Immunomimetic Designer Cells Protect Mice from MRSA Infection

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Graphical Abstract

Highlights

- A closed-loop gene network with bacterial sense-and-destroy actuation
- Direct diagnosis of implant-associated infections through blood biomarkers
- Early prevention of MRSA infection, as well as biofilm formation, in vivo
- Curing acute MRSA infections as an alternative to antibiotic therapy

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In Brief

Encapsulated mammalian cells with a synthetic gene circuit to sense and respond to MRSA infection could provide potential prophylactic, diagnostic, or therapeutic options for medical implant-associated infections.

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Immunomimetic Designer Cells Protect Mice from MRSA Infection

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SUMMARY

Many community- and hospital-acquired bacterial infections are caused by antibiotic-resistant pathogens. Methicillin-resistant Staphylococcus aureus (MRSA) predisposes humans to invasive infections that are difficult to eradicate. We designed a closed-loop gene network programming mammalian cells to autonomously detect and eliminate bacterial infections. The genetic circuit contains human Toll-like receptors as the bacterial sensor and a synthetic promoter driving reversible and adjustable expression of lysostaphin, a bacteriolytic enzyme highly lethal to S. aureus. Immunomimetic designer cells harboring this genetic circuit exhibited fast and robust sense-and-destroy kinetics against live staphylococci. When tested in a foreign-body infection model in mice, microencapsulated cell implants prevented planktonic MRSA infection and reduced MRSA biofilm formation by 91%. Notably, this system achieved a 100% cure rate of acute MRSA infections, whereas conventional vancomycin treatment failed. These results suggest that immunomimetic designer cells could offer a therapeutic approach for early detection, prevention, and cure of pathogenic infections in the post-antibiotic era.

INTRODUCTION

Infectious disease is the leading cause of mortality and morbidity worldwide, claiming about 15 million lives each year (Fauci and Morens, 2012). Staphylococcus aureus is one of the most frequent pathogens, accounting for 11.6 million outpatient and emergency room visits and nearly 500,000 hospitalizations per year in the United States alone (Klein et al., 2007; McCaig et al., 2006). This organism causes a variety of human diseases, ranging from skin and soft tissue infections to life-threatening infections, such as bacteremia, pneumonia, osteomyelitis, endocarditis, meningitis, and sepsis (Tong et al., 2015). The high morbidity associated with S. aureus has further increased with the rising prevalence of strains that exhibit broad antibiotic resistance, such as methicillin-resistant S. aureus (MRSA) (Dantes et al., 2013), which causes more deaths annually (~19,000) than any other single infectious agent in the United States; indeed, the number of deaths caused by MRSA exceeds that associated with HIV/AIDS, hepatitis, and influenza combined (Boucher and Corey, 2008). As these staphylococcal strains are highly virulent and are increasingly becoming resistant to every clinically available antibiotic (Stryjewski and Corey, 2014), alternative therapies are urgently needed.

One particularly important unmet medical need for anti-S. aureus therapies is to treat implant-associated infections (IAIs) (Darouiche, 2004). IAIs account for half of the 2 million cases of nosocomial infections that occur each year in the United States (Darouiche, 2004) and are one of the most feared and difficult-to-treat medical complications, causing high morbidity and mortality, and leading to substantial healthcare costs (Kapadia et al., 2016). S. aureus is the leading cause of IAI and is particularly adept at infecting foreign bodies within the human host (Del Pozo and Patel, 2009). This organism is able to persist on implant surfaces, forming biofilms, which are sessile communities of microcolonies encased in an extracellular matrix that adheres to biomedical implants (Bjarnsholt et al., 2013). Infections associated with biofilms are difficult to treat due to the presence of biomaterials that can reduce the inoculum of S. aureus required to establish an infection by a factor of more than 100,000 (Puhto et al., 2014), and it is estimated that sessile bacteria in biofilms are over 1,000-fold less sensitive to antibiotics than their planktonic counterparts (Sutherland, 2001). Therefore, most implants that are infected by S. aureus have to be surgically removed to achieve a definite cure, leading to a poor patient outcome and considerable economic burden (Darouiche, 2004).
Human innate immune response is the first line of defense against infectious microbes (Akira et al., 2006). Early recognition of S. aureus is initiated by pattern recognition receptors (PRRs) on epithelial cells and innate phagocytic cells (Fournier and Philpott, 2005). Toll-like receptor 2 (TLR2) has emerged as the most important of these PRRs in detecting extracellular S. aureus (Fournier and Philpott, 2005). TLR2 recognizes lipoproteins, lipoteichoic acid, and peptidoglycan embedded in the staphylococcal cell envelope by forming heterodimers with TLR1 (Jin et al., 2007) or TLR6 (Kang et al., 2009), and the pathogen recognition is facilitated by a CD14 co-receptor (Nilsen et al., 2008). Upon stimulation, TLR2 and TLR1 or TLR6 initiate downstream signaling events that lead to the translocation of nuclear factor κB (NF-κB) and the production of proinflammatory cytokines and chemokines that recruit phagocytes to the site of infection for the disposal of pathogens (Akira et al., 2006). However, S. aureus is a well-adapted pathogen that has evolved many mechanisms for thwarting the human immune response, ranging from blocking neutrophil chemotaxis, lysing leukocytes, and avoiding phagocytosis to resisting phagocytic killing and surviving within host cells (Foster et al., 2014). In this study, instead of using the detect-defect-destroy policy employed by the innate immunity, we apply a direct sense-and-destroy strategy based on engineering of a synthetic genetic circuit that expresses lysostaphin under the regulation of human TLR2, TLR1, TLR6, and CD14. Lysostaphin is a bacteriocin that kills most known staphylococcal species (von Eff et al., 2003). It is an endopeptidase that enzymatically cleaves the specific cross-linking polyglycine bridges in the cell walls of staphylococci (Schindler and Schuhardt, 1964). The bacterialcidal efficacy of lysostaphin was reported to be higher than those of human native antimicrobials and broad-spectrum antibiotics, including penicillin (Schaaffner et al., 1967), oxacillin (Kiri et al., 2002), and vancomycin (Plancencia et al., 2009). It is also effective against biofilms (Kokai-Kun et al., 2009; Hertlein et al., 2014) and has been widely tested in various animal models (Dajcs et al., 2000; Hertlein et al., 2014; Kokai-Kun et al., 2003, 2007; Patron et al., 1999) and in humans (Davies et al., 2010; Harris et al., 1967; Stark et al., 1974).

Here, we show that this synthetic gene network can be implemented into a variety of rodent and human cells, providing an autonomous, self-regulated staphylococcal sensing-and-eliminating function. We also show that in a foreign-body infection mouse model, for which therapeutic options are extremely limited, this closed-loop synthetic gene network operating inside microencapsulated designer-cell implants could effectively protect mice from biofilm-forming MRSA infection and cure acute MRSA infection.

