Preconditioning with Lipopolysaccharide or Lipoteichoic Acid Protects against Staphylococcus aureus Mammary Infection in Mice

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Staphylococcus aureus is one of the most causative agents of mastitis and is associated with chronic udder infections. The persistency of the pathogen is believed to be the result of an insufficient triggering of local inflammatory signaling. In this study, the preclinical mastitis model was used, aiming to evaluate if lipopolysaccharide (LPS) or lipoteichoic acid (LTA) preconditioning could aid the host in more effectively clearing or at least limiting a subsequent S. aureus infection. A prototypic Gram-negative virulence factor, i.e., LPS and Gram-positive virulence factor, i.e., LTA were screened whether they were able to boost the local immune compartment. Compared to S. aureus-induced inflammation, both toxins had a remarkable high potency to efficiently induce two novel selected innate immunity biomarkers i.e., lipocalin 2 (LCN2) and chitinase 3-like 1 (CHI3L1). When combining mammary inoculation of LPS or LTA prior to a local S. aureus infection, we were able to modulate the innate immune response, reduce local bacterial loads, and induce either LCN2 or CHI3L1 at 24 h post-infection. Clodronate depletion of mammary macrophages also identified that macrophages contribute only to a limited extend to the LPS/LTA-induced immunomodulation upon S. aureus infection. Based on histological neutrophil influx evaluation, concomitant local cytokine profiles and LCN2/CHI3L1 patterns, the macrophage-independent signaling plays a major role in the LPS- or LTA-pretreated S. aureus-infected mouse mammary gland. Our results highlight the importance of a vigilant microenvironment during the innate immune response of the mammary gland and offer novel insights for new approaches concerning effective immunomodulation against a local bacterial infection.

Keywords: intramammary injection, Staphylococcus aureus, mastitis, lipopolysaccharide, lipoteichoic acid, chitinase 3-like 1, lipocalin 2

INTRODUCTION

Antibiotic-resistant bacteria remain a serious problem in modern health care, food safety, and animal health, resulting in an urgent need for alternatives to replace the conventional curative treatment (1, 2). The ideal strategy would be that (mis)used antibiotics become obsolete or can at least be partly replaced by preventive treatment. The latter strategy modulates the immune system
by a primary insult that enhances the activity of a beneficial antibacterial host response which, in turn, clears the bacterial hazard when it presents itself (3). In contrast to previous literature, it is now more accepted that next to the adaptive immunity, the innate immunity can also harbor some immunological memory (4).

Mastitis is a disease featured by either chronically or acutely inflamed mammary glands and is still a major challenge in the dairy sector as it affects both milk quality and animal welfare (5, 6). *Staphylococcus aureus* (*S. aureus*) in the udder of dairy cows causes harmful clinical symptoms and is associated with poor cure rates (7). The insufficient recognition of the infecting *S. aureus* in the mammary compartment is believed to hinder the host in raising an adequate innate immune response and the pathogen may, therefore, persist in the tissue (8, 9). In addition, these Gram-positive bacteria may infiltrate the mammary epithelial cell (MEC) layers and induce complicating factors such as biofilms formation (10, 11). Altogether, these observations are recognized to be a reason why classical antibiotic therapies have difficulties in eradicating contagious infections from a herd. Moreover, this currently leads to excessive use of antibiotics and even drug-resistant variants of mastitis pathogens (12–14). Commercial efforts have been made to create *S. aureus* mastitis vaccines e.g., Lysigin (Boehringer Ingelheim Vetmedica, Inc.) and Startvac (Hipra, Inc.) as an alternative to antibiotics. Unfortunately, many of these strategies have shown limited efficacy under field conditions (15).

The mammary gland is an immune receptive organ that harbors different kinds of pathogen recognition receptors (PRRs) on local cells (16). It is accepted that the outcome of the inflammatory reaction in this niche strongly depends on the efficiency of activating local PRR-expressing cells through the recognition of pathogen-associated molecular patterns (PAMPs) (17). Gram-negative mastitis pathogens contain several highly potent PAMPs that trigger multiple PRRs during an intramammary infection and result in a cytokine storm associated with acute symptoms at the cow's udder. These events eventually induce either a fast recovery or a negative outcome (18, 19). By contrast, the comprehensive identification of Gram-positive pathogens by the innate immune system is impeded due to a smaller number of highly potent immunodominant proteins that typically mediate much milder symptoms and chronic infections (20–23). Unraveling the inflammatory mechanisms that are triggered by different bovine mastitis pathogens and boost the host's immunity for low immunopotent bacterial strains such as *S. aureus* remain, therefore, mandatory. Such experiments in cows are severely limited with respect to availability of immunological research tools and requested infrastructure. Complementing *in vitro* bovine models, *in vivo* mouse models were developed to tackle these major drawbacks (24). The latter preclinical models allow the dynamic interaction between mammary gland cell populations and have been successfully used with different mastitic bacteria, including bovine *S. aureus* isolates, focusing on the early innate immune response that is highly conserved between both mammalian species (25).

Until recently, the main focus with regard to PRR-containing mammary cells was on the MECs as these were expected to be the key cell type responsible for sensing of mastitis pathogens (17, 26). Nevertheless, novel data indicate that the mammary compartment is an immunological microenvironment where multiple innate immune cells are present that are also able to sense virulence factors of pathogens (27). The differential role of the sentinel cell types responsible for immunosurveillance during an infection is gaining increasing importance in the mammary gland immunity field (28–32). It is also currently, albeit partly, known that the mammary epithelium communicates with other PRR-containing cells in the mammary gland for bacterial recognition, e.g., by shedding PAMPs such as CD14 and likely also through microRNA (33, 34). However, whether appropriate activation of this microenvironment may trigger subsequent antibacterial or anti-inflammatory responses to a secondary insult remains largely unknown.

