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## NON-CONTACT INDUCTION HEATING AND SAAP-148 ELIMINATE PERSISTERS WITHIN MRSA BIOFILMS MIMICKING A METAL IMPLANT INFECTION

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### Abstract

Implant-associated infections are the primary cause of complications following orthopaedic surgery. Due to biofilm and persister formation, current treatments, *i.e.* surgical debridement followed by antibiotics, often fail. There is an urgent need for alternative strategies to combat such infections. Therefore, the present study investigated the effects of non-contact induction heating (NCIH), the antimicrobial peptide SAAP-148 and combinations thereof on bacterial counts in 7 d mature biofilms and in persister-enriched biofilms of methicillin-resistant *Staphylococcus aureus* (MRSA) on titanium-aluminium-niobium (TAN) discs. Enrichment of persisters was achieved by daily exposure of mature biofilms to high doses of rifampicin and ciprofloxacin for 3 consecutive days. To heat up the TAN discs, a miniaturised induction heater was built and successfully validated. Using this apparatus, NCIH resulting in surface temperatures up to 85 °C eradicated all the bacteria in immature biofilms but not in mature biofilms, whereas persisters were already eliminated at surface temperatures  $\geq 70$  °C. SAAP-148 at concentrations  $> 25.6$   $\mu\text{mol/L}$  reduced the persister counts in antibiotics-exposed, mature biofilms. As surface temperatures  $> 60$  °C can have detrimental effects on the surrounding tissues, the maximum temperature of NCIH used in combination with SAAP-148 on persisters was set to 60 °C. Results revealed that this combination was slightly more effective than the peptide or NCIH alone in eliminating biofilm-embedded persisters. NCIH and SAAP-148 can be applied both invasively and non-invasively in various treatment scenarios. Together, combinations of NCIH and SAAP-148 might be a promising treatment strategy to combat metal-implant-associated infections.

**Keywords:** Implant-associated infection, methicillin-resistant *Staphylococcus aureus*, biofilm, persisters, titanium-aluminium-niobium discs, non-contact induction heating, antimicrobial peptide, SAAP-148, prosthetic joint infection.

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List of Abbreviations			
		PBS	phosphate-buffered saline
		PEMF	pulsed electro-magnetic field
		SAAP	synthetic anti-microbial and anti-biofilm peptide
AMF	alternating magnetic fields	TAN	titanium-aluminium-niobium
BHI	brain heart infusion	TSB	tryptone soy broth
BM2	biofilm medium 2		
CFU	colony-forming unit		
DAIR	debridement-antibiotic therapy and implant retention		
DMSO	dimethyl sulphoxide		
MBC	minimal bactericidal concentration		
MRSA	methicillin-resistant <i>Staphylococcus aureus</i>		
NCIH	non-contact induction heating		

### Introduction

An implant-associated infection is a major complication of orthopaedic surgery and is responsible for implant failure. Current treatments often fail to combat these

persistent infections, which results in high morbidity and mortality rates, and a major economic burden (Kapadia *et al.*, 2014; Springer *et al.*, 2017). Infections of (prosthetic) implants are predominantly caused by *Staphylococci*, such as MRSA, that form a biofilm on the implant surface (Benito *et al.*, 2016; Pulido *et al.*, 2008). The extracellular polymeric matrix of the biofilm protects the biofilm-embedded bacteria from the actions of antibiotics, effector cells and molecules of the immune system (Achermann *et al.*, 2014). Also, the various bacterial subpopulations within the biofilm show different metabolic activities, which alter their response to antibiotics, *i.e.* subpopulations with low metabolic activity tolerate antibiotics and will therefore persist within the biofilm (Bjarnsholt *et al.*, 2011). Upon discontinuation of the antibiotic treatment, these persisting bacteria can cause infection relapse (Harms *et al.*, 2016), thus explaining the failure of the antibiotic treatment (Spoering *et al.*, 2001). Other treatments that are used to remove implant-associated infections include DAIR and revision surgery (McConoughey *et al.*, 2014; Vaz *et al.*, 2020). Unfortunately, these treatments are invasive, often not successful and associated with additional complications.