RESULTS

Design of a Synthetic Gene Circuit for Infectious Microbial Sensing

Infectious Microbial Sensing Gene Network (Infect<sub>sen</sub>) consists of a synthetic bacteria-responsive signaling cascade in which the constitutive co-expression of human TLR2 (hTLR2; pYL30, P<sub>hCMV</sub>HTLR2-pA), human TLR1 (hTLR1; pYL126, P<sub>hCMV</sub>HTLR1-pA), human TLR6 (hTLR6; pYL25, P<sub>hCMV</sub>HTLR6-pA), and human CD14 (hCD14; pYL25, P<sub>hCMV</sub>HCD14-pA) senses extracellular cell-wall components of gram-positive bacteria and triggers a MYD88-dependent signaling pathway that results in activation and translocation of AP-1 and NF-κB into the nucleus (Akira and Takeda, 2004). By connecting the signaling activation of hTLR2, hTLR1, hTLR6, and hCD14 to a carefully tuned chimeric promoter P<sub>AP-1 × 5-IFN-κB × 5-NF-κB</sub> containing AP-1 and NF-κB response elements, the presence of extracellular microbial components could be directly linked to the expression of a specific target gene (Figure 1A). Since human embryonic kidney 293 cells (HEK293) cells already express hTLR1 and hTLR6 endogenously (Godefroy et al., 2014), we characterized the microbial sensor device in HEK293 cells without ectopic expression of hTLR1 and hTLR6. To optimize the gene network, we first screened for synthetic NF-κB-responsive promoters with different NF-κB-binding sites and minimal promoters (Figure S1A). We then validated the signaling activation through AP-1 by incorporating AP-1 binding elements (Figure S1B). Subsequently, to obtain the best overall transgene induction, we compared a series of promoters containing different combinations of AP-1/NF-κB-binding sequences upstream of several minimal promoter variants (Figure S1C). We also showed that system activation is greatly influenced by the expression strength of hTLR2 driven by different constitutive promoters (Figure S1D). In summary, the combinatorial assembly of 5 repeats of AP-1 and NF-κB followed by an IFNβ minimal promoter (pYL3, P<sub>AP-1 × 5-IFN-κB × 5-NF-κB</sub>) yielded the greatest signal transduction in response to stimulation of pYL30 (P<sub>hCMV</sub>HTLR2-pA), endogenous hTLR1 and 6, and pYL25 (P<sub>hCMV</sub>HCD14-pA) (Infect<sub>sen</sub>) (Figure S1).

The versatility of the optimized gene circuit was assessed by co-transfection of pYL30, pYL126, pYL125, pYL25, and pYL3 into a set of mammalian and human cell lines. Consistent secreted alkaline phosphatase (SEAP) induction by lipoteichoic acid (LTA) indicated that the system was functional in all tested cell lines, including stem-cell-derived hMSC-hTERT, suggesting broad applicability of this gene control device (Figure 1B). Different expression profiles among cell lines may be due to differences in the transfection efficiency, availability of receptor and adaptor proteins, cellular composition of downstream signaling effectors and regulators, and protein expression and secretion capacities. Considering the basal expression level, maximum expression level and induction fold, we selected HEK293 for further characterization (Figure 1B). Infect<sub>sen</sub> cells were highly sensitive to either cell-wall components of S. aureus: LTA, PGN (Figure 1C), or heat-killed whole S. aureus (HK_SA) (Figure 1D). The transcriptional activation of SEAP expression was regulated dose dependently and initiated at 10<sup>3</sup> cells/mL of HK_SA (Figure 1D). A lower inoculum of 10<sup>2</sup> colony-forming unit (CFU)/mL of live S. aureus was already sufficient to activate gene expression (Figure 1E), owing to the replication of bacteria in the cell culture medium. All transcriptional activations, however, were greatly diminished without hTLR2 expression (Figures 1C–1E), indicating that the system is hTLR2-dependent. Furthermore, the metabolism or viability of the cells as scored by constitutive SEAP expression (pSEAP2-control, P<sub>hCMV</sub>SEAP-pA) was not impacted over the entire concentration range of bacterial components or whole bacteria examined (Figures 1C and 1D).

Characterization of the gene expression kinetics of Infect<sub>sen</sub> cells, cultivated for extended periods of time in medium.
containing specific concentrations of staphylococcal cell-wall components (LTA, PGN) or whole bacteria (HK_SA), revealed consistently low leakiness and a robust, dose-dependent induction of SEAP over 3 days (Figures 2 A–2C), indicating a direct correlation between transgene expression and the microbial concentration and duration of infection. In-depth real-time monitoring of the *Staphylococcus*-responsive production of intracellular fluorescent reporter gene turboGFP further confirmed the high sensitivity to different concentrations of heat-killed, methicillin-resistant *S. aureus* (HK_MRSA) and rapid transgene switching of Infect sen (Figure 2 D). Importantly, the switching time of the system is strongly dependent on the bacterial dosage; the higher the inoculum, the faster the transgene is switched on (Figure 2 E). Furthermore, when Infect sen was exposed to HK_MRSA at varying doses for different pulse durations, high bacteria levels rapidly activated transgene expression, whereas lower levels required a longer induction pulse period to reach significant SEAP production (Figure 2 F). Finally, gene expression could be switched on and off in response to the presence or absence of heat-killed *S. aureus* (Figure 2 G).

**Design and Killing Dynamics of an Infectious Microbial Protection Device**

To obtain staphylococcal killing effects, we replaced SEAP with lysostaphin as a therapeutic output under the control of human TLR2, TLR1, TLR6, and the CD14 co-receptor (Infect pro). Functional expression of Lysostaphin in eukaryotic cells was achieved by introducing two base-pair substitutions converting Asn125 and Asn232 to glutamine residues from the mature lysostaphin sequence (Kerr et al., 2001). Since signal peptide influences protein production in mammalian cells (Knappskog et al., 2007), we screened six signal peptides of different origin and selected that derived from *Gaussia* luciferase in a humanized form for maximal lysostaphin production (Figure S2A). The enzymatic activity of HEK293-expressed lysostaphin was determined by bactericidal plate assay (Figure S2 B) and levels of lysostaphin.
expression were quantified in cell culture supernatant, showing dose-dependent responsiveness upon HK_SA induction (Figure S2C). Further modifications of the therapeutic output can be achieved by fusing the human immunoglobulin G (IgG)-derived Fc (fragment crystallizable) region to the C terminus of lysostaphin to increase its serum half-life (Czajkowsky et al., 2012) (Figures S2 D and S2E) or by co-expressing lysostaphin and TRL1068, a human monoclonal antibody that binds and disrupts biofilm (Estelles et al., 2016), to specifically target biofilm-associated infections (Figures S2F–S2H).

To ensure consistent performance over time, we constructed a stable chromosomal integration of pYL25/pYL30/pYL75 (PAP-1 x 5-NF-kB x 5-IFN-beta×tGFP-dest1-pA) in HEK293 cells. Stable Infect*pro monocalonal cell populations were selected and profiled for lysostaphin expression after induction with LTA (Figure S3). The monoclonal no. 73 cell population of Infect*pro was the best-performing transgenic cell line, combining high induction fold of lysostaphin expression, superior absolute lysostaphin levels, and improved sensitivity compared to transient transfection of the same genetic components (Figure S3).

The killing dynamics of Infect*pro toward live staphylococci were first evaluated using Staphylococcus carnosus, an apathogenic strain of the Staphylococcus genus. S. carnosus has a similar peptidoglycan composition to S. aureus (Schleifer and Fischer, 1982), rendering this organism an excellent candidate for the verification of lysostaphin activity (Strauss et al., 1998). Stimulation of Infect*sen cells with live S. carnosus at increasing concentrations exhibited a similar SEAP induction profile to

![Figure 2. Expression Kinetics, Responsiveness, and Reversibility of Infect*sen](image)

(A–C) Induction kinetics of SEAP expression with increasing concentrations of LTA (A), PGN (B), and HK_SA (C) in HEK293 cells transfected with pYL25/pYL30/pYL58 (PAP-1 x 5-NF-kB x 5-IFN-beta×tGFP-dest1-pA) and induced by heat-killed MRSA (HK_MRSA). Plot are means ± SD (n = 4).