The aim of this paper was to analyze the antibacterial capacity of the mammary microenvironment against *S. aureus* mastitis using an intraductal inoculation mouse model. First, the innate immune response of the mammary compartment resulting from exposure to virulence factors lipopolysaccharide (LPS) and lipoteichoic acid (LTA) or the well characterized bovine mastitis isolate *S. aureus* Newbould 305 was investigated. Second, we assessed immunomodulation and the antibacterial response of the complete mammary microenvironment by LPS or LTA on a subsequent insult with *S. aureus*. Third, we verified whether the preconditioning by either LPS or LTA to a *S. aureus* infection in our mouse model was unique for the local macrophages or whether other MECs are of essence for this beneficial response.

For the analysis of the host immune response in the different experiments, we preferred to measure lipocalin 2 (LCN2) and chitinase 3-like 1 (CHI3L1) to complement local cytokine profiles, rather than classical acute phase protein, such as serum amyloid A and haptoglobin. Both LCN2 and CHI3L1 are increasingly reported as key players in a wide range of inflammatory diseases. LCN2 binds and sequesters the iron-scavenging siderophore enterobactin, preventing Gram-negative bacterial iron acquisition (35, 36). Recently, it has been shown that secreted human LCN2 cloned into bovine mammary epithelial stem cells actively inhibits growth of mastitis-causing *S. aureus* as well (37). CHI3L1, on the other hand, belongs to the family of chitinase-like proteins and was initially reported in host antifungal responses as an important scavenger of chitin that is present in the fungal cell wall (38). However, further research demonstrated that the enzymatically inactive CHI3L1 has prominent functions in bacterial adhesion and invasion in host tissue, bacterial clearance, and innate immunity processes (39–43).

**MATERIALS AND METHODS**

**Intramammary Inoculation Model**

Hsd:ICR (CD1) outbred mice were purchased from a commercial supplier (ENVIGO, The Netherlands). The mice were
fed ad libitum in a controlled environment that included light and dark cycles (12 h light: 12 h darkness). All the protocols that involved animals were approved by the Committee on the Ethics of Animal Experiments of the Faculty of Veterinary Medicine, Ghent University (permit number 2014/107 and 2015/127). The experimental design of this study is shown in Figure 1. Eight-week-old female mice were mated with 10-week-old male mice. To ease the intraductal accessibility of the lactating mothers, the 10-day-old pups were weaned 2 h before the first inoculation. If possible the pups were transferred to foster mothers, the remaining pups were culled 2 h before the first inoculation. If possible the pups were transferred to foster mothers, the 10-day-old pups were weaned 2 h before the first inoculation. If possible the pups were transferred to foster mothers, the remaining pups were culled by CO₂ in a closed compartment as approved by the ethical committee of the faculty. All intraductal injections (100 µl) using a 32-gauge blunt needle were performed in the fourth gland pair of each of the mice under isoflurane anesthesia combined with the long-acting analgesic buprenorphine (10 µg/kg Vetergesic, Patheon UK Ltd., UK) and core body temperature was post-operative monitored with a rectal thermometer. Next to the ones described below, different agents were injected in the mammary gland, such as clodronate and PBS liposomes (Van Rooijen, The Netherlands). Inoculation of mammary glands with clodronate-containing liposomes depletes local alveolar macrophages and indirectly also decreases the recruitment of PMNs following intramammary LPS inoculation (27). At relevant time points post-inoculation (p.i.) mice were sedated by an intraperitoneal administered mixture of ketamine (100 mg/kg Anesketin, Eurovet Animal Health BV, Bladel, The Netherlands) and xylazine (10 mg/kg; Xylazini Hydrochloridum, Val d’Hony-Breyne, Belgium) and, subsequently, euthanized through cervical dislocation. Mammary glands were harvested to determine local bacterial loads, cytokine levels, or histopathology.

**Bacterial Products**

The mastitis isolate S. aureus Newbould 305 (ATCC 29740) was grown in brain heart infusion (Thermo Scientific, Belgium) broth for 5 h from a frozen stock. The CFU/mL was determined after 24 h of incubation at 37°C from the 1/100 diluted bacterial preculture by spectrophotometric measurements at 600 nm (OD600) and were confirmed a posteriori by plating on tryptic soy agar (Thermo Scientific, Belgium) plates. The pathogen was suspended in PBS to achieve the required concentration (10⁻¹⁻¹⁰ CFU/100µl). This 100 µl solution was then used to inoculate the mammary gland in the different mouse experiments (Figure 1). Ultrapure LPS 0111:B4 (InvivoGen, USA) and ultrapure LTA from S. aureus (InvivoGen, USA) were suspended in endotoxin-free water and diluted to the required concentration in PBS. These 100 µl LPS and LTA solutions were injected 24 h prior to the intramammary infection with S. aureus (Figure 1).

**Cell Culture**

Peritoneal macrophages were isolated as described in Walachowski et al. (44) and plated out in DMEM/F12 (Gibco) with 10% fetal calf serum for 2 h according to Mosser and Edwards (45). The non-attached cells are washed out and the sticky macrophages were trypsinized, washed, and checked for their F4/80 (clone CI:A3-1AbD Serotec) positivity on a flow cytometer (Cytoflex, Beckman Coulter). Subsequently, macrophages were activated by exposing them in vitro to 10,000 EU LPS [referred to as M(LPS)], 10,000 EU LTA [referred to as M(LTA)], or PBS [referred to as M(PBS)] for 24 h (45). Cells were then washed twice with PBS. As shown in the experimental design flow (Figure 1), these treated macrophages were injected in the mammary gland 24 h prior to a mammary infection with S. aureus.

