Article

**Staphylococcus Aureus Protection-Related Type 3 Cell-Mediated Immune Response Elicited by Recombinant Proteins and GM-CSF DNA Vaccine**

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Abstract: *Staphylococcus aureus* mastitis remains a major challenge for dairy farming. Here, 24 mice were immunized and divided into four groups: G1: control; G2: Granulocyte Macrophage Colony-Stimulating Factor (GM-CSF) DNA vaccine; G3: FOF1 ATP synthase subunit α (SAS), succinyl-diaminopimelate (SDD), and cysteinyl-tRNA synthetase (CTS) recombinant proteins; and G4: SAS+SDD+CTS plus GM-CSF DNA vaccine. The lymphocyte subpopulations and the intracellular interleukin-17A (IL-17A) and interferon-γ production in the draining lymph node cells were immunophenotyped by flow cytometry. The immunophenotyping and lymphocyte proliferation was determined in spleen cells cultured with and without *S. aureus* stimulus. Immunization with *S. aureus* recombinant proteins generated memory cells in draining lymph nodes. Immunization with the three recombinant proteins plus GM-CSF DNA led to an increase in the percentage of IL-17A+ cells among overall CD44+ (memory), T CD4+, CD4+ T CD44+ CD27+, γδ TCR, γδ TCR+ CD44+ CD27+ and TCRVγ4+ cells. Vaccination with *S. aureus* recombinant proteins associated with GM-CSF DNA vaccine downregulates Tn2 immunity. Immunization with the three recombinant proteins plus the GM-CSF DNA led to a proliferation of overall memory T, CD4+ and CD4+ TEM cells upon *S. aureus* stimulus. This approach fostered type 3 immunity, suggesting the development of a protective immune response against *S. aureus*.
Keywords: vaccine; Staphylococcus aureus; T cell response; mastitis; bovine

1. Introduction

Bovine mastitis is the disease with the greatest global impact on dairy farming, causing a pronounced reduction in milk production and quality. Several etiologic agents are implied in mastitis, but Staphylococcus aureus is considered one of the most prevalent mastitis pathogen. It is an extremely critical pathogen due to its pathogenicity, contagiousness, and refractoriness to antimicrobial treatment [1,2], as well as critical issues related to food security and public health [3]. Therefore, to decrease the impact of S. aureus in dairy farming, alternative strategies to control mastitis should be evoked, such as the development of an effective vaccine.

Staphylococcus aureus pathogenicity is a complex issue; thus, a single antigen may not confer protective immunity [4]. Most studies have focused on S. aureus virulence factors for developing vaccines against S. aureus intramammary infections (IMIs) in dairy cows, but this initiative has been unsuccessful so far. Here we used an innovative approach focusing on the animals that apparently had overcome the infection by developing successful anti-pathogen host immune responses. We intended to take advantage of the knowledge derived from detailed analysis and characterization of the immune response that favored the host to identify promising epitopes for immunization [1,2,5,6]. Using this approach, we previously searched S. aureus-derived proteins likely related to protection [2]. Based on these antigenic proteins revealed by serum immunoproteomics approach, we obtained three candidate proteins: F0F1 ATP synthase subunit α (SAS), succinyl-diaminopimelate (SDD), and cysteinyl-tRNA synthetase (CTS) [2].

Some vaccine candidate antigens induced a robust humoral immunity, but they have failed so far [6,7]. Thus, there is growing evidence that S. aureus vaccine strategies aimed at eliciting both appropriate long-lived B- and T-cell responses should be spotlighted [5,8,9]. Therefore, a more comprehensive characterization of T-cell immunity [9] is pivotal to characterize the immune response induced by the proteins mentioned above and to analyze in the context of protective immunity.