(D) Responsiveness of Infect*sen to different bacterial doses and induction time in terms of SEAP expression. pYL25/pYL30/pYL3-transgenic HEK293 cells were cultivated with different concentrations of HK_MRSA and assayed for SEAP expression at indicated time points.

(E) Impact of bacterial concentrations and induction pulses on SEAP expression. pYL25/pYL30/pYL3-transgenic HEK293 cells were incubated with various amounts of HK_MRSA for the indicated periods. Cells were washed once before medium change and sampled at 25 hr for SEAP production.

(F) Reproducibility and reversibility of Infect*sen. pYL25/pYL30/pYL3-transfected HEK293 cells were cultivated in the presence (6 hr) or absence of 10^5 cells/mL HK_SA for 72 hr. Every 24 hr, the culture medium was exchanged, and the cell density was readjusted. All data are means ± SD (n ≥ 3).
that obtained with live *S. aureus* (Figure S4A). A time-kill study of Infect<sub>pro</sub> against *S. carnosus* showed a complete elimination of staphylococcal growth at 6 hr after an initial inoculum of 10<sup>3</sup> CFU/mL (Figure 3A). Constitutive cells transfected with pcDNA3.1(+)/pYL75 (-pYL30), pcDNA3.1(+)/pYL48 (constitutive), or Infect<sub>pro</sub> were incubated with live *S. carnosus* culture at 10<sup>3</sup> CFU/mL and sampled at the indicated time points for bacterial quantification (A) and agar plate imaging (B). Original pictures are shown in Figure S4B.

(C and D) Dose-dependent killing of *S. carnosus*. HEK293 cells transfected with pYL25/pYL30/pYL3 (control) or Infect<sub>pro</sub> were incubated with live *S. carnosus* at various concentrations and sampled at every 24 hr for bacterial counting (C) and plate assay (control, left; Infect<sub>pro</sub>, right) (D). All data are means ± SD (n = 3).

See also Figures S2, S3, and S4.

**Figure 3. Staphylococcal Sense-and-Destroy Function In Vitro**

(A and B) Time course of bactericidal effect of Infect<sub>pro</sub> against *Staphylococcus carnosus*. HEK293 cells transfected with pYL25/pYL30/pYL3 (control), pYL25/pcDNA3.1(+)/pYL75 (-pYL30), pcDNA3.1(+)/pYL48 (constitutive), or Infect<sub>pro</sub> were incubated with live *S. carnosus* culture at 10<sup>3</sup> CFU/mL and sampled at the indicated time points for bacterial quantification (A) and agar plate imaging (B). Original pictures are shown in Figure S4B.

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See also Figures S2, S3, and S4.

Bactericidal Effects *In Vivo*

After validation of both alginate-microencapsulated Infect<sub>sen</sub> and Infect<sub>pro</sub> cells in culture (Figures S5A and S5B), we tested the performance of cell capsules *in vivo*. Mice receiving
microencapsulated Infect<sub>sen</sub> cells showed dose-dependent SEAP expression in the blood in response to PGN and HK_SA (Figure 5A) or live <i>S. carnosus</i> (Figure 5B) after 24 hr. Control cells without hTLR2 expression were devoid of SEAP induction in mice at both low and high inocula of live <i>S. carnosus</i> (Figure S5C).

To determine the efficacy of Infect<sub>pro</sub> toward <i>S. aureus</i>, we employed a murine model of foreign-body infection that closely mimics human implant-associated infection (Nowakowska et al., 2014). In the prophylactic study, each mouse was injected with cell capsules prior to infection with MRSA. Tissue cage fluid from mice receiving Infect<sub>sen</sub> cells showed significant SEAP induction in mice at both low and high inocula of live <i>S. carnosus</i> (Figure 5C), confirming the diagnostic value of the system in vivo. Tissue cage fluid from the Infect<sub>pro</sub> group showed cleared planktonic bacteria in 10 out of 11 mice 1 day after infection; the other mouse started showing infection on day 2 with a consistently lower bacterial number than the average of the control group (Figure 5D). In contrast, cells deficient in lysostaphin expression showed neither bactericidal effects nor clearance (Figure 5D). Consistent results were also obtained for the growth of adherent bacteria on the tissue cage after explantation (Figure 5E). Except for the one infected mouse from each group (Infect<sub>pro</sub> and positive control) that displayed adherent bacterial growth, all other mice in both groups showed no biofilm formation (Figure 5E). The infection prevention rate reached 91% for the Infect<sub>pro</sub> group and 88% for the positive control group because of regrowth of the pathogen in 1 cage of each group after cage explantation. Furthermore, levels of lysostaphin expression in tissue cage fluid were gradually downregulated with the elimination of infection, and were reduced to the background level at the day of explantation (Figure 5F), confirming the closed-loop control characteristic of the Infect<sub>pro</sub> device.

To further evaluate Infect<sub>pro</sub>'s potential for the treatment of acute MRSA foreign-body infections, MRSA-infected mice were implanted with microencapsulated Infect<sub>pro</sub> designer cells. Analysis of the tissue cage fluid of MRSA-infected mice showed that Infect<sub>pro</sub> and constitutive lysostaphin-producing cells (HEKLys) (Figure S6A) were equally effective in clearing planktonic bacteria in all mice (Figure 6A). In contrast, treatment with vancomycin or recombinant lysostaphin only cleared planktonic bacteria in 4 out of 8 and 6 out of 8 infected mice after three days, respectively (Figure 6A). Similar results were also obtained for the treatment of bacterial biofilms in the tissue cages. While all animals treated with Infect<sub>pro</sub> and HEKLys were cleared of
MRSA biofilm, adherent MRSA was still detected in 5 out of 8 mice treated with vancomycin or lysostaphin (Figure 6B). These treatment failures were not associated with the emergence of resistance to vancomycin (there was no increase in minimum inhibitory concentration \([\text{MIC}] \leq 2 \mu\text{g/mL} \); guidelines by the Clinical and Laboratory Standards Institute, 2017) or lysostaphin (Figure S6 B). Remarkably, the cell-based treatment strategies (Infectpro and HEK Lys) achieved 100% cure rates of MRSA foreign-body infections, whereas vancomycin and lysostaphin treatment failed to eliminate infections, achieving only a 38% cure rate in each case (Figure 6C). (D-F) Quantification of bacterial load and lysostaphin in mice receiving no treatment (no treatment, \(n = 8\)), implanted with microencapsulated pYL25/pYL30/pYL3-transgenic HEK293 cells (negative ctrl, \(n = 8\)), microencapsulated pcDNA3.1(+)/pYL48-transfected HEK293 cells (positive ctrl, \(n = 8\)) or Infectpro (\(n = 8\)) prior to MRSA infection. (D) Planktonic MRSA present in tissue cage fluid of mice. (E) Adherent MRSA recovered from tissue cage biofilms in mice. (F) Lysostaphin expression in tissue cage fluid during 8 days. Student’s t test (A–C and F) and Mann-Whitney U test (D and E). All data are means ± SEM. See also Figure S5.