**Quantitative Cytokine Array, LCN2, and CHI3L1 ELISA**

Hundred microliters of mammary gland homogenate were mixed with 300 µl lysis buffer supplemented with protease inhibitors (200 mM NaCl, 10 mM Tris–HCl pH 7.4, 5 mM EDTA, 1% Nonidet P-40, 10% glycerol, 1 mM oxidized l-glutathione, 100 µM PMSF, 2.1 µM leupeptin, and 0.15 µM aprotinin) to extract cellular proteins. The suspensions rested overnight at −20°C, were centrifuged at 12,250 × g for 1 h, and finally the...
supernatant was centrifuged for another 30 min to obtain a clear pellet. Protein concentration of the lysole samples was determined as previously described (46). Cytokine quantification (ProcartaPlex from eBioscience), LCN2 (Biotecnne, USA), and CHI3L1 ELISA (Biotecnne, USA) on the mammary gland lysates were all performed according to company guidelines or previously reported (47).

**In Vivo Imaging of Activated PMN**

When PMN are activated in inflamed tissues, myeloperoxidase (MPO) gets released and catalyzes the reaction of H$_2$O$_2$ and chloride ions (Cl$^-$) to produce hypochlorous acid (HOCI) (48). In turn, HOCI can oxidize luminol (5-amino-2,3-dihydrophthalazine-1,4-dione), which triggers chemiluminescence. MPO can also directly utilize superoxide anion as a substrate for peroxidase-catalyzed oxidation of luminol (45). To visualize PMN activation in our model, we injected 10 mg/ml luminol (Sigma-Aldrich, USA) i.p. and detected the bioluminescent signal with the IVIS lumina II (Caliper) (49).

**Histopathology**

Two mammary glands per condition were fixed in buffered 3.5% formaldehyde for 24 h at room temperature, dehydrated, and embedded in paraffin wax. Sections were deparaffinized, hydrated, and stained with hematoxylin and eosin. Subsequently, sections were dehydrated and mounted with a cover glass.

**Statistical Analysis**

Statistical differences were determined on normalized data with an independent sample T-test (SPSS 20.0, IBM Corporation, USA). The statistical differences for multiple comparisons were determined using ANOVA tests. Statistical significance between different groups was determined through Tukey post hoc testing.

**RESULTS**

Both LPS, and to a Lesser Extent Also LTA, Induce Mammary Immune Cell Recruitment and Local IL-8 Induction

The effect of intramammary challenge with different doses (0.01–10,000 EU) of either LPS or LTA was compared in mice at 24 h p.i. (Figure 2A). Mammary glands subjected to LPS at 10,000 EU had significantly higher IL-8 levels than glands exposed to a same dose of LTA 24 h p.i. (Figure 2B). While significant differences were detected in local IL-8 levels after a low-dose challenge of LPS (10 EU) compared to PBS (sham)-inoculated glands, IL-8 levels only increased in LTA challenged glands starting at a 100-fold higher dose (1,000 EU). This observation was in line with the histological data, showing the recruitment of immune cells in the alveolar lumen at a later time point for the mammary glands exposed to LTA compared to the mammary glands injected with LPS (Figure 2C).

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**FIGURE 2** | Effect of an intramammary injection with lipopolysaccharide (LPS) and lipoteichoic acid (LTA) on neutrophil recruitment in the intraductal mouse model. (A) Schematic presentation of the set-up. The white mammary glands represent the PBS (sham)-inoculated mammary glands, whereas the patterned glands represent the glands inoculated with LPS or LTA. (B) IL-8, a prototype chemoattractants for PMN influx was determined in mammary gland tissue 24 h p.i. of different doses of LPS and LTA for inoculum dose 0.01, 0.1, 1, 10, and 10,000 EU LPS/LTA: n = 3 mice/6 glands; for inoculum dose 100 and 1,000 EU LPS/LTA: n = 2 mice/4 glands. Represented data were pooled from two independent experiments. Values are expressed as mean ± SEM. Significance between LPS-treated glands and PBS (sham)-inoculated mammary glands or significance between LTA-treated glands and PBS (sham)-inoculated mammary glands is illustrated without brackets. Values are expressed as mean ± SEM. **P < 0.01 and ***P < 0.001 versus sham. Significance between LPS- and LTA-inoculated glands is illustrated with brackets. Values are expressed as mean ± SEM. ***P < 0.001. (C) Mammary tissue was evaluated histologically p.i. with different doses of LPS or LTA. Immune cells in the alveolar lumen are indicated with black arrows and enlarged in the images below. Scale bar, 100 µm. p.i., post-inoculation.
LPS Treatment More Effectively Induces Local LCN2 and CHI3L1 than LTA Treatment, Partially Concomitant with Mammary Immune Cell Recruitment

Lipopolysaccharide was more effective than LTA to significantly induce both mammary LCN2 and CHI3L1 compared to PBS (sham)-inoculated mammary glands (Figure 3). Nevertheless, it was also observed that local LCN2 and CHI3L1 levels not fully mimicked the previously mentioned IL-8 levels. Indeed, dissimilar to the IL-8 patterns, 10-fold higher doses of both LPS (≥100 EU) and LTA (10,000 EU) were needed to significantly induce CHI3L1 in the mammary gland compared to sham-inoculated glands. Furthermore, no significant difference in LCN2 and CHI3L1 levels could be observed between the LPS and LTA inductions at the highest dose (10,000 EU). Interestingly, recruitment of neutrophils was observed in the mammary alveoli at the time point that both the LCN2 and CHI3L1 levels in the LTA and LPS groups were significantly increased compared to the PBS (sham)-inoculated glands.