Novel non-invasive strategies, such as heating up metal implants by using high-frequency AMF (also known as induction heating), have been proposed to reduce or even eradicate 24 h bacterial biofilms (Chopra *et al.*, 2017; Pijls *et al.*, 2017). The principle of AMF to generate heat in metal objects has been used on several biomedical technologies as it specifically heats up metals and does not directly harm the surrounding tissue (Christiansen *et al.*, 2017). Recently, the effect of heat generation by AMF has been tested in an *in vitro* model of *Pseudomonas aeruginosa* and MRSA biofilms on metal implant mimics (Chopra *et al.*, 2017). Exposure for longer than 3 min to ~80 °C results in elimination of bacteria within biofilms and increased sensitivity of bacteria to antibiotics. Similarly, heating up titanium alloys by NCIH results in a temperature of 80 °C after 3.5 min and almost total eradication of the bacterial load of MRSA 24 h biofilms (Pijls *et al.*, 2020). Moreover, a synergistic effect could be observed upon combination of 60 °C NCIH and antibiotics, resulting in a significant elimination of viable bacteria within biofilms (Pijls *et al.*, 2020). Another alternative strategy for implant-associated infections is the use of antimicrobial peptides such as SAAP (Riool *et al.*, 2017). SAAPs were developed based on the blueprint of the human antimicrobial peptide LL-37 and exert a lower rate of resistance development due to their multifunctional mechanisms of action (Fjell *et al.*, 2011; Scott *et al.*, 2002). Notably, SAAP-148 has shown promising anti-biofilm activity by totally eradicating MRSA biofilms from polystyrene surfaces, wounded *ex vivo* human skin and superficially wounded murine skin (de Breij *et al.*, 2018). Moreover, SAAP-148 is more effective than several other antimicrobial peptides in eliminating bacteria, including persisters,

in mature MRSA biofilms on various surfaces, including TAN discs (Scheper *et al.*, 2021). The aim of the present study was to assess the effect of NCIH and NCIH in combination with SAAP-148 on bacteria in biofilms on metal surfaces mimicking a metal implant-associated infection. For this purpose, persisters in 7 d mature biofilms were obtained by high-dose antibiotic treatment and subsequently exposed to NCIH, SAAP-148 and combinations thereof. As the TAN discs were designed to fit in the wells of a 96-well plate, a custom-built NCIH device was developed to heat up the TAN discs. The combination of NCIH and SAAP-148 was expected to combat the biofilm-embedded persisters with a high efficacy by attacking the biofilm from both the basal (implant) and top (peri-implant) side.

## Materials and Methods

### Antibiotics and SAAP-148 peptide

Ciprofloxacin (Sigma-Aldrich, 86393-32-0) was diluted in MilliQ water (5.12 mg/mL) and rifampicin (Sigma-Aldrich, 13292-46-1) in DMSO (4 mg/mL) and stored at -20 °C until use. SAAP-148 peptide (sequence LKR<sup>W</sup>VWKR<sup>V</sup>FKLLKRYWRQLKKPVR) was synthesised by solid phase chemistry using an automated multiple peptide synthesiser (SyroII, MultiSyntech, Witten, Germany) as described elsewhere (de Breij *et al.*, 2018). The molecular mass of the N-terminal acetylated and C-terminal amidated peptide was confirmed by mass spectrometry and the purity of the peptide exceeded 95 %, as determined by reverse phase high-performance liquid chromatography and detection at 214 nm. The lyophilised peptides were stored at -20 °C until use.

### TAN discs

Medical grade titanium-7 % aluminium-6 % niobium (TAN; iso5832/11) discs (diameter 5 mm, height 1.5 mm) with a handle allowing gentle removal of the discs from the wells made specifically to fit in 96-well plates were a kind gift of Dr TF Moriarty (AO Research Institute, Davos, Switzerland). For re-use, TAN discs were rinsed and submerged overnight in 70 % EtOH, after which they were air-dried and autoclaved.

### NCIH

TAN discs were exposed to a PEMF of 101 kHz at a maximum of 135 W from a custom-built induction heater (Fig. 1a). The induction heater featured a solenoid type coil with 10 turns of 6 mm copper tubing and it was designed to accommodate a 50 mL plastic centrifuge tube. This tube was equipped with a silicon mould in the centre to fit a 2 mL micro-tube with the TAN disc inside. This set-up was designed to consistently place the TAN disc in the centre of the PEMF generated by the solenoid coil, therefore allowing temperature control by adjusting the heating time when the association between time and

temperature was known. The relationship between time and temperature was determined in triplicate using two fibre-optic temperature sensors connected to a fibre-optic signal conditioner (Neoptix, ReFlex-4; Neoptix Inc, Quebec City Canada; Fig. 1b). Fibre-optic temperature sensors were not affected nor heated by the PEMF, therefore ensuring reliable temperature measurements in the presence of a PEMF.

