Research Article

Galleria mellonella as a Model to Study the Effect of Antimicrobial Surfaces on Contamination by Staphylococcus aureus

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Abstract

Objective: The role of the inanimate environment in the transmission of healthcare-associated infections (HCAIs) is increasingly recognized. Hence, several self-disinfecting surfaces have gained attention as new infection control strategies. The objective of this study was to assess the antimicrobial efficacy of membrane-active polycation (maPK) surfaces against Staphylococcus aureus (S. aureus) applying the in vivo Galleria mellonella (G. mellonella) contamination model.

Methods: In the first set-up maPK surfaces were contaminated with S. aureus and G. mellonella larvae were placed on the surface for 45 min. In the second set-up larvae were contaminated with S. aureus and transferred to maPK surfaces. Finally, the cross-over rates to larvae and maPK-surface were determined. Additionally, log reduction values (LRVs) of S. aureus on contaminated maPK and larvae were calculated.

Results: The first experimental set-up revealed a significantly higher cross-over rate of S. aureus to the larvae from contaminated control surface (~2 x 10³ colony forming units (cfu)/mL) when compared to maPK surface (~5.1 x 10¹ cfu/mL).
In the second set-up the cross-over rate of *S. aureus* from larvae to the control surface was significantly higher (~5.1 x 10³ cfu/plate) than to the maPK surface (~2.1 x 10¹ cfu/plate).

In both experiments LRVs were superior on maPK with 98.1% (p=0.0117) and 99.5% (p=0.0001) reduction, respectively.

**Conclusions:** This in vivo study reveals the ability of maPK surfaces to reduce bacterial contamination and highlights the potential power to avoid cross-contamination from environmental surface.

**Keywords:** Healthcare-associated infections; Antimicrobial surfaces; Environmental contamination; Cross-transmission; *Staphylococcus aureus*; *Galleria mellonella*

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**Citation**

Introduction
Nosocomial infections represent a major problem complicating patient care and result in increased morbidity and mortality [1]. There is growing evidence that exogenous sources such as surface contamination may serve as an important reservoir for pathogens and hence may contribute to their transmission [2-4]. High level of microbial contamination with relevant pathogens including multidrug-resistant microorganisms distant and close to patients is reported [5-7]. Environmental persistence has been shown for Staphylococcus aureus (S. aureus), methicillin-resistant S. aureus, vancomycin-resistant enterococcus, Acinetobacter baumannii, and Clostridium difficile [8, 9]. Cleaning and disinfection procedures display basic measures in infection control, however, multiple studies show gaps in decontamination practices [10, 11]. Additionally, surfaces can rapidly become recontaminated after cleaning and disinfection [12].

Antimicrobial surfaces have been proposed as an additive tool to reduce environmental contamination and possibly prevent healthcare-associated infections (HCAIs) [3, 13]. Polycationic coated surfaces which are microbicidal upon contact were demonstrated to kill microbes in vitro by causing physical damage to the microbe’s cell envelope [14]. We recently showed antimicrobial activity of a surface coated with membrane-active polycations (maPK) against S. aureus ATCC 6538 in vitro [15]. However, whether there is a real clinical benefit is still a matter of debate. For various antimicrobial surfaces implementation of antimicrobial surfaces in the clinical setting has demonstrated several problems resulting in a reduced efficacy compared to in vitro results.

The aim of this follow-up study was to further validate the biocidal capacity of the maPK surface in an in vivo model. For this purpose we used Galleria mellonella larvae and evaluated whether maPK surface prevents growth and cross-over of S. aureus ATCC 6538 from surface to larvae and vice versa.

Methods
Bacterial strains
S. aureus (ATCC 6538) cultured on Columbia agar containing 5% sheep blood (CBA; Becton Dickinson, Heidelberg, Germany) at 35 ± 2°C was used as test-organism. The bacterial suspension was prepared by suspending a 24 h-old colony into 2 mL aqua destillata and adjusting it to a standard turbidity equivalent of 0.5 density units on the McFarland Scale (approximately 1.5 x 10^8 colony forming units (cfu)/mL) using DensiCHECK™ Plus (bioMérieux, France).