S. aureus Effectively Recruits Immune Cells during Mammary-Induced Inflammation Independently of LCN2

In contrast to sterile mammary gland-induced inflammation, inoculation of 1,000 CFU S. aureus Newbould 305 (S. aureus) in the murine mammary gland resulted in high bacterial counts (9.2 ± 0.3 log10 CFU/g mammary gland), accompanied by a prominent local host response at 24 h.p.i. (Figure 4A). This inflammatory response was characterized by the production of multiple cytokines and chemokines as comparing tissue samples of a S. aureus- and a PBS (sham)-inoculated murine mammary gland (data not shown). In addition, the significant increase of IL-8 levels at 24 p.i. in the S. aureus-infected glands compared to sham-inoculated control glands could be demonstrated (Figure 4B, left). This latter observation could be confirmed upon histological examination through a pronounced presence of immune cells in the lumen of the mammary alveoli with S. aureus, compared to sham-inoculated mammary glands (Figure 4B, right). Interestingly, the IL-8 levels showed a median fold increase of 14.0 between S. aureus- and sham-inoculated mammary glands, whereas a 34.1 and 14.6 median fold increase was observed for 10,000 EU LPS- or LTA-exposed mammary glands, respectively, compared to the control glands. Although LCN2 levels in these infected glands were not significantly different from the LCN2 levels in sham-inoculated control glands, local induction of CHI3L1 levels by S. aureus could still be observed at 24 h.p.i. (Figure 4C). More specifically, the median fold increase of local CHI3L1 levels in S. aureus-inoculated glands compared to sham-inoculated control glands was 2.88, which is an order of magnitude lower compared to the CHI3L1 levels following exposure to 10,000 EU LPS or LTA (8.16 and 5.13, respectively).

Preconditioning of the Complete Mammary Gland Compartment with LPS or LTA Strongly Immunomodulates S. aureus Mastitis, Reducing Bacterial Loads, Recruiting PMN, and Increasing Local CHI3L1 or LCN2 Levels, Respectively

As shown in Figure 1, mice were intramammary injected with PBS, LPS (10,000 EU), or LTA (10,000 EU) 24 h prior to S. aureus infection. With this setup, we wanted to verify whether LPS or LTA preconditioning is able to boost the murine mammary gland microenvironment against a S. aureus infection (Figure 5A). All mice pretreated with either LPS or LTA showed a significant decrease in local bacterial loads 24 h.p.i. with S. aureus compared to PBS (sham)-pretreated mammary glands (Figure 5B). Histology also showed that all conditions were characterized by the presence of PMN in the alveoli of the mammary glands at 24 p.i. with S. aureus (Figure 5C). However, on the molecular level, this observation during LPS and LTA pretreatment was accompanied with a significant reduction in mammary IL-8 levels compared to sham preconditioning (Figure 5D).

As previously shown, preconditioning with PBS prior to S. aureus infection induced low levels of CHI3L1, but did not induce LCN2 levels in the mammary gland. Preconditioning with

FIGURE 3 | Effect of an intramammary injection with lipopolysaccharide (LPS) and lipoteichoic acid (LTA) on local lipocalin 2 (LCN2) and chitinase 3-like 1 (CHI3L1) levels in the intraductal mouse model. LCN2 and CHI3L1 levels were determined in mammary gland tissue 24 h.p.i. of different doses of LPS and LTA (n = 3 mice/6 glands for inoculum dose 0.01, 0.1, 1, 10, and 10,000 EU LPS/LTA; n = 2 mice/4 glands for inoculum dose 100 and 1,000 EU LPS/LTA). Represented data were pooled from two independent experiments. p.i., post-inoculation. Values are expressed as mean ± SEM (**P < 0.01 and ***P < 0.001 versus sham).
10,000 EU LPS or LTA prior to S. aureus infection, on the other hand, boosted either one of these innate immune biomarkers (Figure 5E). More specifically, a significant increase in mammary CHI3L1 levels was detected with LPS preconditioning prior to S. aureus infection, whereas a significant increase in mammary LCN2 levels was observed with LTA preconditioning prior to S. aureus infection.

Local cytokine profiling showed that the inflammatory response following infection with S. aureus was influenced by the LPS and LTA preconditioning at an inoculum dose of 10,000 EU. IL-1 alpha, IL-1 beta, IL-17A, IL-6, MCP-1, and TNF-alpha followed the same pattern as the neutrophil chemoattractant IL-8 profile and were significantly reduced by LPS and LTA preconditioning compared to PBS preconditioning in the S. aureus-infected mammary glands (Figure 5F). Mammary RANTES/CCL5 levels were significantly increased by LPS preconditioning compared to the other conditions, while LTA preconditioning significantly reduced local BAFF levels compared to PBS preconditioning (Figure 5G).

**Alveolar Macrophages Have an Immunomodulatory Function during LPS and LTA Preconditioning against S. aureus-Induced Mastitis**

Alveolar macrophages are a major innate immune cell type in the mammary gland that contain PRRs on the cell surface to sense bacterial antigens (such as LPS or LTA). Macrophages are known to be a crucial cell type during immune tolerance, e.g., endotoxin tolerance. It is a protective mechanism against excessive inflammation where macrophages change their responsiveness to LPS following a first exposure to LPS (50). To test their immunomodulatory potential of macrophages in the mammary gland, isolated murine peritoneal macrophages were exposed ex vivo either to 10,000 EU LPS [referred to as M(LPS)], 10,000 EU LTA [referred to as M(LTA)] or PBS [referred to as M(PBS)]. The response of these macrophages was subsequently verified with the two previously used innate immune biomarkers LCN2 and CHI3L1. Activated macrophages, i.e., M(LPS) as well as M(LTA),...
Preconditioning against S. aureus in a Mouse Mastitis Model

Figure 5

A

B

C

D

E

F

FIGURE 5 | Continued
were found to secrete significantly more LCN2 and CHI3L1 than non-activated macrophages, i.e., M(PBS) (Figure 6A). Of notice, the secreted levels of both these innate immune biomarker proteins showed a significantly higher increase with M(LPS) than with M(LTA).