### Culture of MRSA

Frozen stocks of MRSA LUH14616 (sequence type 247, NCCB 100829, kindly provided by Dr S Croe, Department of Medical Microbiology, Maastricht University Medical Centre, Maastricht, the Netherlands) in 20 % glycerol (Thermo Fisher Diagnostic BV, 1200.36.001) were thawed to spread on Trypcase Soy Agar + 5 % sheep blood plates (Biomérieux, Marcy-l'Étoile, France, 43009) and cultured overnight at 37 °C. Subsequently, several colonies were taken from these plates and cultured in TSB medium (Oxoid, Basingstoke, Hampshire, UK, CM0129) for 2.5 h at 200 rpm and 37 °C. The mid-logarithmic phase bacteria were harvested by centrifugation (1,000 ×g for 10 min), washed with PBS (pH 7.4) and then, based on the optical density at 600 nm, diluted to  $1 \times 10^7$  CFU/mL in the desired medium.

### MRSA biofilm and persister formation

MRSA was diluted in BHI broth (Oxoid, CM1135) for 7 d biofilms and in BM2 medium [62 mmol/L potassium phosphate buffer (pH 7) with 7 mmol/L  $(\text{NH}_4)_2\text{SO}_4$ , 200 mmol/L  $\text{MgSO}_4$ , 10 mmol/L  $\text{FeSO}_4$ , 40 % glucose + 50 % casamino acids] for 24 h biofilms. Briefly, 100  $\mu\text{L}$  of bacterial suspension was pipetted into 96-well plates containing TAN discs. After 24 h or 7 d, the wells were washed and the discs with biofilms were gently transferred to polystyrene 96-well plates or polypropylene 96-well plates for exposure to antibiotics or SAAP-148, respectively. Plates with 24 h biofilms were sealed with non-breathable plastic film sealers (AmpliStar Adhesive Plate Sealers, Westburg, Leusden, the Netherlands, WB 2-3830). Mature biofilms on plates were sealed with breathable rayon film sealers (VWR, 391-1262). For enrichment in persisters, mature biofilms were washed daily with PBS and then 100  $\mu\text{L}$  of BHI suspension with 10× the MBC of ciprofloxacin (1,280  $\mu\text{g}/\text{mL}$ ) and rifampicin (10  $\mu\text{g}/\text{mL}$ ) were added. After 24 h, the antibiotics were refreshed daily for up to 5 d. Controls received similar concentrations of BHI suspension in MilliQ water. Biofilms were allowed to form in a plastic box at 37 °C, to maintain a humidified environment. Medium controls were used to monitor any possible contamination.

### Effect of treatments on bacterial counts in biofilms

Biofilms were exposed to NCIH at various target temperatures (60 to 85 °C), to SAAP-148 peptide or to combinations of NCIH and SAAP-148 at various dilutions in PBS for 2 h. For these combinations,

biofilms were first exposed to the peptide and subsequently to NCIH. Prior to treatment, planktonic bacteria were removed from the wells by 2 washes with PBS. After treatment, TAN discs with biofilms were transferred to 96-well plates containing 100  $\mu\text{L}$  of PBS and sonicated (10 min at 40 kHz). Serial dilutions of bacterial suspensions were plated on Mueller Hinton agar plates (Oxoid, CM0337) and CFU was counted after incubation overnight (37 °C).

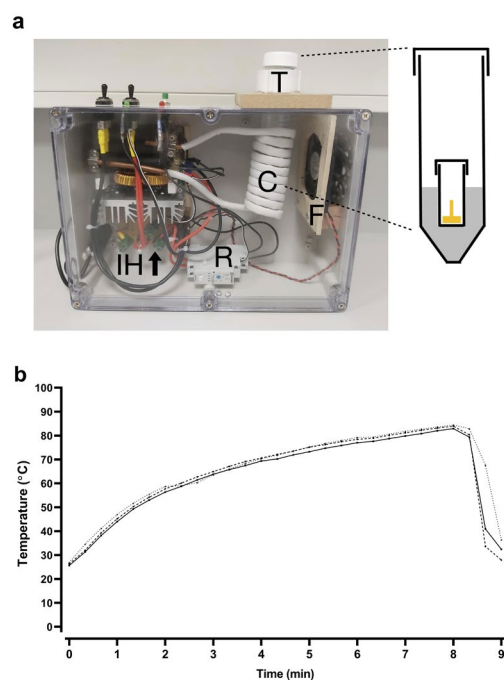
### Data analysis

Kruskal-Wallis tests followed by Mann-Whitney U tests (GraphPad Software 9) were performed to determine the significance of the difference in CFUs between treated and control samples.  $p < 0.05$  was considered statistically significant.