Preparation of the larvae and evaluation of their cuticle microbiota
Sixth-instar larvae (SAGIP, Bagnacavallo, Italy) weighing 0.3-0.4 g were kept at 18°C in the dark prior to use. Since the study protocol intended the use of S. aureus as test organism in the wax moth larvae model the natural occurrence of this microorganism on the exoskeletal cuticle of the insect was assessed in a pilot experiment. For this purpose, a modified protocol of the roll-plate culture method by Maki et al. [16] was performed. Briefly, five randomly chosen larvae were each rolled back and forth across the surface of CBA (Becton Dickinson, Heidelberg, Germany) and Mannitol salt agar (MSA; Oxoid, Wesel, Germany) at least three times. After incubating the plates for 24h at 35 ± 2°C microorganisms were identified with standard microbial methods, including Matrix-assisted laser desorption
ionization time-of-flight mass spectrometry (MALDI-TOF®️, Bruker, Bremen, Germany).

**Determination of S. aureus burden on larvae’s cuticle after contamination**

As a further pilot experiment insects’ bioburden on the cuticle after *S. aureus* contamination was ascertained. Larvae (n=5) were contaminated by immersing the latter into an inoculum containing approximately 1.5 x 10⁸ cfu/mL of *S. aureus* for 5 sec. To determine residual burden of *S. aureus* on the cuticle immediately after dipping, excess fluid was blotted off and larvae were immersed separately into sterile phosphate buffered saline (PBS) for 15 sec by slightly shaking. The suspension was serially diluted and plated onto MSA (Oxoid, Wesel, Germany) agar. The plates were incubated at 35 ± 2°C for 24h, bacterial load was quantified in cfu/mL and larvae were disposed.

**Assessment of maPK using the Galleria mellonella model to simulate real life conditions**

To evaluate the antimicrobial potency of the surface two test proceedings were used (Figure 1).

![Flow chart regarding the principle of the two test procedures](image)

**Figure 1**: Flow chart regarding the principle of the two test procedures

maPK: membrane-active polycation

In the first set-up maPK combined with an acrylate dispersion applied on melamine resin layer (Fritz Egger GmbH & Co. OG, Austria) was contaminated with *S. aureus* (1.5 x 10⁸ cfu/mL) on a 30 mm radius as described previously [15]. Samples containing acrylate without maPK were used as control. Subsequently, five larvae were simultaneously transferred onto the contaminated maPK and control surfaces, respectively,
and removed after 45 min. Residual microbial contamination on the surface was determined by contact Rodac™ plates containing MSA (Oxoid, Wesel, Germany). To assess the crossover-rate of *S. aureus* from surface to larvae, insects were finally immersed in sterile PBS each for 15 sec by slightly shaking. Suspensions were serially diluted with PBS and spread on MSA (Oxoid, Wesel, Germany).

In the second set of experiments the vice-versa situation was examined by colonizing the larvae and measuring the spread of *S. aureus* from the larvae to the surface. Superficial colonization of *G. mellonella* was simulated by immersing the larvae (n=5) into an inoculum containing approximately $1.5 \times 10^8$ cfu/mL of *S. aureus* for 5 sec. Colonized larvae were placed to maPK and control surface for 45 min. Residual microbial contamination on the larvae’s cuticle was determined by immersing the larvae in PBS each for 15 sec by slightly shaking, followed by spreading the solution on MSA plates as described above. The cross-over rate from the contaminated larvae to the surface was investigated by contact Rodac™ plates with MSA (Oxoid, Wesel, Germany).

All agar plates were incubated at 35 ± 2°C for 24h and bacterial load was quantified in cfu/plate.

**Definition and analysis**

The bacterial cfu counts were $\log_{10}$ transformed to log densities (LD), and each LD was normalized to the initial bacterial LD, resulting in a log reduction value (LRV) that could then be used to compare reduction of *S. aureus* on larvae, maPK and control surfaces (mean LRVΔ). Percent reduction (PR) was evaluated by the formula $PR=1-10^{-LRV\Delta}$ [17]. A p-value of < 0.05 was considered statistically significant.

**Results**

**Microbial colonization on the larvae’s cuticle (pilot experiment)**

Investigation of the larvae’s (n=5) exoskeletal cuticle revealed no *S. aureus*. This proved both, the suitability of the model for our use and *S. aureus* as an appropriate test organism. We could ensure that all *S. aureus* detected in following assays originated from intended contamination with *S. aureus*. Furthermore four bacterial genera including both, gram-positive (coagulase-negative staphylococci, average of $1 \times 10^3$ cfu/plate; enterococci, average of $4.5 \times 10^2$ cfu/plate, *Bacillus* spp., average of $1.5 \times 10^2$ cfu/plate) and gram-negative (*Escherichia coli*, average of $4 \times 10^2$ cfu/plate) bacteria were found on the larval cuticle.