In a next step, the immune modulating effect of these M(LPS)- versus M(LTA)-activated macrophages was studied in a mammary gland environment. As shown in Figure 1, M(PBS), M(LPS), or M(LTA) were first transferred into mammary mammary glands, followed by an inoculation with 10⁴ CFU S. aureus bacteria or a PBS (sham) inoculation 24 h later (Figure 6B). The transfer of M(PBS), M(LPS), or M(LTA) did not significantly decrease the bacterial loads in the murine mammary glands at 24 p.i. with S. aureus. Nevertheless, there were 40 and 43% of the mammary glands in, respectively, the M(LPS) and the M(LTA) transfer group that had a decreased bacterial load p.i. with S. aureus (Figure 6C). In comparison, the number of glands with a decrease in bacterial loads in the M(PBS)-transferred glands was 17%.

The non-significant difference in bacterial loads between the different transfer groups p.i. with S. aureus can be reasoned by the lack of PMN recruitment induced by the activated macrophages. Indeed, based on histology, M(PBS), M(LPS), or M(LTA) transfer followed by sham inoculation 24 h later did not induce any recruitment of PMN at 48 h post-macrophage transfer (Figure 6D). The absence of PMN in the alveolar lumen could also be observed based on the low mammary IL-8 levels in the glands that received M(PBS), M(LPS), or M(LTA) followed by sham inoculation (Figure 6E, left). By contrast, the mammary glands that were infected with S. aureus at 24 h post-macrophage transfer displayed a significantly increased IL-8 response compared to M(PBS) and PBS (sham). Moreover, statistical analysis showed that the M(PBS), M(LPS), and M(LTA) transfer induced a different IL-8 response in the infected mammary glands. To visualize these differences in IL-8 levels more easily, the effect of macrophage transfer followed by S. aureus inoculation in the murine mammary gland was standardized with the IL-8 levels measured following macrophage transfer and subsequent sham inoculation (Figure 6E, right). This analysis showed that immune tolerance was limited with M(LPS) while the transfer of M(LTA) followed by S. aureus infection significantly decreased the IL-8 response in the challenged murine mammary glands compared to M(PBS) and M(LPS) transfer and subsequent S. aureus infection.

Next, we investigated whether LPS- and LTA-activated macrophages modulate the response of both LCN2 and CHI3L1 in vivo to a similar extent as in vitro. Standardizing the LCN2 and CHI3L1 levels with M(PBS), M(LTA), and M(PBS) transfer in infected versus non-infected mammary glands showed that M(LPS) transfer significantly boosted both local LCN2 and CHI3L1 levels compared to M(PBS) or M(LTA) transfer upon S. aureus infection (Figure 6F). By contrast, but similarly to what was found in vitro, M(LTA) induced lower local LCN2 levels following S. aureus inoculation, compared to the M(LPS) transfer group. Local CHI3L1 levels, however, were not induced by the M(LTA) transfer group upon S. aureus infection (Figure 6F).

Local cytokine profiling further identified a partly and differential immunosuppressive effect due to the activated macrophages (Figure 6G). A significant decrease in the levels of MCP-1, IL-17A, and IL-1 alpha was observed after M(LTA) transfer in the mammary glands p.i. with S. aureus compared to M(PBS) transfer. M(LPS) transfer only induced a significant decrease in IL-17A levels compared to M(PBS) transfer. Other cytokines that showed elevated levels at 24 h p.i. with S. aureus included IL-6, IL-1 beta, RANTES/CCL5, and BAFF remained unaltered with M(LPS) and M(LTA) transfer compared to M(PBS) transfer (data not shown).
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**Figure 6** Continued
Effect of clodronate treatment in the murine mammary gland against Staphylococcus aureus-induced mastitis. Histology showed that intramammary inoculation of clodronate-containing liposomes could decrease the recruitment of PMN in the alveoli of the mammary gland upon challenge with LPS, whereas inoculation of PBS-containing liposomes followed by LPS challenge resulted in a typical strong influx of PMN (Figure 7B). This distinction was not clearly observed when comparing both liposome groups upon challenge with LTA. Luminol-dependent bioluminescence is able to visualize detection of MPO activity and its use in vivo is specified for active PMN rather than macrophages. In vivo imaging with luminol confirmed the effect of clodronate treatment in the murine mammary gland upon LPS challenge by a decreased MPO activity compared to the mammary glands that were treated with PBS-containing liposomes (Figure 7C). Of note, the low recruitment or activity of PMN by LTA compared to LPS as seen previously, impaired the visualization of the bioluminescent signal from luminol in both conditions. These observations indicate that clodronate treatment has mainly an effect on LPS- rather than LTA-induced inflammation. This was also indirectly shown through the stronger decrease in mammary gland IL-8 levels following clodronate treatment in the LPS-challenged group compared to the LTA-challenged group (Figure 7D).