**DISCUSSION**

Bacterial resistance is overwhelming our ability to cope with infection, and development of new antimicrobials is failing to meet the need (Marston et al., 2016). In fact, addressing the threat of antimicrobial resistance is a perpetual challenge, because antimicrobial resistance is a predictable outcome of all antimicrobial use (Fauci and Marston, 2014). It is not just “inappropriate” use that selects for resistance. Rather, the speed with which resistance spreads is driven by microbial exposure to all antimicrobials (Spellberg et al., 2013). Thus, interventions that aim to prevent infections from occurring in the first place will reduce the need for antimicrobials and slow the appearance of resistance, thereby prolonging the useful lives of all available antimicrobials (Spellberg et al., 2013). Although vaccines hold great promise for restricting the need for antibiotics and preventing antibiotic-resistant infections, no antistaphylococcal vaccine has yet successfully passed clinical trials (Mohamed et al., 2017). Much of the difficulty stems from the lack of biomarkers for protective human immunity, the problems of translating findings from animal models into clinical trials, and safety concerns associated with unbalanced immune stimulations (Proctor, 2012). Moreover, it appears to be impossible to create a single universal vaccine against the broad range of virulence factors and diseases produced by a plethora of bacterial strains (Proctor, 2012).
prosthetic heart valves (Wang et al., 2007) and aortic grafts are especially high among patients with cardiovascular implants, such as pacemakers. Infections are often life threatening. Furthermore, infectpro effectively addresses the challenge of diagnosing implant-associated infections. This finding suggests potential clinical usefulness of the system, because once infection is established, especially in the case of biofilm-associated implant infections, standard antibiotic therapies (e.g., vancomycin, the first-line antibiotic for treatment of MRSA infections) are often no longer effective, and surgical procedures are required to extract and replace the infected device (Del Pozo and Patel, 2009). The mortality associated with MRSA infections is particularly high among patients with cardiovascular implants, such as prosthetic heart valves (Wang et al., 2007) and aortic grafts (Smids et al., 2016). Although less severe, infections associated with orthopedic devices and ventricular shunts often result in serious disabilities, and infections of mammary and penile implants can cause major disfigurement and psychological trauma (Darouiche, 2004). On the other hand, prevention of antibiotic-resistant infections before symptoms appear would be particularly beneficial to patients with compromised immune functions (e.g., due to HIV, organ transplantation, cancer, and/or treatment with immunosuppressive medication), in whom infections are often life threatening. Furthermore, infectpro effectively addresses the challenge of diagnosing implant-associated infection. Standard microbiological diagnosis of implant-associated infections requires large sample size, long culture time, and isolation of the organisms from periprosthetic tissue and/or explanted implants (Trampuz et al., 2007). We show here that application of our cellular device enables tissue fluid or blood to be used for measurement of reporter protein levels to enable detection and diagnosis of infection at the site of infection.

The availability of a closed-loop gene network to regulate therapeutic agent release in mammalian cells should also facilitate implementation of novel antiinfectives. Protein therapeutics that are less prone to antimicrobial resistance (Fitzgerald-Hughes et al., 2012), (2) can have narrow antibacterial spectra, thereby sparing non-pathogenic members of the human microbiome, which is essential to human health and infection control (Khosravi et al., 2014), and (3) they can have immunomodulatory and/or endotoxin-neutralizing effects (Finlay and Hancock, 2004) in addition to microbial killing. Furthermore, protein peptides are amenable to modifications that increase their potency (Peng et al., 2017) and half-life (Dennis et al., 2002) or can be used in combination therapy to achieve synergistic effects (Desbois et al., 2013) and multifunctional properties (Becker et al., 2016), hence reducing the chances of resistance development (Holmes et al., 2016). However, key unresolved issues regarding toxicity, stability, and method of delivery of the intact agent to sites of action are major reasons for the lack of success in systemic applications (Fitzgerald-Hughes et al., 2012). Indeed, we found that in contrast to cell-based treatment with infectpro or HEKlys, injection of recombinant lysostaphin was not sufficient to cure bacterial infections in mice, probably because of the poor pharmacokinetics of lysostaphin (Walsh et al., 2003), a general disadvantage of protein therapeutics. Our cellular device provides a means to achieve dosing control and adjustment on a self-regulated basis, as well as infection-targeted direct delivery, affording high local tissue levels of active agent without systemic effects, together with access to difficult-to-reach regions such as avascular joint or bone prostheses.

Although alginate-poly-(L-lysine)-alginate microcapsules can remain functional for at least several years (Elliott et al., 2007) and have been licensed for human clinical trials (ClinicalTrials.gov NCT01379729, NCT00790257, NCT01739829), key challenges facing the translation of designer cell-based therapies into first-in-human applications include treatment economics, the use of autologous cells, non-invasive monitoring of cell...
stability and performance over extended periods of time, and strategies to replace or remove cells when required (Orive et al., 2015). However, these challenges do not seem insurmountable, and already several strategies, such as safety switches (Di Stasi et al., 2011; Santos et al., 2019), encapsulation material with improved biocompatibility (Vegas et al., 2016a, 2016b), and real-time implant monitoring (Barnett et al., 2011; Catena et al., 2010), are being introduced to improve the safety and efficacy of cellular grafts and to allow the maintenance of cellular function for extended periods of time in humans.

We believe that the gene circuit described here could serve as a pioneering model that would be readily adaptable into different clinical settings and applicable to populations at high risk of infections, including immunologically compromised individuals, as well as patients undergoing surgery or with indwelling medical devices. We envision that this cell-based anti-MRSA approach could serve as a valuable alternative to current antibiotic therapies and an example to stimulate development of next-generation anti-infectives in the post-antibiotic era.

STAR METHODS

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SUPPLEMENTAL INFORMATION

Supplemental Information includes six figures and one table and can be found with this article online at https://doi.org/10.1016/j.cell.2018.05.039.

A video abstract is available at https://doi.org/10.1016/j.cell.2018.05.039#mmc2.

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AUTHOR CONTRIBUTIONS


DECLARATION OF INTERESTS

The authors declare no competing interests.

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REFERENCES


### KEY RESOURCES TABLE

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CONTACT FOR REAGENT AND RESOURCE SHARING

Requests for further information or resources and reagents should be directed to and will be fulfilled by the Lead Contact, Martin Fussenegger (fussenegger@bsse.ethz.ch). Plasmids containing the human TRL1068 cDNA used and generated in this study are subject to restrictions under a simple material transfer agreement (MTA) with Trellis Bioscience.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Mice
Dose-dependent transgene expression in mice induced by PGN, HK_SA or live S. carnosus were performed according to the directives of the European Community Council (2010/63/EU), approved by the French Republic and performed in 2012 by Ghislaine Charpin-El Hamri (license no. 69266309) at the Université de Lyon, F-69622 Lyon, France. Six to eight week-old native healthy female wild-type OF1 mice (oncins France souche 1) were kept in a pathogen-free environment in a 12 hr light/dark cycle (light from 6 am to 6 pm) in a temperature-controlled room (24°C). Six to seven mice were randomly assigned to experimental groups, which were not involved in previous procedures and housed in standardized cages with unrestricted access to regular mouse chow and water.