**Residual *S. aureus* burden on the larvae’s cuticle after contamination (pilot experiment)**

After contaminating the larvae (with $1.5 \times 10^8$ cfu/mL *S. aureus*) the residual burden of *S. aureus* was determined. An average number of $5.1 \times 10^4$ cfu/mL *S. aureus* was identified in the suspension assuming that this amount of microorganisms was present on the cuticle of the larvae after contaminating.

**Significant reduction of *S. aureus* burden on contaminated maPK surfaces exposed to larvae - Test set-up 1**

Each experiment was repeated six times, and maPK surfaces were contaminated with *S. aureus* and subsequently larvae were exposed to these surfaces. Bacterial load on larvae after exposure to the contaminated surfaces revealed a colony count of ~$5.1 \times 10^3$ cfu/mL (representing the contamination rate of the larvae’s cuticle) for maPK compared to a count of ~$2 \times 10^3$ cfu/mL after exposure to control surface (Table 1). Thus, a significantly higher cross-over rate of *S. aureus* from the control surface to larvae than from maPK surface to larvae was observed.
**Method** | **Surface Type** | **Recovery of *S. aureus* on:**
--- | --- | ---
 |  | larvae

*S. aureus* contamination of surface (set-up 1)

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<tr>
<th></th>
<th>maPK</th>
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<tr>
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<td>control</td>
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*S. aureus* contamination of larvae (set-up 2)

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<th>maPK</th>
<th>2.1 x 10¹ **</th>
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<td>control</td>
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**Table 1:** Comparison of bacterial load on surfaces and larvae in both set-ups after exposure.

**Note:** maPK membrane-active polycation. *Colony forming units per mL; **Colony forming units per plate

Determination of residual microbial contamination of the surface by contact plates revealed a mean LRV of 2.56 for the antimicrobial surface compared to a mean LRV of 0.83 for the control surface (Table 2). These results represent a significant microbial growth reduction of 98.1% and a difference in LRVs of 1.73 (p=0.0117) for maPK compared to control.

**Method** | **Surface Type** | **Mean LRV** | **SD** | **Mean LRVΔ** * | **Mean PR** | **p-value**
--- | --- | --- | --- | --- | --- | ---
**Set-up 1** | maPK | 2.56 | 0.378 | 1.73 | 98.1% | 0.0117 ** |
|  | control | 0.83 | 0.467 |  |  |  |
**Set-up 2** | maPK | 3.46 | 0.148 | 2.35 | 99.5% | 0.0001 ** |
|  | control | 1.11 | 0.237 |  |  |  |

**Table 2:** Summary of the antimicrobial efficacy of maPK versus control surfaces in both set-ups against *S. aureus*

**Note.** maPK membrane-active polycation, LRV log reduction value, SD standard deviation, PR percent reduction

* Presents the difference between LRVs of each experiment and was used to calculate the mean PR

**Indicating significance (p < 0.05).**
Significant reduction of pathogen cross-over from contaminated larvae to maPK surface - Test set-up 2

Transmission of the microorganism from colonized larvae to surfaces was simulated and each experiment was repeated six times.

Investigation of the surfaces after contact with contaminated larvae revealed a higher prevalence of *S. aureus* on the control surface compared to maPK (~5.1 x 10^3 versus ~2.1 x 10^1 cfu/plate), and substantiates a higher transmission of *S. aureus* from larvae to control (Table 1).

Determination of *S. aureus* on larvae after exposure to the surfaces revealed a LRV of 3.46 for larvae transferred to maPK and a LRV of 1.11 for larvae exposed to control (Table 2). Comparison of the two LRVs showed a difference of 2.35 (p=0.0001) representing a significant PR of 99.5% of the test-organism on larvae in case of exposure to maPK surface.

Discussion

We recently demonstrated in vitro antimicrobial activity of maPK surfaces against *S. aureus* ATCC 6538 [15]. This follow-up investigation confirmed the previously described activity in a *Galleria mellonella* larvae model used to mimic a more “real life” condition by applying a living organism on the surface. Here, we demonstrate that larvae exposed to the contaminated surface showed a lower rate of microbial transference in case of exposure to the antimicrobial active surface. This data was supported by a significant reduction (98.1%; p=0.0117) of the bacterial load on the antimicrobial surface after contaminating the surface with *S. aureus* and exposure of *Galleria mellonella* larvae on this surface.