To verify whether other PRR-expressing mammary cells—next to macrophages—may be activated and have immunomodulatory effects during a S. aureus infection, mammary glands of mice were challenged with a clodronate- or PBS-containing liposome treatment followed by either LPS or LTA challenge (both 10,000 EU) and a S. aureus infection (Figures 1 and 8A). Local depletion of macrophages by clodronate-containing liposomes did not significantly increase the local bacterial loads when LPS or LTA was intramammarily inoculated prior to S. aureus (Figure 8B). However, the innate immune response following

**Figure 6** The effect of ex vivo activated macrophages on an intramammary infection with Staphylococcus aureus in mice. (A) Lipocalin 2 (LCN2) and chitinase 3-like 1 (CHI3L1) levels were determined in medium of harvested peritoneal macrophages 24 h post-exposure to PBS [M(PBS)], lipopolysaccharide (LPS) [MLPS], or lipoteichoic acid (LTA) [M(LTA)] (n = 3 for each group). Represented data were collected from one experiment. (B) Schematic presentation of the set-up. Macrophages ex vivo exposed to PBS (white glands), LPS or LTA (patterned glands) were transferred into mammary glands and inoculated with PBS (sham) or infected with S. aureus. (C) Bacterial loads 24 h.p.i. with S. aureus in mammary glands that received M(PBS), MLPS, or M(LTA) transfer [M(PBS); n = 12 glands; M(LPS); n = 14 glands; M(LTA); n = 10 glands]. (D) Representative images of alveoli of mammary glands that received M(PBS), MLPS, or M(LTA) transfer followed by PBS (sham). Transferred macrophages are indicated by an arrow. Scale bar, 100 μm. (E) Quantitative levels of IL-8 determined in mammary gland tissue that received M(PBS), MLPS, or M(LTA) transfer followed by an inoculation with PBS (sham) or S. aureus (left) [PBS inoculations: n = 13 glands for M(PBS), n = 12 glands for M(LPS), n = 12 glands for M(LTA); S. aureus inoculations: n = 12 glands for M(PBS), n = 6 glands for M(LPS), n = 4 glands for M(LTA)]. Normalized quantitative IL-8 values between macrophage transferred glands that received PBS (sham) or S. aureus. (F) Normalized quantitative LCN2 and CHI3L1 levels between M(PBS), MLPS, or M(LTA) transfer [M(PBS); n = 12 glands for M(PBS), n = 14 glands for M(LPS), n = 10 glands for M(LTA); CHI3L1: n = 12 glands for M(PBS), n = 12 glands for M(LPS), n = 12 glands for M(LTA)]. (G) Normalized quantitative MCP-1, IL-17A, and IL-1 alpha levels between M(PBS), MLPS, or M(LTA) transfer [M(PBS); n = 17 glands; MLPS; n = 12 glands; M(LTA); n = 16 glands]. Represented data were pooled from three independent experiments. p.i., post-inoculation. Values are expressed as mean ± SEM (P < 0.05, **P < 0.01, and ***P < 0.001).

**Figure 7** Preconditioning with LPS and LTA against S. aureus-induced mastitis. Preconditioning with LPS and LTA (both 10,000 EU) was intramammary inoculated prior to S. aureus infection to verify whether other PRR-expressing mammary cells—next to macrophages—may be activated and have immunomodulatory effects during a S. aureus infection, mammary glands of mice were challenged with a clodronate- or PBS-containing liposome treatment followed by either LPS or LTA challenge (both 10,000 EU) and a S. aureus infection (Figures 1 and 8A). Local depletion of macrophages by clodronate-containing liposomes did not significantly increase the local bacterial loads when LPS or LTA was intramammarily inoculated prior to S. aureus (Figure 8B). However, the innate immune response following

**Macrophage-Independent Signaling Is Important for Bacterial Clearance and Immunomodulation during Mammary Gland Preconditioning with LPS and LTA against S. aureus-Induced Mastitis**

Clodronate liposomes were used to verify the influence of macrophage depletion p.i. LPS or LTA (Figure 7A). Histology showed that intramammary inoculation of clodronate-containing liposomes could decrease the recruitment of PMN in the alveoli of the mammary gland upon challenge with LPS, whereas inoculation of PBS-containing liposomes followed by LPS challenge resulted in a typical strong influx of PMN (Figure 7B). This distinction was not clearly observed when comparing both liposome groups upon challenge with LTA. Luminol-dependent bioluminescence is able to visualize detection of MPO activity and its use in vivo is specified for active PMN rather than macrophages. In vivo imaging with luminol confirmed the effect of clodronate treatment in the murine mammary gland upon LPS challenge by a decreased MPO activity compared to the mammary glands that were treated with PBS-containing liposomes (Figure 7C). Of note, the low recruitment or activity of PMN by LTA compared to LPS as seen previously, impaired the visualization of the bioluminescent signal from luminol in both conditions. These observations indicate that clodronate treatment has mainly an effect on LPS- rather than LTA-induced inflammation. This was also indirectly shown through the stronger decrease in mammary gland IL-8 levels following clodronate treatment in the LPS-challenged group compared to the LTA-challenged group (Figure 7D).

To verify whether other PRR-expressing mammary cells—next to macrophages—may be activated and have immunomodulatory effects during a S. aureus infection, mammary glands of mice were challenged with a clodronate- or PBS-containing liposome treatment followed by either LPS or LTA challenge (both 10,000 EU) and a S. aureus infection (Figures 1 and 8A). Local depletion of macrophages by clodronate-containing liposomes did not significantly increase the local bacterial loads when LPS or LTA was intramammarily inoculated prior to S. aureus (Figure 8B). However, the innate immune response following