Mouse model of foreign body infection (John et al., 2011; Nowakowska et al., 2014) was established with the approval of the Kantonal Veterinaireamt Basel-Stadt (permit no. 1710). Experiments were conducted according to the regulations of Swiss veterinary law. Healthy wild-type female C57BL/6 mice at 13 weeks age (JanvierLabs) kept under specific-pathogen-free conditions (Biosafety level 2) in the animal house of the Department of Biomedicine, University Hospital Basel, were used for the MRSA infection experiments. They were housed in a 12 hr light/dark cycle (light from 7 am to 7 pm) in a temperature-controlled room (24°C) with free access to regular mouse chow and water. Mice were randomly assigned to experimental groups, which were not involved in previous procedures.

Cell types and culture conditions
Human embryonic kidney cells (HEK293, female), baby hamster kidney cells (BHK-21, male), human cervical adenocarcinoma cells (HeLa, female), human fibrosarcoma cells (HT-1080, male) and human bone marrow stromal cells transgenic for the catalytic subunit of human telomerase (hMSC-TERT, male) were cultivated in Dulbecco’s modified Eagle’s medium (DMEM, Life Technologies, cat# 31966047) supplemented with 10% (v/v) fetal calf serum (FCS, Sigma-Aldrich, St. Louis, MO, USA, cat# F7524) and 1% (v/v) penicillin/streptomycin solution (Biowest, Nuaille, France; cat. no. L0022-100). Chinese hamster ovary cells (CHO-K1, female) were grown in ChoMaster HTS (Cell Culture Technologies GmbH, Gravesano, Switzerland, art. no. CHTS-8) supplemented with 5% (v/v) FCS and 1% (v/v) penicillin/streptomycin solution. All cell lines were cultivated at 37°C in a humidified atmosphere of 5% CO₂ in air. Cell line authentication information was supplied by ATCC. Information on the sex of the cell line was obtained from ATCC, https://web.expasy.org/cellosaurus/CVCL_0317, and Shah et al. (2014).

Staphylococcal strains and growth conditions
S. carnosus were cultivated aerobically in Luria Broth (LB) medium with shaking at 37°C. Methicillin-susceptible S. aureus (MSSA) and methicillin-resistant S. aureus (MRSA) were aerobically grown in tryptic soy broth (TSB) (Becton Dickinson and Company, Allschwil, Switzerland) overnight at 37°C without shaking.

METHOD DETAILS

Cell transfection
For transfection of HEK293 and hMSC-TERT cells, a 1:3 DNA:PEI mixture of 0.525 μg DNA (pYL25:pYL30:pYL126:pYL125:pYL3 = 1:2:1:1:2) and 1.575 μL PEI (1 mg/mL in water, pH 7.0) was incubated for 15 min at room temperature before being added dropwise to 2 × 10⁵ cells/mL seeded in cell culture plates 18 hr before transfection. After 7 hr, the medium was changed and cells were induced with different ligands. BHK-21, CHO-K1, HeLa and HT-1080 cells were seeded at 1 × 10⁵ cells/mL and transfected with a 1:4 DNA:PEI ratio. Transgene expression was profiled at indicated time points after adding inducer reagents.

Bacterial inoculum and quantification
For in vitro sense-and-destroy study, MRSA overnight culture was washed twice with PBS (Bichsel, Interlaken, Switzerland) before being inoculated into cell culture medium. For the in vivo prevention study, MRSA overnight culture was washed twice with PBS before infecting mice at desired concentrations. For the in vivo acute infection, MRSA overnight culture was washed twice with 0.9% saline (Bichsel, Interlaken, Switzerland) for the inoculation of mice with adjusted concentrations as required. To prepare bacteria in the exponential-growth phase, the overnight culture was diluted 1:100 and further incubated for 6 hr at 37°C, followed by two washes with 0.9% saline, and the concentration was adjusted as required.

The CFU (colony forming unit) per milliliter (mL) was determined by plating aliquots from appropriate dilutions on MHA agar plates, followed by colony counting after 24 hr of incubation at 37°C.
Generation of stable cell lines
The polyclonal HEK_hTLR2 population, transgenic for stable expression of human TLR2, was constructed by co-transfecting 2 × 10^5 HEK293 cells/mL with 1 µg/mL pYL98 (ITR-P_hEF1α-hTLR2-pA:RPBSA-BFP-P2A-PuroR-pA-ITR) and 50 ng/mL of the Sleeping Beauty transposase expression vector pCMV-T7-SB100 (P_hCMV-SB100X-pA). After selection with 5 µg/mL puromycin for two passages, the surviving population was FACS-sorted into three different subpopulations based on blue-fluorescence intensities. The subpopulation with low 15% BFP intensity showed the best induction profile and was used for follow-up experiments.

The polyclonal HEK_hCD14/hTLR2 population, transgenic for stable expression of human CD14 and human TLR2, was generated by co-transfecting 2 × 10^5 HEK_hCD14/hTLR2 cells/mL with 1 µg/mL pYL97 (ITR-P_hEF1α-hCD14-pA:RPBSA-dTomato-P2A-BlastR-pA) and 50 ng/mL of pCMV-T7-SB100. After selection with 15 µg/mL blasticidin, the surviving population was FACS-sorted into three different subpopulations based on red-fluorescence intensities. The subpopulation with the top 15% dTomato intensity showed the best performance and was used for follow-up experiments.

The monoclonal Infectpro cell line, transgenic for bacteria-stimulated lysostaphin expression, was constructed by co-transfecting HEK_hCD14/hTLR2 cells with 1 µg/mL pYL112 (ITR-P_AP-1 × 5-NF-κB × 5-IFNγmin-hGLuc-Lys-pA:RPBSA-EGFP-P2A-ZeoR-pA-ITR) and 50 ng/mL of pCMV-T7-SB100, followed by selection in culture medium containing 1 mg/mL zeocin. The surviving population with the highest 4% EGFP expression levels was subjected to FACS-mediated single-cell sorting. Clone no. 73 showing optimal LTA-inducible lysostaphin expression was used for all follow-up studies.

HEK_LYS, stably transgenic for constitutive lysostaphin expression, was constructed by co-transfecting 2 × 10^5 HEK293 cells/mL with 1 µg/mL pYL128 (ITR-P_hCMV-hGLuc-Lys-pA:RPBSA-EGFP-P2A-ZeoR-P2A-mRuby-pA-ITR) and 50 ng/mL of the Sleeping Beauty transposase vector pCMV-T7-SB100 followed by stable selection using 1 mg/mL zeocin.

FACS-mediated single-cell sorting
HEK293 cells expressing EGFP (488nm laser, 530/30 emission filter) BFP (405 nm laser, 530/30 emission filter) or dTomato (561nm laser, 586/15 emission filter) were sorted using a Becton Dickinson LSRII Fortessa flow cytometer (Becton Dickinson, Allschwil, Switzerland) while excluding dead cells and cell doublets. Untreated HEK293 cells or parental polyclonal populations were used as negative controls.