## Results

### Calibration of the induction heating equipment

The temperature at the surface of TAN discs was measured every 1 s during a period of 8 min of



**Fig. 1. The relationship between duration of induction heating by the custom-built NCIH equipment and the surface temperature of TAN discs.** (a) The custom-built induction heater and a schematic drawing of the orange TAN disc in the 2 mL micro-tube, placed in a silicon mould (gray) in the centre of the 50 mL plastic centrifuge tube (IH = induction heater circuit; C = coil to generate a pulsed electromagnetic field; T = docking station for 50 mL tube; F = fan for convection cooling of coil and induction heater circuit; R = time release to control heating time). (b) Fibre-optic measurements were performed to determine the relationship between the duration of NCIH and the surface temperature of the TAN disc in PBS. Results are means of duplicates from 3 independent runs plotted every 20 s for 9 min.

NCIH using fibre-optic sensors to establish the relationship between the duration of NCIH and the surface temperature of TAN discs in PBS (Fig. 1b). Results revealed that within the first 2 min the surface temperature rapidly increased from 27 °C to 60 °C. Thereafter, the surface temperature linearly increased by 10 °C every 2 min until 80 °C and, subsequently, increased up to 85 °C during the 2 min interval. After transfer of the TAN discs to wells containing PBS at 37 °C, the temperature of the discs dropped to ambient temperature within 1 min. Based on these data, a time relay was used to control the heating time. The time for induction heating of TAN discs in the presence of 100 µL of PBS were set at 2 min for 60 °C, 4 min for 70 °C, 6 min for 80 °C and 8 min for 85 °C.

#### Effect of heat-treatment duration on the bacterial counts in 7 d mature biofilms

To determine if reaching the temperature rather than the duration of the exposure to that temperature killed the bacteria within the biofilms, the number of viable bacteria within 7 d mature biofilms on TAN discs was determined upon exposure to a range of temperatures (60, 70 or 80 °C) using a heat-block for various durations. Results revealed a heat-dependent reduction in biofilm-embedded bacteria, with a maximal reduction down to 500 CFU/mL after exposure to 80 °C. Longer exposure of biofilms (4 min compared to 2 min) to the heat-block at 60 °C did not show a further reduction in bacterial load. Further, subjecting the biofilms for 4 min at 70 and 80 °C exerted a maximal, negligible 10-fold reduction in CFU/mL compared to 2 min exposure to these temperatures. Data indicated that longer durations at a selected temperature resulted in a steady-state

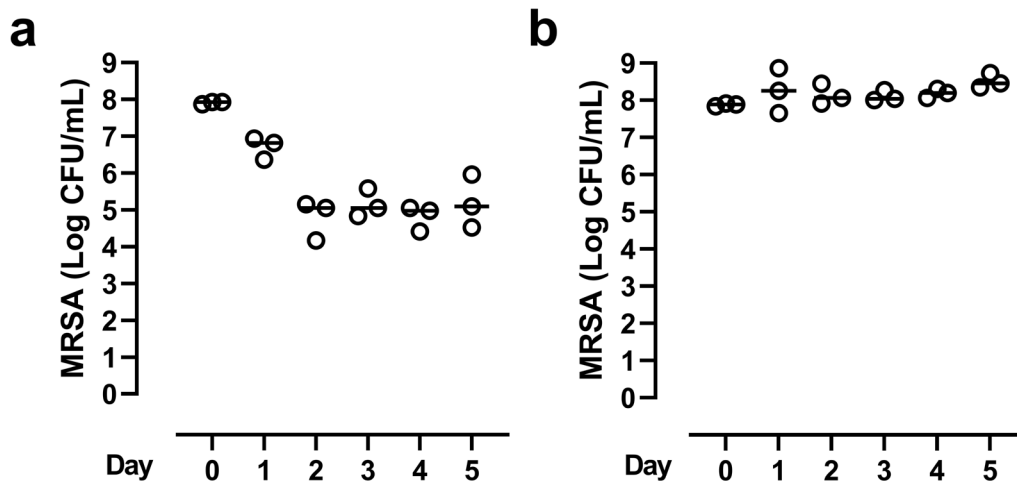
antibacterial effect rather than a further reduction in bacterial counts in the biofilms.

#### Development of the *in vitro* model for persisters in biofilms

To develop an *in vitro* model that partially mimicked metal implant-associated infections, the effects of daily exposure to high doses (10× MBC) of rifampicin and ciprofloxacin on bacteria residing within 7 d mature MRSA biofilms on TAN discs for a period of 5 d were determined. Results revealed that the number of viable bacteria in the biofilms rapidly dropped ( $p < 0.05$ ) from approximately  $1 \times 10^8$  CFU/mL to about  $1 \times 10^5$  CFU/mL during the first 2 d of antibiotic exposure and thereafter the bacteria tolerated the antibiotics for at least an additional 3 d (Fig. 2a). The numbers of viable bacteria in control biofilms did not change ( $p > 0.05$ ) during these 5 d (Fig. 2b). Based on these data, 3 d of antibiotic treatment has been used in further experiments to study the effects of innovative treatments on persisters in biofilms.