**Figure 8** (A) Representative images of alveoli of mammary glands that received M(PBS), M(LPS), or M(LTA) transfer followed by an inoculation with PBS (sham) or S. aureus (left) [PBS inoculations: n = 13 glands for M(PBS), n = 12 glands for M(LPS), n = 12 glands for M(LTA); S. aureus inoculations: n = 12 glands for M(PBS), n = 6 glands for M(LPS), n = 4 glands for M(LTA)]. Normalized quantitative IL-8 values between macrophage transferred glands that received PBS (sham) or S. aureus. (B) Normalized quantitative LCN2 and CHI3L1 levels between M(PBS), MLPS, or M(LTA) transfer [M(PBS); n = 12 glands for M(PBS), n = 14 glands for M(LPS), n = 10 glands for M(LTA); CHI3L1: n = 12 glands for M(PBS), n = 12 glands for M(LPS), n = 12 glands for M(LTA)]. (C) Normalized quantitative MCP-1, IL-17A, and IL-1 alpha levels between M(PBS), MLPS, or M(LTA) transfer [M(PBS); n = 17 glands; MLPS; n = 12 glands; M(LTA); n = 16 glands]. Represented data were pooled from three independent experiments. p.i., post-inoculation. Values are expressed as mean ± SEM (P < 0.05, **P < 0.01, and ***P < 0.001).
S. aureus infection was influenced by this depletion as IL-8 levels significantly increased in infected mammary glands challenged with either LPS or LTA (Figure 8C).

In this experimental setup, LCN2 and CHI3L1 also showed different patterns upon S. aureus infection in the LPS- versus LTA-challenged groups. Intramammary inoculation with clodronate-containing liposomes and subsequent LTA, but not LPS preconditioning significantly increased local LCN2 levels compared to the PBS liposome treatment in infected glands. By contrast, local CHI3L1 levels were decreased in mammary glands...
that received LPS preconditioning and clodronate treatment followed by *S. aureus* infection. Liposomes in the mammary gland did not affect the local CHI3L1 levels p.i. with LTA preconditioning followed by *S. aureus* infection (Figure 8D).

Local cytokine profiling also showed that, similar to the neutrophil attractant IL-8 profile, the macrophage chemoattractant MCP-1 as well as IL-6 were increased by the macrophage depletion process both with LPS and LTA preconditioning followed by...
S. aureus infection (Figure 8E). BAFF was only affected by clodronate treatment in the LPS-pretreated group with S. aureus infection. IL-17A and RANTES levels, respectively, were significantly increased and decreased, respectively, in the LTA-pretreated group with clodronate-depleted macrophages and S. aureus infection.

**DISCUSSION**

*Staphylococcus aureus* mastitis is contagious and does not respond well to classical antibiotic therapies (12–14). This Gram-positive pathogen is characterized by biofilm formation and intra-epithelial localization, which hinder access of therapeutic agents (11). Consequently, *S. aureus* udder infections have led to the overuse of antibiotics that increases drug resistance (51–53). Therefore, either novel vaccination or other innovative preventive measures with negligible side-effects are highly anticipated as they have remained unsuccessful in Gram-positive mastitis (7).

The hosts’ recognition of pathogen-associated factors and its innate immune response determines the outcome of bovine mastitis (54, 55). The persistent character of *S. aureus* udder infections is believed to result from a limited mammary cytokine production compared to the “cytokine storm” raised against *E. coli* (20, 21). This implies that *E. coli* antigens are either more easily recognized by mammary PRR or that Gram-negative virulence factors have a higher potency to trigger a strong local immune response. In this study, the dose-dependent stimulation with LPS versus LTA allowed controlled investigation of the potency and differential activation of PRRs through monitoring of local IL-8 production and immune cell infiltration at 24 h p.i. Histology demonstrated that the highest inoculum dose of LPS and LTA initiated innate immune cell recruitment. However, the innate immune response was significantly faster and higher stimulated by LPS compared to LTA.

It has been suggested that LPS may influence the course of a subsequent bacterial udder infection (56, 57). More specifically, injection of LPS in the udder reduced the severity of a subsequent experimental *E. coli* mastitis (57). Whether preconditioning enhances the pathogen recognition or the host immune response is not clear. *Staphylococcus aureus* stimulates TLR2 but lacks efficient activation of TLR4 compared to *E. coli* (58, 59). Similarly, as TLR4 activation by LPS is protective against *E. coli* mastitis, this study aimed to boost TLR2-induced immune response of the murine mammary microenvironment by LTA to protect the host against *S. aureus* mastitis. On the other hand, we also explored the possibility that we should compensate for the lack of TLR4 signaling during *S. aureus* mastitis by preconditioning with LPS. Our results confirmed the observations previously suggested for an *in vitro* *S. aureus* infection (60, 61): preconditioning the host significantly reduced bacterial loads at 24 h p.i. We also described the participation of local chemo- or cytokines. Although several major local pro-inflammatory mediators decreased, the host was still able to recruit neutrophils to the lumen of LPS or LTA preconditioned alveoli. The lower local IL-8 levels are, therefore, likely an indication of abolished further neutrophil recruitment rather than of their initial LPS- or LTA-induced recruitment. In addition, local MCP-1 and IL-17A levels are signature cytokines for macrophage recruitment and T helper cell activation, indicating that the immunomodulatory response is not limited to neutrophils. On the one hand, LTA but not LPS preconditioning additionally lowered local BAFF levels, a cytokine necessary for proliferation and differentiation of B cells. On the other hand, LPS but not LTA preconditioning elevated local RANTES/CCL5 levels, in line with the *in vitro* observations of cytokine networks after recurrent exposure of bovine MECs to LPS (56). Overall, our different preconditioning protocols against *S. aureus* mastitis together with the reported sterile mammary inflammation data indicate that the two key Gram-negative and -positive virulence factors have unique traits and differentially immunomodulate the mouse mammary gland.

