to other materials, as just the two materials studied showed clear differences. The same applies for the plasma source used, as other configurations, especially changes in gas flow, energy density and type of plasma produced can lead to other results. Further investigations are therefore necessary in order to verify the efficacy for other microorganisms and under other conditions.

**Conclusion**

Biofilms play a key role in the pathogenesis of many acute and subacute microbial infections, including chronic wound and foreign-body-related infections. CHX and PHMB have been shown to be effective against bacteria in biofilms and could therefore be an interesting option for biofilm treatment. Topical antimicrobial therapy with TTP was also shown to be effective against \( P. \) aeruginosa biofilms on polystyrene and silicone materials. Further research should be done to evaluate the possible role of physical therapies for biofilm-derived infections.

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**Disclosure Statement**

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**References**