#### Effect of induction heating on bacteria in biofilms and persisters

To allow for a comparison with previously published studies on NCIH with 24 h biofilms, the effects of NCIH on bacteria in 24 h MRSA biofilms on TAN discs were first determined after various durations of induction heating. Results revealed a surface temperature-dependent reduction ( $p < 0.05$ ) in bacterial counts, with complete eradication of the bacteria seen at a surface temperature of 85 °C (Fig. 3a). However, a maximum 5-Log reduction in bacterial counts was seen for MRSA in 7 d mature biofilms on TAN discs after heating the surface



**Fig. 2.** Effect of high-dose antibiotics on bacteria in mature biofilms. 7 d mature *Staphylococcus aureus* LUH14616 (MRSA) biofilms on TAN discs were exposed daily for up to 5 consecutive days to a combination of (a) 10× MBC rifampicin and ciprofloxacin or (b) the diluent of the antibiotics as control. After each day, TAN discs with biofilms were transferred to wells containing PBS, sonicated and thereafter the number of viable bacteria was assessed microbiologically by counting the CFU. Results are from 3 independent experiments each performed in triplicate; the solid line indicates the median Log CFU/mL. Significance of the differences between the various surface temperatures was calculated by Kruskal-Wallis test (a:  $p < 0.05$ , b:  $p > 0.05$ ).

to  $\geq 70$  °C (Fig. 3b), indicating that NCIH was not sufficient to eradicate all biofilm-encased bacteria. Regarding the persister model, NCIH of the TAN discs to 60 °C led to a 10-fold reduction in the number of persisters (Fig. 3c), whereas heating to surface temperatures  $\geq 70$  °C resulted in total eradication of persisters within the biofilms ( $p < 0.001$ ).

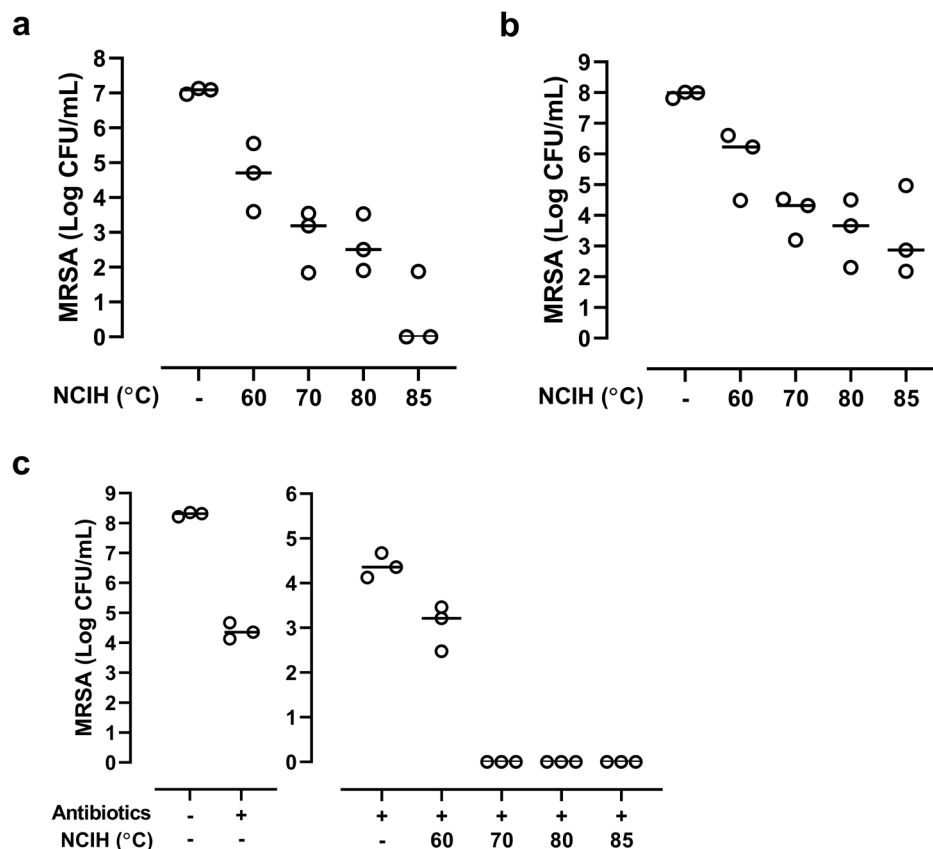
#### Effect of SAAP-148 monotherapy and in combination with induction heating on persisters within biofilms