Quantification of target gene expression
SEAP
Production of the human placental secreted alkaline phosphatase (SEAP) in culture was quantified by means of light absorbance measurements using a p-nitrophenyl phosphate-based assay (Schlatter et al., 2002). In brief, 100 µL of culture medium was heat-inactivated for 30 min at 65°C and centrifuged for 30 s at 15000 g. Subsequently, 80 µL of the heat-inactivated supernatant was transferred to a well of a 96-well plate containing 100 µL 0.2 × SEAP assay buffer (2 M diethanolamine, 2 mM L-homoarginine hydrochloride, 1 mM MgCl₂, pH 9.8). After addition of 20 µL 120 mM 4-nitrophenyl phosphate (pNPP) to each well, the increase in light absorbance was measured at 405 nm every 30 s up to 20 min. Serum levels of SEAP were quantified using a chemiluminescence-based assay (SEAP reporter gene assay).

Fluorescence
The cellular fluorescence of cell populations cultured in an atmosphere of 5% CO₂ in air at 37°C in 96-well plates was profiled every 30 min using a monochromatic Tecan Infinite M200 PRO plate reader (Tecan Group Ltd, Männedorf, Switzerland) at excitation and emission wavelengths of 482/502 nm, respectively. The autofluorescence of mock-transfected cells was subtracted from the fluorescence of cells containing Infect_sen components.

Lysostaphin
Levels of lysostaphin were determined using enzyme-linked immunosorbent assay (ELISA). 96-well EIA/RIA plates were coated overnight at 4°C with 100 µL of 4 µg/mL rabbit polyclonal anti-lysostaphin antibody in coating buffer (0.1 M sodium carbonate, pH 9.5). The plates were then washed with PBS-0.05% Tween 20 (PBST) 3 times, blocked for 4 hr at 37°C with 200 µL 3% BSA in PBS, and again washed 3 times. Serial dilutions of standards (recombinant lysostaphin) and samples (1:5 dilution in dilution buffer (PBS-0.01% Tween 80 and 0.1% BSA) were added to the wells and the plates were incubated at 37°C for 1 hr, then washed with PBST 5 times. Polyclonal anti-lysostaphin antibody conjugated to horseradish peroxidase (HRP) (lightening-link HRP) at 1 µg/mL in dilution buffer (100 µL) was added and incubation was continued for 1 hr at 37°C, followed by washing 10 times with PBST and incubation with TMB for colorimetric detection. Lysostaphin was quantified by reading the absorbance at 730 nm using an Envision 2104 multilabel (Perkin Elmer).

Human IgG ELISA
TRL1068 expression levels were quantified using sandwich ELISA. 96-well EIA/RIA plates were coated with 5 µg/mL goat anti-human IgG Fcγ fragment in coating buffer (0.1 M sodium carbonate, pH 9.5) at 4°C overnight. Plates were blocked with 200 mL per well PBS/3% BSA, pH 7.4, for 1 hr at 22°C. Human IgG standard (0.976-125 ng/mL) and samples (100 µL per well) were diluted (PBS-0.01% Tween 80 and 0.1% BSA) and incubated for 1.5 hr at 22°C. Plates were washed 3 times with 200 mL per well PBS containing 0.05% Tween-20 and then incubated with 100 µL per well of a goat anti-human IgG-alkaline phosphatase antibody (dilution 1:5000) for 1 hr at 22°C. After three washing steps (200 mL PBS/0.05% Tween-20), the plates were developed by incubation with
100 mL per well of pNPP substrate diluted in assay buffer (0.1 M ethanolamine-HCL, 5 mM MgCl₂, pH 9.8; 1 mg/mL) at 22 °C for 6 min. TRL1068 was quantified at 405 nm using an Envision 2104 multilabel reader (Perkin Elmer).

**HU binding assay of TRL1068**

Streptavidin-coated plates (Thermo Scientific Pierce, cat no. 15500) were washed (PBS-0.05% Tween 20) four times prior to use. The plates were then coated with biotinylated epitope from S. aureus histone-like DNA-bindig protein (Trellis Bioscience) at 10 µM in dilution buffer (PBS-0.01% Tween 80 and 0.1% BSA) for 2 hr with shaking at room temperature (RT). Samples from cell supernatant and epitope peptide standard were then added to each well and incubated on an orbital shaker for 30 min at RT. The plates were washed 4 times and incubated with goat anti-human IgG-alkaline phosphatase antibody (dilution 1:1000) on an orbital shaker for 30 min at RT. After washing for 4 times, plates were developed by incubation with pNPP substrate diluted in buffer (0.1 M ethanolamine-HCL, 5 mM MgCl₂, pH 9.8; 1 mg/mL) at RT for 3 min and measured at 405 nm using an Envision 2104 multilabel reader (Perkin Elmer).

**Antimicrobial assays**

**Time-kill study**

0.5 × 10⁶ Infectₜ₀ and HEK293 cells transfected with pYL25/pYL30/pYL3 (Control), pYL25/pcDNA3.1(+)/pYL75 (-pYL30) or pcDNA3.1(+)/pYL48 (Constitutive) were seeded per well in a 48-well plate one day before inoculation. After transfection for overnight, cell culture medium was changed to that without antibiotics, and appropriate amounts of S. carnosus overnight culture were added to each well. At different time points after induction, cell mixtures were collected for bacterial counting by serial dilution plating to determine bactericidal activity. Samples (30 µL) were also dropped on agar plates and incubated for 24 hr at 37°C, and then the plates were imaged.

**Bactericidal effects against MRSA**

3 independent MRSA overnight cultures were washed twice in PBS and diluted as required for inocula triplicates. Infectₜ₀, HEK₂₉₃ or control cells were seeded one day before. After transfection for overnight, cell culture medium was changed to that without antibiotics, and different concentrations of MRSA culture was added to each well for up to 96 hr at 37°C. Bacterial growth was quantified through serial plating, and lysostaphin was quantified by ELISA.

**Spot-on-lawn assay**

After 24 hr incubation, spots were imaged. Cleared spots indicate lytic activity. An exponentially growing culture of S. carnosus was spread on the plate, and 30 µL of lysostaphin-containing sample was spotted on the air-dried plate. Overnight incubation results in growth of a bacterial lawn, whereas antimicrobial activity of a protein is indicated by a clear lysis zone within the lawn, where growth has been prevented.

**Bioassay**

1.5 × 10⁶ CFU of a MRSA culture was added to MHA medium and used to fill bioassay dishes. Samples of 100 µL cell supernatant from transfected HEK293 cells were applied in punched holes in the agar plate. The plate was incubated for 20 hr at 37°C. Bacterial growth was quantified through serial plating, and lysostaphin was quantified by ELISA.

**Emergence of antimicrobial resistance in vivo**

Surviving MRSA were sampled from explanted tissue cages and screened for the emergence of resistance to vancomycin or lysostaphin during in vivo treatment. Colonies collected from bacterial subcultures were suspended to the turbidity of a McFarland 0.5 standard, and spread on Mueller-Hinton agar plates containing 2 µg/mL and 3 µg/mL of vancomycin. Plates were incubated at 37°C and profiled for bacterial growth after 20 hr. Resistance to lysostaphin was determined using a disk diffusion assay (50 µg/disc) (Kusuma and Kokai-Kun, 2005). The plates were incubated at 37°C and profiled for bacterial growth after 20 hr. Inhibition zones were compared between the treatment and control groups.

**Cell encapsulation**

Cells were encapsulated in 400 µm alginate-PLL (poly-L-lysine)-alginate beads using the Inotech Encapsulator Research IE-50R (Buechi Labortechnik AG, Flawil, Switzerland) according to the manufacturer’s protocol. The parameters were as follows: 25 mL syringe operated at a flow rate of 500 units, 200 µm nozzle with a vibration frequency of 1,024 Hz and bead dispersion voltage of 1.2 kV, stirrer speed set at 4.5 units.