T-cell immunity has been increasingly reckoned as a central player in host-pathogen interaction, and T cell-derived interleukin (IL)-17 as a cornerstone of protection in S. aureus infections [8,10–13]. In this regard, the use of a DNA vaccine containing Granulocyte Macrophage Colony-Stimulating Factor (GM-CSF) appears as a powerful immunoadjuvant as it increases the lymphoproliferative response and IL-17 cytokine production [14]. Furthermore, the administration of a recombinant human GM-CSF alone was seen to reduce the rate of new IMIs by 46.7% following a challenge with S. aureus [15].

Aiming for an effective vaccine for S. aureus bovine IMIs, in the present study, we propose evaluating the immune response profile triggered by three recombinant S. aureus vaccine candidate antigens (SAS, SDD, and CTS) in association with GM-CSF DNA vaccine in a mouse model.

2. Materials and Methods

2.1. Ethical Statement

This study was approved by the Animal Research Ethics Committee of the Universidade de São Paulo, Brazil, under protocol number 6276100519.
2.2. Production of S. Aureus Recombinant Proteins

The sequence of amino acids corresponding to the SAS (Genbank gi|446897629), SDD (Genbank gi|486621908), and CTS (Genbank gi|257275504) S. aureus antigens were codon-optimized for Escherichia coli expression. Genes were commercially synthesized by Genscript, USA. The produced synthetic genes were cloned into the pUC57 vector and then sub-cloned into the pET28a expression vector [16]. Recombinant plasmids were used to transform the expression strain E. coli BL21-Start™(DE3) as previously described [17] with modifications. Briefly, transformed E. coli BL21-Start™(DE3) were selected on kanamycin plates. An overnight bacterial culture of three colonies containing the respective expression plasmid was grown in Luria-Bertani medium (LB) in the presence of kanamycin (0.05 mg mL⁻¹) until an optical density (OD) of 0.4 at 600 nm, on a rotary shaker at 37 °C. Then, isopropyl-β-D-1-thiogalactopyranoside (IPTG, Sigma) was added to the culture to a final concentration of 0.4 mM, and the induced cultures were grown for 4 h. Cells were ruptured by ultrasound sonication on ice and debris removed by centrifugation (20,000 x g, at 4°C, for 30 minutes). The recombinant proteins were purified using Ni Sepharose High Performance immobilized metal ion affinity chromatography columns (HisTrap Hp, cat. n. GE17-5248-02, GE Healthcare, Logan, USA), attached to an ÄKTA Pure (GE Healthcare, USA), under denaturing conditions according to the manufacturer’s instructions.

2.3. Production of GM-CSF DNA Vaccine

The amino acid sequence of the GM-CSF (Genbank, gi|145301581) was forwarded to FastBio (Ribeirão Preto, Brazil) for optimization for eukaryotic cells and to synthesize the respective gene. The produced synthetic genes were cloned in pUC57 vector and then were sub-cloned into the pCI-neo mammalian expression vector insert (Promega Incorporation, USA). E. coli DH5α was transformed with pCI-GM-CSF, and immunization plasmids were purified using the ZR plasmid Gigaprep Kit (Zymo Research, USA), according to the manufacturer’s instructions.

2.4. Liposome Preparation and Entrapment of Plasmid DNA

We used a cationic liposome to efficiently deliver the GM-CSF DNA plasmid DNA, which has been associated with induces strong humoral and cell-mediated responses [18]. Briefly, liposome were prepared using dimethyl-di-octadecyl-ammonium (DDAB), cholesterol (molar ratio 1:4) and 1,2-distearoyl-sn-glycero-3-phosphoethanolamine-N-[amino(polyethylene glycol)-2000 (DSPE + PEG2000), 5% of total lipids used. First, 1 mL of chloroform was used to dissolve this lipid solution and then left under a constant N2 flow to evaporate chloroform and develop a phospholipid film on the tube walls. This film was then retained under vacuum for at least one h to eliminate any remaining chloroform. The film was rehydrated in 5 mM Tris-HCl (pH 7.5) at 60 °C for one h and vigorously stirred every 10 min. The obtained opaque solution was then exposed to sonication at high energy until the solution became almost fully transparent. Then, the solution was centrifuged for one h at 100,000 x g. Any residual pellet was removed, and the supernatant containing unilamellar lipid vesicles was used to form the liposome. The genetic material was then added to the liposomes in a molar stoichiometry of 8nM DDAB for each 1 µg of nucleic acids, the molarity being approximately 0.2 pmol.