Since persisters are the main cause of therapy failure in implant-associated infections, the effect of SAAP-148 on persisters in antibiotics-exposed mature biofilms on TAN discs was first confirmed. Results revealed a dose-dependent reduction ( $p < 0.05$ ) in bacterial counts for persisters in a biofilm by peptide concentrations  $> 12.8$   $\mu\text{mol/L}$ , with almost all persisters eradicated at the highest peptide concentration (51.2  $\mu\text{mol/L}$ ) (Fig. 4). Next, the effect of SAAP-148 followed by induction heating to a surface temperature of 60 °C on persisters in biofilms was determined. Results showed that this combination was slightly more effective in reducing persisters in antibiotics-exposed, mature biofilms than the peptide

or NCIH alone. Mann-Whitney U test indicated that the combination of NCIH with 12.8  $\mu\text{mol/L}$  SAAP-148 was significantly ( $p < 0.05$ ) more effective than the peptide or induction heating alone (Fig 4).

#### Discussion

Two innovative, non-antibiotic strategies to combat metal implant-associated infections with MRSA were investigated using a novel *in vitro* model (Scheper *et al.*, 2021). For this model, 7 d mature biofilms on TAN discs were developed and exposed for 3 d to high doses of rifampicin and ciprofloxacin, an antibiotic combination that is commonly used to treat implant-associated infections caused by *Staphylococcus aureus*/MRSA. Control experiments indicated that the surviving bacteria indeed developed tolerance to the subsequent antibiotic treatment. First, the effects of NCIH and of the novel antimicrobial peptide SAAP-148 on bacterial subpopulations including persisters in biofilms were determined. To circumvent undesirable side-effects of these treatments on peri-



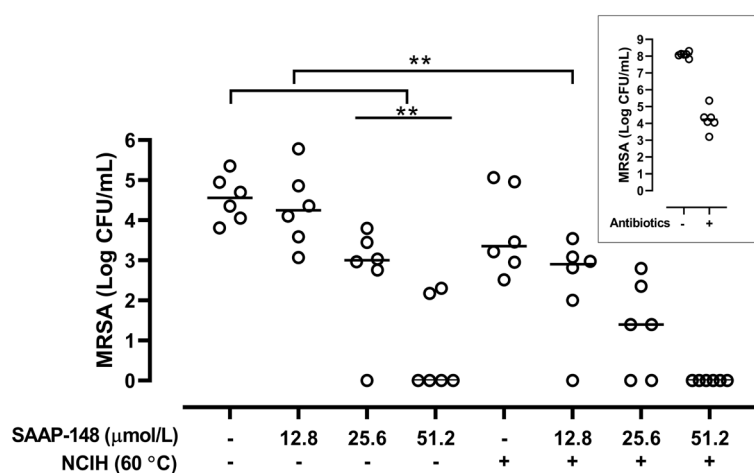
**Fig. 3.** Effect of surface temperature on bacteria, including persisters, residing in biofilms. (a) 24 h *Staphylococcus aureus* LUH14616 (MRSA) biofilms, (b) 7 d mature MRSA biofilms and (c) 7 d mature, 3 d antibiotics ( $10\times$  MBC of rifampicin and ciprofloxacin)-exposed MRSA biofilms on TAN discs were transferred to vials containing PBS for surface heating by NCIH for various intervals. Thereafter, the TAN discs with biofilms were transferred to wells containing PBS and subjected to sonication for microbiological determination of the number of viable bacteria. Results are from 3 independent experiments each performed in quadruplicate; the solid line indicates the median Log CFU/mL. Significance of the differences between the various surface temperatures was calculated by Kruskal-Wallis test (a,c:  $p < 0.0001$ , b:  $p < 0.05$ ).

implant tissues, the second strategy focused on establishing the possible additive effects of combining NCIH with SAAP-148 on the counts of persisters in biofilms.

Results showed a temperature-dependent reduction in bacterial counts in immature biofilms with complete bacterial eradication seen at a surface temperature of 85 °C, whereas bacterial counts in mature biofilms decreased in a temperature-dependent fashion up to 70 °C and thereafter remained unchanged. The observed effect of NCIH on immature biofilms is in line with an earlier observation that almost all bacteria within 24 h biofilms are eliminated by a heat shock of 85 °C (Pijls *et al.*, 2020). The difference in susceptibility of bacteria in mature and immature biofilms to induction heating could be related to the amount of protective extracellular matrix, which is considerably larger in mature biofilms. Further, the present study focused predominantly on (persister-enriched) mature biofilms since others have shown that 24 h biofilms may not be optimal to study the effects of novel strategies and drugs on bacteria in mature biofilms (Dice *et al.*, 2009). In agreement, Cazander *et al.* (2010) reported that at least 6-8 d are required to develop the mature biofilm profile that is observed in clinical implant-associated infections. Since the persister subpopulation is the main cause of treatment failure in metal implant-associated infections, the findings that NCIH resulting in temperatures  $\geq 70$  °C completely eradicated all bacteria in persister-enriched mature biofilms are of particular importance. The difference in the efficacy of NCIH on bacteria in mature biofilms and persisters within biofilms may be related to the bacterial numbers in these models; the number of persisters being 10,000-fold lower than the number