The second experimental set-up demonstrated efficacy of the antimicrobial surface also when larvae were contaminated with *S. aureus* and transferred to the surface by a reduced transference of microorganisms from the larvae to the antimicrobial active surface, and a significantly reduced residual colonization (99.5%; p=0.0001) of the larvae.

To the best of our knowledge, this is the first study that investigated the biocidal efficacy of surfaces in a more practical assessment involving this model.

Since the transmission of nosocomial pathogens from hospital surfaces is evident and even potentiated by unsatisfactory adherence to infection control guidelines [18-20], the implementation of antimicrobial surfaces as an approach to minimize contamination in the hospital setting is eligible [8, 21].

Several studies have shown effectiveness and potential of self-disinfecting surfaces in in-vitro assays [22, 23]. However, when implementing these surfaces in the hospital setting, various problems became evident, which were not predictable after their in-vitro evaluation, demonstrating that the transfer of data obtained by laboratory models into real-life setting is challenging. For example, copper surfaces showed antimicrobial potency with long term effectiveness in vitro [24]. However, impact of organic soiling on the antimicrobial effect of copper and acquired copper resistance in gram-positive and in gram-negative bacteria has been reported and thus limiting its application in the hospital setting [25, 26]. Micro-patterned surfaces, designed to prevent microbial adhesion to the surface were evaluated in vitro and in the clinical setting and demonstrated inhibition of surface contamination and transfer of clinical relevant organisms in both settings [21, 27]. In this case the issue...
of cost, applicability and stability was not considered in the initial studies. Hence, translational pre-clinical in vivo trials, like the *Galleria* model, might be useful to further validate the biocidal capacity of surfaces prior to their implementation in the hospital setting [17].

We demonstrate, that maPK surfaces possess antimicrobial efficacy both, in vitro and in vivo, although surfaces displayed a stronger killing effect in vitro (LRV of 8 in vitro versus 1.73 and 2.35 in vivo). Besides demonstrating a reduction of bacterial surface contamination in the larvae model we also show a reduction of bacterial transfer from the larvae to the surface and vice-versa in the present study. These observations lead to the hypothesis that maPK surfaces might also reduce pathogen transfer from and onto surfaces via through healthcare workers’ (HCWs) hands, which is a major concern for microbial transmission in the hospital setting [18, 19].

*Galleria* larvae are mainly used as alternative model organisms to simulate systemic forms of disease and have been adapted for a broad range of bacterial and also fungal pathogens [28].

These organisms bear several advantages in comparison to other invertebrate models, i.e. they are easy to keep, cheap to purchase, can be maintained at temperatures up to 37°C and the larvae are not subject to the ethical limitations of mammalian models [28].

Limitations of the present study include that contact-plate sampling was performed with agar plates which didn’t contain neutralizing compounds, but earlier data displayed no influence. Furthermore, the experiments were performed only with *S. aureus* ATCC 6538 using one inoculum. The effectiveness against other relevant microorganisms and a broader range of inocula has to be investigated.

In addition, this study did not address the impact of potential compromising factors on biocidal activity of the maPK surface, like organic soiling or microbial resistance to the biocide [26]. The main challenge ahead is to evaluate whether our tested surface has any in vivo impact in terms of reducing HCAI.

**Conclusion**

In conclusion, the maPK surface showed antimicrobial efficacy in a larvae, in vivo model. Thus, this surface may offer a complementing tool to reduce microbial burden in the hospital setting and delay recontamination in a more sustained manner compared to periodic cleaning and disinfection alone.

**List of abbreviations**

- CBA: Columbia agar 5% sheep blood
- HCAI: Healthcare-associated infection
- HCW: Healthcare worker
- LD: Log density
- LRV: Log reduction value
- maPK: membrane-active polycation
- MSA: Mannitol salt agar
- PBS: Phosphate buffered saline
- PR: Percent reduction
- SD: Standard deviation

**Declarations**

**Ethics approval and consent to participate**

The use of *G. mellonella* does not require ethic approval.

**Consent for publication**

Not applicable.

**Availability of data and material**

All data generated or analyzed during this study are included in this published article.

**Competing interests**

The authors declare that they have no competing interests.
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Author’s contributions
IW conducted the experiments and reviewed the manuscript, UB and VN provided the larvae and reviewed the manuscript. HH provided the test surfaces and reviewed the manuscript. CLF, GH and DO reviewed the manuscript. AM analyzed the examinations, interpreted the data and drafted the manuscript. All authors read and approved the final manuscript.

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References