2.5. Animals and Immunization

Twenty-four C57BL/6J mice (six-week-old, female) were purchased from the Centro de Bioterismo of Faculdade de Medicina, Universidade de São Paulo. After three weeks of adaptation, the animals (nine weeks old) were divided into four groups (six animals per group) and immunized as defined in Figure 1. Here, 50 µg of pCI-GM-CSF plasmid DNA resuspended in 100 µL of sterile liposomal formulation were administered subcutaneously under the skin of the interscapular area near the draining lymph nodes. The recombinant SAS, SDD, and CTS proteins (20 µg each protein) in 100 µL of saponin adjuvant
(Quil-A®, cat. n. 8047-15-2, Invitrogen, San Diego, California, USA) were administered intramuscularly in the deltoid muscles (Figure 1).

A)

![Timeline of vaccination](image)

**Figure 1. Scheme of the timeline of vaccination (A) and experimental groups (B).** SAS: F0F1 ATP synthase subunit alpha recombinant *S. aureus* protein; SDD: succinyl-diaminopimelate desuccinylase recombinant *S. aureus* protein; CTS: cysteiny1-tRNA synthetase recombinant *S. aureus* protein; GM-CSF: pCI-granulocyte and macrophage colony-stimulating factor plasmid DNA (GM-CSF DNA vaccine). The unvaccinated group just received the liposome or saponin adjuvant.

On day 44 [14 days after the last administration of the recombinant protein(s)], the animals were anesthetized intramuscularly with xylazine (10 mg kg⁻¹) and ketamine (50 mg kg⁻¹) solution and euthanized by cervical dislocation. Then the draining lymph nodes (accessory axillary and proper axillary) and spleen were removed to obtain the cells for further immunophenotyping and lymphocyte proliferation evaluation, respectively. The accessory axillary and proper axillary lymph nodes from the side of vaccination were pooled in each animal.

**B)**

<table>
<thead>
<tr>
<th>VACCINE</th>
<th>G1</th>
<th>G2</th>
<th>G3</th>
<th>G4</th>
</tr>
</thead>
<tbody>
<tr>
<td>SAS+SDD+CTS</td>
<td>-</td>
<td>-</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>GM-CSF</td>
<td>-</td>
<td>+</td>
<td>-</td>
<td>+</td>
</tr>
</tbody>
</table>

2.6. Obtaining the Spleen and Lymph Nodes Cells

The spleens or lymph nodes of donor mice were aseptically removed, and mechanically disrupted, and homogenized by syringe plunger into cell strainer (cat. n. Z742102, Sigma Aldrich, St. Louis, USA) on the top of a 50 mL tube with 10 mL of RPMI 1640 medium (cat. n. R7638, Sigma Aldrich, St. Louis, USA) supplemented with 5% heat-inactivated fetal bovine serum (Gibco, USA), 100 U mL⁻¹ penicillin, 100 μg mL⁻¹ streptomycin, and Fungizone 0.25 μg mL⁻¹ (cat. n. 15240-096, Gibco) (complete medium).

The spleen and lymph node cells were centrifuged at 380 x g, 10 min, at 4 °C. Thereafter, the hypotonic lysis of erythrocytes was performed by adding 1,000 μL of 0.2% NaCl
for 20 s; then, isotonicity was restored by adding 1,000 µL of 1.6% NaCl. Cells were further centrifugated and suspended in a complete medium. Cell viability was checked with the trypan blue exclusion test (cat. n. T8154-100ML, Sigma Aldrich, St. Louis, USA).