**Animal testing**

**Transgene induction**

To test the bacteria-controlled transgene expression in mice, HEK293 cells transfected with pYL25/pYL30/pYL3 were encapsulated and intraperitoneally injected into OF1 mice (2-3 × 10⁶ cells/mouse). Immediately after implantation, PGN, HK_SA or live S. carnosus was administered intraperitoneally at specific dosages. After 24 hr, mice were sacrificed, blood samples were collected retroorbitally, and SEAP levels were quantified in the serum.

**Prevention of foreign body infection**

To test the diagnostic and therapeutic efficacy of cell implants against MRSA in mice, C57BL/6 mice were anesthetized followed by the subcutaneous implantation of sterile cylindrical Teflon tissue cages (32 × 10 mm; volume: 1.9 mL) with 130 regularly spaced holes (Angst + Pfister AG, Zurich, Switzerland). Upon complete wound healing, cages were tested for sterility by plating percutaneously aspirated tissue cage fluid from the lumen of the tissue cage on Columbia sheep blood agar plates. Prophylactic cell-based treatment
was started at 16 hr prior to infection with an injection of 3-5 × 10^6 microencapsulated cells contained in 400 μL serum-free DMEM into the lumen of each cage. On the day of infection, 400 μL serum-free DMEM containing 3-5 × 10^6 microencapsulated cells and 1205 ± 215 CFU of MRSA were injected into each cage. On days 1, 2, 3, 6, 8 post-infection, tissue cage fluid was collected to measure the levels of lysostaphin and to assess planktonic bacterial load by plating on blood agar plates. On day 8, mice were sacrificed, mouse serum was collected via heart puncture, and tissue cage of each mouse was explanted under aseptic conditions.

**Curing of foreign body infection**

For the acute infection study, mice were infected with 201 ± 58 CFU/cage of MRSA 2 hr before treatment with Infectpro (3 × 10^7 cells, maintained for 10 hr in medium containing 10 μg/mL PGN), HEKlys (3 × 10^7 cells), recombinant lysostaphin (5 mg/kg, once intraperitoneal) or vancomycin (200 mg/kg, every 12 hr subcutaneously). A vancomycin treatment schedule of 200 mg/kg every 12 hr was chosen to provide pharmacokinetic-pharmacodynamic parameters exceeding those reached in human treatment schemes: 1 g every 12 hr or a target AUC/MIC value of 400 (Crandon et al., 2010; Rybak et al., 2009). The explanted tissue cages were washed twice with phosphate-buffered saline followed by 30 s vortexing, sonication for 3 min at 130 W and another 30 s vortexing to release adherent bacteria from the biofilm. Quantification of adherent bacteria was performed by plating appropriated dilutions on blood agar plates. The presence of any regrowth of adherent bacteria was examined by further incubation of the cage in tryptic soy broth (TSB) (Becton Dickinson and Company, Allschwil, Switzerland) for 48 hr at 37°C. Any visualization of a positive culture of MRSA was defined as treatment failure. The prevention and cure rate was defined as the percentage of cages without growth in the individual treatment group. Concentrations of lysostaphin in the tissue cage fluid were determined by ELISA as described above.

**QUANTIFICATION AND STATISTICAL ANALYSIS**

Statistical parameters including the definitions and exact values of n, distributions and deviations, statistical test and statistical significance are reported in the Figures and corresponding Figure Legends. A two-tailed, unpaired Student’s t test and the Mann-Whitney U test were applied to determine the statistical significance of differences among groups using Graphpad Prism. Differences are considered statistically significant at p < 0.05; *p < 0.05; **p < 0.01; ***p < 0.001; ****p < 0.0001; ns, not significant.
Figure S1. Optimization of Bacteria-Regulated Transgene Expression with Different Promoter Configurations, Related to Figure 1

(A) Selection of NF-κB binding sites and minimal promoters. SEAP-expressing vectors driven by different promoters containing 5 repeats of NF-κB binding site (GGGGACTTTCC) (pYL43, P_{NF-κB}^3_5min-SEAP-pA; pYL45, P_{NF-κB}^3_5-IFNβmin-SEAP-pA; pNifty-SEAP, P_{NF-κB}^3_5-ELAM-SEAP-pA) or 5 repeats of NF-κB' binding sequence (GGGAATTTCC) (pYL44, P_{NF-κB'}^3_5min-SEAP-pA; pYL46, P_{NF-κB'}^3_5-IFNβmin-SEAP-pA) were co-transfected with pYL30/pYL25 (P_{hCMV-hTLR2-pA}/P_{hCMV-hCD14-pA}) in HEK293 cells. SEAP activity was assayed in the supernatant of cells after 24 hr in the presence of 5 μg/mL PGN.

(B) Validation of signal transactivation through AP-1-responsive promoters. pYL30/pYL25-transgenic HEK293 cells were co-transfected with NF-κB-responsive reporter plasmids with or without AP-1 binding site (pYL3, P_{AP-1}^3_5-NF-κB^3_5-IFNβmin-SEAP-pA; pYL45, P_{NF-κB}^3_5-IFNβmin-SEAP-pA; pYL47, P_{AP-1}^3_5-IFNβmin-SEAP-pA). SEAP activity was profiled at 24 hr after exposing cells with or without 5 μg/mL PGN.

(C) Screening of AP-1/NF-κB-responsive promoters for best transgene induction. HEK293 cells were co-transfected with pYL30/pYL25 and SEAP-encoding plasmids controlled by composite promoters varying in binding sequences, number of repeats and minimal promoters (pYL3, P_{AP-1}^3_5-NF-κB^3_5-IFNβmin-SEAP-pA; pYL83, P_{NF-κB}^3_5-NF-κB' IFNβmin-SEAP-pA; pYL84, P_{NF-κB}^3_5-NF-κB' IFNβmin-SEAP-pA; pYL85, P_{NF-κB}^3_5-NF-κB' IFNβmin-SEAP-pA; pYL86, P_{NF-κB}^3_5-NF-κB' IFNβmin-SEAP-pA; pYL99, P_{NF-κB}^3_5-NF-κB' IFNβmin-SEAP-pA; pYL100, P_{NF-κB}^3_5-NF-κB' IFNβmin-SEAP-pA; pYL101, P_{AP-1}^3_5-NF-κB^3_5-IFNβmin-SEAP-pA; pYL102, P_{AP-1}^3_5-NF-κB^3_5-IFNβmin-SEAP-pA; pYL103, P_{AP-1}^3_5-NF-κB^3_5-IFNβmin-SEAP-pA). SEAP activity was profiled at 24 hr after exposing cells with or without 0.5 μg/mL LTA.

(D) Screening for constitutive promoters driving hTLR2 expression. pYL25/pYL3-transgenic HEK293 cells were cotransfected with a plasmid encoding hTLR2 expression unit driven by different promoters (pYL30, P_{hCMV}^3-hTLR2-pA; pYL37, P_{hTLR2}^3-hTLR2-pA; pYL38, P_{hTLR2}^3-hTLR2-pA).