of bacteria in mature biofilms. As reported earlier (Scheper *et al.*, 2021), SAAP-148 peptide at the highest non-haemolytic concentration (de Breij *et al.*, 2018) effectively reduces persister counts in antibiotics-exposed, mature biofilms, but does not eradicate all persisters.

Both NCIH and SAAP-148 should not result in adverse effects, such as cytotoxicity. In this context, temperatures  $> 60$  °C were considered undesirable due to the risk of transmission, to some extent, of heat to the peri-implant tissue. This is based on the observation that induction heating of a titanium memory rod up to 60 °C in rats does not cause necrosis (Müller *et al.*, 2014). Chopra *et al.* (2017) reported that *in vivo* AMF exposure reaching temperatures up to 100 °C results in (irreversible) damage to tissues closely surrounding the metal implant. The present study revealed that exposure of antibiotics-exposed, mature biofilms to SAAP-148 for 2 h followed by induction heating to 60 °C was slightly more effective than the heat shock or the peptide alone, with the highest concentration of the peptide in combination with induction heating completely eradicating all persisters in biofilms. Of note, the effect of the combination did not depend on the order in which the peptide and induction heating were applied (results not shown). A possible explanation for this enhanced effect of NCIH in combination with SAAP-148 on persisters in mature biofilms could be that the treatments attacked the bacteria in the biofilms from different sides, *i.e.* NCIH from the basal side (implant) and the peptide from the top (peri-implant). In addition, the modes of action of these treatments also differ completely. Based on these findings, it was concluded that combinations of NCIH and SAAP-148 could become a promising



**Fig. 4. Effect of SAAP-148 in combination with NCIH on persisters within MRSA biofilms.** 7 d mature *Staphylococcus aureus* LUH14616 (MRSA) biofilms on TAN discs were exposed daily to high-dose antibiotics (10 $\times$  MBC of rifampicin and ciprofloxacin) and subjected for 2 h to increasing concentrations of SAAP-148 with or without subsequent NCIH to 60 °C. Insert indicates the effect of antibiotics on bacteria in the mature biofilm. All biofilms were subjected to sonication for microbiological determination of the number of viable bacteria. Results are from 6 independent experiments each performed in quadruplicate with a solid line indicating the median Log CFU/mL. Significance of the differences between the various conditions was calculated by Kruskal-Wallis ( $p < 0.0001$ ) and subsequently Mann-Whitney U tests. \*\*  $p < 0.01$ .

strategy for the treatment of metal implant-associated infections.

To heat up the TAN discs in a controlled fashion, a miniaturised version of the NCIH apparatus was built and successfully validated. This equipment was designed to reach a target temperature rather than to maintain the temperature for a prolonged period, as was the case in previous NCIH studies (Pijls *et al.*, 2020). The rationale for this approach was twofold. First, a closed feedback temperature control system for non-invasive heating at 60 °C may be difficult to achieve clinically (Cheng *et al.*, 2018), indicating that the NCIH model featured here represents a practical approach. Secondly, temperature rather than duration of that temperature may be more effective against bacteria. In agreement, experiments investigating the effects of duration of exposure to a temperature using a heat block revealed that longer durations at a selected temperature resulted in a steady-state antibacterial effect rather than a further reduction in bacterial counts in biofilms. In line with this, recent studies on thermal stability of proteins among different species (*e.g.* human, bacterial and fungi) have suggested that stability of key proteins of these pathogens is affected by lower temperatures than that of human proteins (Leuenberger *et al.*, 2017; O'Toole *et al.*, 2015). The future medical application of NCIH could be divided into non-invasive and invasive use. During the non-invasive use, the metal implant is heated non-invasively through the skin, while great care is taken to avoid excessive heating to areas where the implant is fixed to the bone or areas that are in close contact with crucial anatomical structures such as nerves and major blood vessels. This may be performed in a segmental way with tailored heating protocols for each segment: segmental induction heating (Pijls *et al.*, 2018). In the future, the non-invasive application of induction may first help patients for whom surgical treatment is not possible and who, for instance, receive suppression antibiotic therapy. NCIH will most likely be part of a multi-modality treatment that also involves antibiotics and adjuvants (non-antibiotic chemical agents). Alternatively, NCIH can be used invasively, *e.g.* during surgery, to increase the effectiveness of the surgical procedure such as in DAIR: NCIH kills micro-organisms by heat and it could be used to heat the implant and parts of the implant that cannot be reached and mechanically cleaned. The soft tissue can be kept away from the heated part of the implant to protect it from thermal damage. Furthermore, surgery allows for direct temperature control by direct temperature measurement (*e.g.* thermocouple) or infrared thermal imaging. However, it should be mentioned that the current *in vitro* model for metal implant-associated infections suffers from some limitations. First, all experiments were performed with only a single strain (although a clinical isolate) of MRSA (de Brij *et al.*, 2018). Another major limitation is the absence of peri-implant tissue and local factors, such as exudate or wound fluid. These limitations may