2.7. Immunophenotyping of ex-vivo Lymph Nodes Cells

The lymph node cells were phenotyped using 0.5 µL of each of the fluorescent-conjugated monoclonal antibodies (mAbs; Table 1) for cell surface CD4 (clone GK1.5), CD8 (clone 53-6.7), CD19 (clone 1D3), γδ TCR (clone GL3), TCRVγ4 (clone GI2), CD44 (clone IM7) and CD27 (clone LG.3A10) by incubating the cells for 30 min at room temperature in the dark. After the incubation period, the cells were washed with 0.01 M phosphate-buffered saline, pH 7.4 (PBS), and spun at 250 x g at 4°C for 8 minutes. Further, the intracellular production of IL-17A and interferon (IFN-γ) were determined fixing and permeabilizing the cells as previously described [19], staining with 0.5 µL of fluorescein-conjugated anti-mouse IL-17 and anti-mouse IFN-γ antibodies (Table 1), respectively, and incubating for one hour at 4 ºC. After the incubation period, 500 µL of the permeabilization solution (PBS + 0.1% saponin + 0.09 azide + 1% heat-inactivated fetal bovine serum) was added and the samples centrifuged at 250 x g at 4°C for 8 minutes. Then, the samples were resuspended in 300 µL PBS with 1% SFB and analyzed by flow cytometry (BD LSRFortessa™ X-20 flow cytometer, Becton Dickinson Immunocytometry System™, San Diego, USA). Here, 100,000 events were acquired per sample. FlowJo Tree Star software (FlowJo - Treestar 10.5.3 for Windows, Tree Star Inc., Ashland, OR, USA) was used to analyze the data. Non-stained control and single-stained samples as compensation controls, negative control stained with conjugated isotype control antibody, and cells stained with fluorescence minus-one (FMO) controls were included. Doublets were excluded using forward scatter (FSC) area versus FSC height.

2.8. Preparation of S. aureus Inoculum

An udder adapted S. aureus isolate originated from a case of chronic subclinical IMI was used [17]. *Staphylococcus aureus* (spa typing t605) inoculum was prepared as previously described [20] with minor modifications. Bacteria were resuspended in RPMI-1640 (cat. n. R7638, Sigma Aldrich, St. Louis, USA) with 10% heat-inactivated fetal bovine serum (Cultilab, Campinas, Brazil). To determine the bacteria concentration in the inoculum, an aliquot of the bacterial suspension was further cultured on BH1 agar plates in dilution series, and colony numbers (colony-forming units mL⁻¹) were determined. Then, the bacterial concentration was adjusted to the final inoculum dose (2 x 10⁸ staphylococci mL⁻¹) to obtain a multiplicity of infection (MOI) = 10. The inoculum was heat-inactivated at 60 ºC for one hour, after which 100 µL of the solution was plated on a plate containing blood agar and incubated at 37°C for 24 h to confirm inactivation.

2.9. Immunophenotyping and Lymphocyte Proliferation of Cultured Spleen Cells

For lymphocyte activation of spleen cells, 2 x 10⁸ viable cells per well were cultured at 37 ºC and 5% CO₂ for 96 h in 96-wells flat-bottom plates with or without (control) 10 µL of heat-inactivated *S. aureus* (2 x 10⁸ CFU mL⁻¹). After the incubation period, the cells were harvested from the 96-well plates and transferred to 5 mL tubes, round bottom, 12 x 75 mm, for flow cytometry, and centrifuged at 250 x g at 4°C for 8 minutes. Then, the supernatant was collected and stored for cytokines measurement, and the cells were phenotyped using fluorescent-conjugated mAbs (Table 1).