Cell were induced with 5 μg/mL PGN and assayed for SEAP activity after 24 hr. Fold induction of SEAP expression levels is shown above each figure bar. All data are means ± SD (n = 3).
Figure S2. Lysostaphin Secretion, Expression, and Further Modifications, Related to Figure 3

(A) Screening for optimal signal peptide that facilitates lysostaphin secretion. HEK293 cells were transfected with plasmids encoding constitutive lysostaphin expression directed by various signal peptides for secretion: hGLuc (pYL48, P_hCMV-hGLuc-Lys-pA); hGH (pYL33, P_hCMV-hGH-Lys-pA); hIL2 (pYL49, P_hCMV-hIL2-Lys-pA); hIL2* (pYL51, P_hCMV-hIL2*-Lys-pA); halbumin (pYL50, P_hCMV-halbumin-Lys-pA); mlgl (pYL52, P_hCMV-mlgl-Lys-pA). Lysostaphin levels were quantified by ELISA after 24 hr.

(B) Spot-on-lawn assay for staphylolytic activity. Lytic zones developed on a lawn of S. carnosus at 24 hr after sample application. Samples (30 µL) were derived from HEK293 cells transfected with pYL25/pYL30/pYL75 (TLR2 (+)), pcDNA3.1(+)/pYL48 (Constitutive), pYL25/pcdNA3.1(+)/pYL75 (TLR2 (-)) and pYL25/pYL30/pYL3 (Blank), and induced with 10^4 and 10^6 cells/mL of HK_SA for 48 hr.

(C) Lysostaphin expression in HEK293 cells transfected with pYL25/pYL30/pYL75 (P AP-1 NFKB-3 IFNBmin-hGLuc-Lys-pA) or pcDNA3.1(+)/pYL48 (Constitutive), and induced with 10^4 and 10^6 cells/mL of HK_SA for 48 hr.

(D) Expression of lysostaphin fusion proteins with human IgG-Fc fragment (hFc). HEK293 cells were transfected with lysostaphin (Lys; pYL48, P_hCMV-hGLuc-Lys-pA), Lysostaphin C-terminally fused with hFc (Lys-hFc; pYL72, P_hCMV-hGLuc-hFc-Lys-pA) or Lysostaphin N-terminally fused with hFc (hFc-Lys; pYL69, P_hCMV-hGLuc-Lys-hFc-pA), and cultivated for 24 hr before sampling for lysostaphin expression.

(E) Antimicrobial activity of lysostaphin-fusion proteins. Bioassay toward MRSA was performed with cell supernatants from Lys (pYL48)-, Lys-hFc (pYL72)-, or hFc-Lys (pYL69)-transgenic HEK293 cells after 24 hr.

(F) The influence of linker sequence on the expression of human TRL1068. HEK293 cells were transfected with pYL115/pYL119 (PhCMV-hTRL1068-HC-pA/PhCMV-hTRL1068-LC-pA), pYL93 (P_hCMV-hTRL1068-HC-RKRFF2A-LC-pA), pYL116 (P_hCMV-hTRL1068-HC-RKRFF2A-LA-pA) or pYL118 (P_hCMV-hTRL1068-HC-RKRFF2A-LA-pA), and sampled after 24 hr for the determination of hTRL1068 expression.

(G) Validation of hTRL1068 activity. Titration ELISA assay was used for the verification of the binding of the epitope (AARKGRNPQTGKEID) contained in Staphylococcus aureus histone-like DNA-binding protein. Samples were derived from HEK293 cells transfected with pYL115/pYL119, pYL93, pYL116, or pYL118 after 24 hr.

(H) Dose-dependent regulation on hTLR1068 activity. Titration ELISA assay of TRL1068 binding to the epitope contained in Staphylococcus aureus histone-like DNA-binding protein. Cell supernatants were taken from non-transfected HEK293 cells (Control) or cells transfected with pYL25/pYL30/pYL117 (P hNF-kB/-NF-kB/-IFNBmin-hTRL1068-HC-RKRFF2A-LC-pA) (TRL1068) and induced with LTA at increasing concentrations for 72 hr. All data are means ± SD (n ≥ 3).
Figure S3. Stably Transgenic Cell Lines of Infectpro, Related to Figure 3

Transgene expression of monoclonal HEK293 cells stably expressing pYL97/pYL98/pYL112 (ITR-P_hEF1a-hCD14-pA:PRPBSA-dTomato-P2A-BlastR-pA/ITR-P_hEF1a-hTLR2-pA:PRPBSA-BFP-P2A-PuroR-pA-ITR/ITR-PAP-1 x 5-NF-kB x 5-IFN-b-min-hGLuc-Lys-pA:PRPBSA-EGFP-P2A-ZeoR-pA-ITR) (Infectpro). Stable cells were cultivated with LTA at the indicated concentrations for 24 hr, and then lysostaphin levels in the cell culture supernatant were quantified. All data are means ± SD (n = 3).
Figure S4. *S. carnosus* Sense-and-Killing Dynamics, Related to Figure 3
(A) *S. carnosus*-controlled SEAP expression. HEK293 cells transfected with pYL25/pYL30/pYL3 (+pYL30) or pYL25/pcDNA3.1(+)/pYL3 (−pYL30) were incubated overnight with *S. carnosus* culture at increasing concentrations. SEAP expression was assayed after 24 hr. All data are means ± SD (n = 3).
(B) Bactericidal effect of Infect pro against *Staphylococcus carnosus* over time. HEK293 cells transfected with pYL25/pYL30/pYL3 (Control: upper left), pYL25/pcDNA3.1(+)/pYL75 (−pYL30: upper right), pcDNA3.1(+)/pYL48 (Constitutive: lower left) or Infect pro (Infectpro: lower right) were incubated with live *S. carnosus* culture at 10^3 CFU/mL and sampled at the indicated time points for plate assay.
Figure S5. Control Experiments, Related to Figure 5

(A) *In vitro* SEAP induction profiles of cells after encapsulation. HEK293 cells transfected with pYL25/pYL30/pYL3 were encapsulated and induced with increasing doses of LTA or heat-killed MSSA (HK_MSSA). SEAP activity was assayed after 24 hr. Data are means ± SD (n = 3).

(B) *In vitro* lysostaphin induction profiles of cells after encapsulation. HEK293 cells transfected with pYL25/pYL30/pYL75 were cultivated with heat-killed MRSA (HK_MRSA) or LTA at the indicated concentrations. Lysostaphin expression levels in the culture supernatant were determined after 24 hr. Data are means ± SD (n = 3).

(C) Transgene expression in mice induced by live *S. carnosus* (n = 7). Mice implanted intraperitoneally with microencapsulated pYL25/pcDNA3.1(+)/pYL3-transgenic HEK293 cells received intraperitoneal injection of *S. carnosus* culture at the indicated concentrations. Blood SEAP activity was assayed after 24 hr. Data are means ± SEM; significant difference by two-tailed t-test (ns: not significant).
Figure S6. Control Experiments, Related to Figure 6

(A) Lysostaphin production in HEK293 cells stably expressing pYL128 (ITR-P_{CMV}-hGLuc-Lys-pA;P_{MFBSA}-ZeoR-P2A-mRuby-pA-ITR) (HEK_{Lys}) versus blank HEK293 cells (Control) after 24 hr. Data are means ± SD.

(B) Inhibition zones of MRSA strains surviving from the treatment groups (Control, n = 6; LYS, n = 5) using a disk diffusion assay. Data are means ± SD; significant difference by two-tailed t test (ns: not significant).