Mammary sentinel cell types—each featuring a distinct immune competence—govern the host response to virulence factors in the mammary gland (16). Some mammary epithelial tissue exposure experiments have described an effect of LPS and LTA on the innate immune response, but how these key virulence factors modulate the mammary gland microenvironment has been scarcely studied *in vitro* and especially *in vivo* (16, 27, 32). The validated murine intramammary inoculation model previously demonstrated that local macrophages are of key importance for the LPS–TLR4 axis-mediated neutrophil recruitment into the alveolar lumen, while this is far less the case for TLR2-mediated signaling (27, 62). In general, macrophages can be programmed to different priming and tolerance states by varying dosages of LPS (63). Partly corroborating the observations from our *in vivo* sterile mammary inflammation data, we demonstrated that macrophages alone affect the mammary environment albeit to a limited degree. Indeed, *in vitro* preconditioning of alveolar macrophages with either LPS or LTA, and their transfer into sterile mammary glands did not induce a neutrophil influx.

**FIGURE 8 | Continued**

Effect of clodronate liposome treatment on lipopolysaccharide (LPS) and lipoteichoic acid (LTA)—preconditioning to *Staphylococcus aureus* in the murine mammary gland. (A) Schematic presentation of the experimental set-up. The treatment groups of Figure 7 received an infection with *S. aureus* (black glands) 24 h p.i. (B) The bacterial loads in mammary glands that received clodronate before LPS or LTA preconditioning prior to a *S. aureus* infection (*n* = 7) glands for the PBS liposome groups (i.e., macrophage +; *n* = 10 glands for the clodronate liposome groups (i.e., macrophage –)). (C) The IL-8 levels in mammary glands that received clodronate injection before LPS or LTA preconditioning prior to a *S. aureus* infection (PBS liposomes: *n* = 8 for the LPS preconditioning group, *n* = 6 for the LTA preconditioning group; clodronate liposomes: *n* = 10 for both the LPS and LTA preconditioning group). (D) Lipocalin 2 (LCN2) and chlorhina 3-like 1 (CHI3L1) levels in mammary glands that received clodronate injection before LPS or LTA preconditioning prior to a *S. aureus* infection (PBS liposomes: *n* = 7 (LCN2) and *n* = 8 (CHI3L1) for the LPS preconditioning group; clodronate liposomes: *n* = 8 (LCN2) and *n* = 10 (CHI3L1) for both the LPS preconditioning group, *n* = 10 (LCN2/CHI3L1) for the LTA preconditioning group). (E) MCP-1, IL-17A, RANTES/CCL5, BAFF, and IL-6 levels in mammary glands that received clodronate injection before LPS or LTA preconditioning prior to a *S. aureus* infection (*n* = 7 glands for the PBS liposome groups; *n* = 10 glands for the clodronate liposome groups). Represented data were collected from one experiment. Values are expressed as mean ± SEM. ns, not statistically significant; p.i., post-inoculation (⁎*P* < 0.05, **⁎*P* < 0.01, and ***⁎*P* < 0.001).
Moreover, upon *S. aureus* infection, activation of these differently preconditioned macrophages reduced the bacterial loads only to a limited extent compared to sham-exposed macrophages. Of relevance, the different TLR trigger changed the response of macrophages as LTA-dependent macrophage signaling was more efficient in reducing the inflammatory signaling compared to the LPS-dependent macrophage signaling.

Injection of macrophages treated with LPS or LTA into the gland prior to infection attenuates the *S. aureus*-induced cytokine release when compared to mice receiving PBS-treated macrophages. However, the pretreated transferred macrophages are not sufficient to reduce the bacterial burden, suggesting that LPS and LTA have to act directly on cells in the mammary gland. MECs have been proposed as special non-professional sentinel cells that are responsible for macrophage-independent local signaling (64). To investigate the role of the mammary epithelium in the LPS- and LTA-mediated immunomodulatory response against *S. aureus*, macrophages were depleted in our preconditioning mastitis model by clodronate-containing liposomes. This local treatment did not change the antibacterial host response, but strongly immunomodulated a subsequent mammary *S. aureus* infection. Consequently, the antibacterial response after preconditioning with either LPS or LTA likely originated from macrophage-independent signaling by the mammary epithelium. Moreover, these data extend those obtained with the preconditioned macrophages. Indeed, macrophages again immunomodulated the mammary compartment during preconditioning as clodronate depletion influenced the local cytokine profile. Depletion of local macrophages was able to further distinguish the differential LPS versus LTA immunomodulatory signaling prior to a *S. aureus* infection. More specifically, our data showed that macrophages inhibited BAFF levels upon LPS preconditioning, while inhibiting IL-17A upon LTA preconditioning. In addition, macrophages had a pro-inflammatory effect by induction of RANTES/CCL5 upon LTA preconditioning prior to *S. aureus* infection. However, based on the methods used in this study, it is not possible to verify whether the changes in cytokine, chemokine, and antimicrobial peptide levels following clodronate pretreatment and subsequent LPS/LTA preconditioning followed by a *S. aureus* infection are either due to an enhanced protein production by the mammary epithelium or the consequence of a reduced uptake by the macrophages after their depletion. Gene expression profiles on isolated mammary cell types would be of added value to distinguish between both phenomena, albeit these were not assessed in this study. In essence, mammary macrophage response may be induced by different mammary cell types and may aid in the evaluation of preconditioning protocols.

**ETHICS STATEMENT**

This study was carried out in accordance with the FELASA guidelines and recommendations. The protocol was approved by the Committee on the Ethics of Animal Experiments of the Ghent University.

**AUTHOR CONTRIBUTIONS**

KB, JS, and KD were essential for the conception and design of the work; the acquisition, analysis, and interpretation of data for the work; and drafted the work. TB and EM revised it critically for important intellectual content. KB, JS, KD, TB, and EM gave final approval of the version to be published and agreed to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.
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