Furthermore, the intracellular production of interleukin (IL)-17A and interferon (IFN)-γ, in addition to the antigen-specific in vitro lymphocyte proliferation marker [19], were determined. Ki67 is a nuclear protein that acts in the regulation of the cell division process. This protein is expressed during all active phases of cell division; thus, it has been widely used to evaluate cell proliferation [19]. After centrifugation, the cells were fixed, and further permeabilized [19]. Afterward, 0.5 µL of fluorescent-conjugated anti-mouse IL-17A and IFN-γ (Table 1) and 10 µL of diluted (2 µL of ki67 diluted in 198 µL of
permeabilization solution) rabbit anti-Ki67 antibody solution (cat n. ab15580, abcam, Cambridge, UK) was added and incubated for one hour at 4 °C. After the incubation period, 500 μL of the permeabilization solution was added, and the samples were centrifuged at 250 x g at 4°C for 8 minutes. After the incubation period, 500 μL of the permeabilization solution was added, and the samples were centrifuged at 250 x g at 4°C for 8 minutes. After centrifugation, the samples were resuspended in 300 μL of PBS with 1% SFB and analyzed by flow cytometry (BD LSRFortessa™ X-20 flow cytometer, Becton Dickinson Immuno cytometry System™, San Diego, USA). For this assay, 100,000 events were examined in each sample. Flow Jo Tree Star software (FlowJo - Treestar 10.5.3 for Windows, Tree Star Inc., Ashland, OR, USA) was used to analyze the data. Non-stained control and single-stained samples were prepared as compensation controls. Negative control samples were stained with conjugated isotype control antibodies. In addition, cells were stained with fluorescence minus-one (FMO) controls. Doublets were excluded using forward scatter (FSC) area versus FSC height.

Table 1. Monoclonal antibodies (mAbs) used for immunophenotyping of splenocytes and lymph nodes lymphocytes by flow cytometry.

<table>
<thead>
<tr>
<th>mAbs</th>
<th>Fluorescent probes</th>
<th>Target</th>
<th>Clone</th>
<th>Host</th>
<th>Concentration (mg mL⁻¹)</th>
<th>Cat. n.</th>
</tr>
</thead>
<tbody>
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<td>APC-Cy7</td>
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<td>GK1.5</td>
<td>Rat</td>
<td>0.2</td>
<td>552051</td>
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<tr>
<td>Anti-CD8</td>
<td>BV510</td>
<td>Mouse</td>
<td>53-6.7</td>
<td>Rat</td>
<td>0.2</td>
<td>563068</td>
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<tr>
<td>Anti-CD44</td>
<td>BV421</td>
<td>Mouse</td>
<td>IM7</td>
<td>Rat</td>
<td>0.2</td>
<td>563970</td>
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<tr>
<td>Anti-CD27</td>
<td>BV750</td>
<td>Mouse</td>
<td>LG.3A10</td>
<td>Hamster</td>
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<td>747399</td>
</tr>
<tr>
<td>Anti-CD19</td>
<td>PE-Cy7</td>
<td>Mouse</td>
<td>1D3</td>
<td>Hamster</td>
<td>0.2</td>
<td>552854</td>
</tr>
<tr>
<td>Anti-γδ TCR</td>
<td>BV650</td>
<td>Mouse</td>
<td>GL3</td>
<td>Hamster</td>
<td>0.2</td>
<td>563993</td>
</tr>
<tr>
<td>Anti-TCRVγ4</td>
<td>FITC</td>
<td>Mouse</td>
<td>GL2</td>
<td>Hamster</td>
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<tr>
<td>Anti-IL-17A</td>
<td>Alexa 700</td>
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<tr>
<td>Anti-IFN-γ</td>
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<td>Mouse</td>
<td>XM1G1.2</td>
<td>Rat</td>
<td>0.2</td>
<td>554412</td>
</tr>
</tbody>
</table>

1BD Pharmingen™ (San Diego, EUA); IL-17A: interleukin-17A; IFN-γ: interferon-γ; FITC: Fluorescein isothiocyanate; APC-Cy7: Allophycocyanin-cyanine 7; PE: Phycoerythrin; PE-Cy7: R-Phycoerythrin-cyanine 7; BV510: Brilliant Violet 510; BV421: Brilliant Violet 421; BV650: Brilliant Violet 650; BV750: Brilliant Violet 750.