be circumvented in future experiments by using TAN discs in 12-well plates containing cultures of relevant cell types, such as myeloid-derived suppressive cells (Bosch *et al.*, 2020), inflammatory macrophages and wound fluid collected from patients undergoing prosthetic joint or hip replacement surgery. While the current experiments have focused on the effects of the combination treatment on persisters in mature biofilms, the effects of this combination on bacteria in mature biofilms remain unexplored. Lastly, it is unknown whether bacteria in mature biofilms can become tolerant to induction heating in the unlikely event of receiving multiple heat shocks as a treatment.

## Conclusion

Neither NCIH nor SAAP-148 peptide at clinically desirable conditions could eradicate all MRSA subpopulations residing within antibiotics-exposed, mature biofilms on TAN discs. Combinations of SAAP-148 and induction heating to 60 °C were slightly more effective than these approaches alone. Of note, the combination of the highest non-haemolytic concentration of SAAP-148 and induction heating to 60 °C resulted in complete eradication of all bacterial subpopulations in MRSA biofilms. Furthermore, NCIH and the peptide can be applied both invasively and non-invasively. Therefore, combinations of NCIH and SAAP-148 may be useful as a treatment for patients with implant-associated infections in whom the current treatment has failed. Moreover, such combinations could also be applied at an earlier stage of the treatment, *e.g.* during debridement of the implant, or even when replacing the current treatment involving debridement and antibiotics.

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One of the authors (BGP) is listed as a co-inventor on a provisional patent application from the Leiden University Medical Centre (WO2020/067898). The Leiden University Medical Centre is also holder of patent number WO2015088344 (co-inventors JWD and PHN) relating to the peptide described in the present study.

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### Discussion with Reviewer

**Jean-Paul Pirnay:** Do you plan to perform animal studies in the near future? If so, which model would you consider?

**Authors:** *In vitro* data indicated that SAAPs and NCIH are highly promising strategies to eliminate implant-associated biofilm infections. Clearly, the next step will be to study the efficacy of these strategies in animal models. SAAP-145 injected alongside infected subcutaneously inserted titanium implants is effective in reducing bacterial counts on the implants, but not in the peri-implant tissue (Riool *et al.*, 2017). In addition, bacterial colonisation of titanium implants coated with this peptide incorporated in a polymer lipid encapsulation is considerably less than that of uncoated implants, whereas the bacterial counts in the peri-implant tissue do not differ between the coated and uncoated implants (Riool *et al.*, 2017). Obviously, these studies suffer from several limitations, including the biofilms in these models being immature. The efficacy of non-contact induction heating has not yet been studied *in vivo* and/or in animal models. However, we have recently obtained a grant to study the efficacy of NCIH and other strategies to combat implant-associated infections in animal models. To address this aim, we will develop an animal model for metal

implant-associated mature biofilm infections or adapt current animal models (Zimmerli *et al.*, 1982, additional reference).

**Jean-Paul Pirnay:** Are you developing a device suitable for human therapy?

**Authors:** It is our intention to translate this technology from *in vitro* research to human therapy that can be used clinically. In the near future, we envision a device that heats the infected metal implant during a surgical debridement procedure, killing and weakening the micro-organisms in the biofilm with induction heat, so that it adds to the present standard of care. Presently, patients with infected implants will undergo long and extensive treatments consisting of multiple surgical procedures and antibiotic courses for several months (Tande and Patel, 2014, additional reference). These treatments often involve extensive revision procedures, which may not be possible in patients with high comorbidity factors. Furthermore, increasing antibiotic resistance of bacteria raises concern and limits choices of antibiotics. Therefore, it is very important that new treatments are developed to cure biofilm-infected implants (Anguita-Alonso *et al.*, 2005; Ravi *et al.*, 2015, additional references).

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**Editor's note:** The Guest Editor responsible for this paper was Henny Van der Mei.