2.10. Cytokines Measurement

The production of cytokines IL-2, IL-4, IL-6, IL-10, IFN-γ, IL-17A, and tumor necrosis factor (TNF)-α was determined in the supernatant of the cultured spleen cells with or without S. aureus stimulation using the BD Cytometric Bead Array Mouse Tpm1, Tn1 and Tn17 Cytokine (cat. n. 560485, BD Bioscience™, San Jose, USA) using a flow cytometer (BD LSRFortessa™ X-20 flow cytometer, Becton Dickinson Immuno cytometry System™, San Diego, USA), as manufacturer’s instructions. The FCAP Array™ v3.0 software (Softflow™, Pécs, Hungary) was used to analyze the data.

2.11. Statistical Analysis

Statistical analysis was performed using GraphPad Prism 9 (GraphPad Software, Inc., San Diego, USA). Data are presented as percentages of distinct T cell populations that produced IL-17A or IFN-γ (percentage of stained cells) and geometric mean fluorescence intensity (GMFI) that provides an accurate measurement of the brightness of stained cells. The GMFI of 17A or IFN-γ was determined among IL-17+ or IFN-γ+, indicating the intensity of cytokine production per cell that produced the respective cytokine. To assess the percentage of each lymphocyte subpopulation, the percentage of proliferative cells (Ki67+), as well as the percentage and the GMFI of IL-17A or IFN-γ, a stimulation index (SI), was calculated by dividing the percentage of positive cells upon S. aureus stimulation per percentage of positive cells under unstimulated control condition [21–25]. The data were first tested for normality of the distribution using the Kolmogorov-Smirnov and Shapiro-Wilk
tests. If the distribution were normal, the data of the experimental groups were subjected to a one-way ANOVA analysis followed by the Tukey test. Variables with non-parametric distributions were analyzed using the Kruskal-Wallis test followed by the Dunn’s test. Results are reported as mean + standard error of the mean. \( P \leq 0.05 \) was considered significant unless otherwise indicated.

3. Results

3.1. Immunization with S. Aureus Recombinant Proteins Generate Memory Cells in Draining Lymph Nodes

CD44 is a prominent activation marker that differentiates memory and effector T cells from their naïve counterparts [26]. A higher percentage of CD44+ cells in draining lymph nodes was found in animals that just received S. aureus recombinant proteins (data not shown). The memory T cells were segregated into two distinct populations: a CD44high CD62Llow that exert a rapid effector function (so-called effector memory, TEM), which have the poor proliferative capacity, and a CD44high CD62Lhigh population with no immediate effector function (central memory, TCM) that possess high proliferative potential [27,28]. Those animals that only received the combination of S. aureus recombinant proteins showed an enhanced percentage of T CD4+ CD44+ CD27+, i.e., TCM T CD4+ cells (Figure 2 A and B), but a reduced percentage of T CD8+ CD44+ CD27+, i.e., effector T CD8+ cells (Figure 2 B).

![Figure 2](image-url)

**Figure 2.** The antidromic trend of the percentage of T CD4+ CD44+ CD27+ cells and T CD8+ CD44+ CD27+ cells in draining lymph nodes from mice vaccinated with three-recombinant S. aureus proteins. A) A representative gating hierarchy (successive gating scheme as depicted by arrows): lymphocytes (gate I), single cells are identified by plotting forward scatter area against forward scatter-height (gate II), single lymphocyte cells (gate III), T CD4+ cells (gate IV), FMO controls for CD44 (gate V) and CD27 (gate VI), and representative plots used to discriminate and identify two distinct T CD4+ memory cell populations: a CD44high CD27low population, and a CD44high CD27high population unstimulated (gate VII) and stimulated with S. aureus (gate VIII). B) Percentage of T CD4+ CD44+ CD27+ cells and T CD8+ CD44+ CD27+ cells in draining lymph nodes among groups. FMO: fluorescence minus one; F0F1 ATP synthase subunit α (SAS), succinyl-diaminopimelate
3.2. TCRVγ4⁺ Cells are Responsible for the Greatest Contribution for Type 1 and 3 Cell-Mediated Immunities in the Draining Lymph Nodes

An emerging concept is that unconventional T cells, such as γδ T cells, may confer protective immunity against *S. aureus* [13,29]. Here, we observed that γδ TCR cells and its subpopulation TCRVγ4⁺ cells in the studied draining lymph nodes were associated with the highest percentage of IL-17A⁺ cells in the draining lymph nodes (*P* ≤ 0.0001; Figure 3), highlighting the critical importance of this lymphocyte population in triggering type 3 cell-mediated immunity [13,30]. Furthermore, the TCRVγ4⁺ cell population exhibited a higher percentage of IFN-γ⁺ cells than all αβ T lymphocytes subpopulations, while γδ TCR⁺ cells just differed from T CD8⁺ lymphocytes (Figure 3).

![Figure 3](image-url)

**Figure 3.** Cellular sources of IFN-γ and IL-17A in draining lymphocyte populations upon in vitro *S. aureus* stimulation. The γδ TCR cells and its subpopulation TCRVγ4⁺ cells are the primary sources of both IL-17A and IFN-γ in draining lymph nodes rather than αβ T cells. *P* < 0.05; **P* < 0.01; **** *P* < 0.0001 (Kruskal-Wallis test followed by Dunn’s test).

3.3. Immunization with *S. Aureus* Recombinant Proteins Associated with GM-CSF DNA Vaccine Drives Type 3 Recall Immunity that Largely Relies on γδ T and TCRVγ4⁺ Cells

There is increasing evidence that type 3 immunity mediated by cells that produce IL17-A and IL17-F is crucial for mammary gland protective immunity [7], consequently a promising target for vaccine development. Observing that the three *S. aureus* recombinant antigens plus GM-CSF DNA vaccine induced appropriate immune response at the draining lymph node, we proceeded to evaluate the systemic recall immune response using splenic cells from immunized mice stimulated in vitro with *S. aureus*. No effect of immunization on the induction of B cell response was revealed (data not shown). To quantify the intensity of response upon *S. aureus* stimulation, T cell responses are normalized by dividing the percentage of positive cells upon *S. aureus* stimulation by the percentage of positive cells under unstimulated control conditions, thereby creating a stimulation index. Thus, when in vitro cultures of spleen cells were stimulated with *S. aureus* from vaccinated mice with the three recombinant proteins plus the GM-CSF DNA vaccine, an increase in the percentage of IL-17A⁺ cells among overall CD44⁺ (memory) cells (35.34 %), T CD8⁺ cells (54.36 %), CD4⁺ TEM (CD44⁺ CD27⁻) cells (142.87 %), and TCRVγ4⁺ cells (56.75 %) were found (Figure 4). This effect was due almost to no significant impact on the IL-17A⁺ among the distinct T cells subsets in spleen cell culture under *S. aureus* stimulus compared with unstimulated conditions in animals from unvaccinated, GM-CSF DNA vaccine and those vaccinated with the *S. aureus* recombinant proteins alone (*P* ≤ 0.05), while mice immunized with the three-antigens *S. aureus* recombinant proteins associated with the GM-CSF DNA vaccine generated a pronounced enhancement in the percentage of IL-17A⁺ cells in all...
abovementioned T cell populations, primarily on γδ T cells and TCRVγ4+ cells. Nevertheless, no vaccine effect on GMFI of IL-17A production and type 1 (IFN-γ) immune cells was observed.

Figure 4. IL-17A+ T cells in mice immunized with three S. aureus recombinant proteins associated with the GM-CSF DNA vaccine. Data presented as stimulation index, normalizing T cell responses, dividing the percentage of IL-17A+ cells upon S. aureus stimulation by the percentage of IL-17A+ cells under unstimulated control condition. A) CD44+ (memory) cells, B) T CD4+ cells, C) T CD4+ TEM (CD44+ CD27−) cells, D) γδ TCR cells, E) γδ TCR TCM, F) TCRVγ4+ cells. * P < 0.05; ** P < 0.01; *** P < 0.001 [One-way ANOVA followed by Tukey test (Figures A, D and F) and Kruskal-Wallis test followed by Dunn’s test (Figures B, C and E)].

3.4. TCRVγ4+ Cells and T CD8+ Cells are the Major Contributors for Type 1 Immunity, while γδ TCR Cells Mainly Support Type 3 Immunity in Splenocyte Cell Culture

Here, we observed both TCRVγ4+ cells and T CD8+ cells are the primary source of both IFN-γ, followed by γδ TCR cells (Figure 5). On the other hand, γδ TCR cells represent the major producer of IL-17A, followed by TCRVγ4+ cells and T CD8+ cells (Figure 4). The T CD4+ cells marginally produce IFN-γ and IL-17A, although the major population found splenocytes cell culture in both unstimulated and S. aureus stimulated conditions (Figure 5).
**Figure 5.** Cellular sources of IFN-γ and IL-17A in splenocyte populations upon *in vitro S. aureus* stimulation. The TCRVγ4+ cells and T CD8+ cells are the primary sources of IFN-γ, while γδ TCR is the major source of IL-17A, under splenocyte cell culture in unstimulated and *S. aureus* stimulated conditions. *P* < 0.05; **P** < 0.01; ***P** < 0.001; ****P** < 0.0001 [Kruskal-Wallis test followed by the Dunn’s test (A, B and D) and One-way ANOVA followed by Tukey test (D)].

### 3.5. Immunization of Mice with *S. Aureus* Recombinant Proteins Associated with GM-CSF DNA Vaccine Downregulates Th2 Immunity

The cytokine measurements in the supernatant of the unstimulated (control) and *S. aureus* stimulated cultured spleen cells are shown in Supplementary Figure 1. In the present study, the IL-4 concentration in the supernatant of the splenocytes culture upon *S. aureus* stimulation was lower in animals vaccinated with the GM-CSF DNA vaccine (*P* = 0.08), those immunized with the three-recombinant antigens (*P* = 0.03) and immunized with the three-recombinant antigens together with the GM-CSF vaccine (*P* = 0.06) compared with unvaccinated control animals (*P* = 0.50). Moreover, the IL-4 concentration in the supernatant of the splenocytes culture upon *S. aureus* was higher in the non-immunized control mice than in those animals that received the three *S. aureus* recombinant proteins alone (*P* = 0.01).

No difference in the concentration of TNF-α and IL-6 in the supernatant of the cell culture among groups and between unstimulated and *S. aureus* stimulated conditions were detected. The concentrations of IL-10, IL-17A, and IFN-γ were below the limit of the detection. In addition, *S. aureus* stimulation leads to a decrease in the concentration of IL-2 in the supernatant of splenocytes culture only in mice that received the GM-CSF DNA vaccine together with the recombinant proteins (*P* = 0.05).
3.6. The Combination of S. aureus Recombinant Proteins and GM-CSF Vaccine Induces a Higher Proliferation of Memory T Cells and T CD4+ Cells

In the present study, when we compared the percentage of proliferative cells between S. aureus stimulated and unstimulated conditions within groups, we observed that animals that received the S. aureus recombinant proteins plus GM-CSF DNA vaccine lead to an increase in the percentage of proliferative (ki67+) overall memory T cells (CD44+: P = 0.02), CD4+ cells (P = 0.01) and CD4+ TEM (CD44+ CD27; P = 0.02) when stimulated with S. aureus that was not sustained in other experimental groups.

4. Discussion

Vaccination strategies may provide memory T cells to fight developing pathogens in the host and respond quickly upon reencounter with the pathogen [28,30–32]. Within this context, a hallmark finding of our study is the high production of IL-17A by diverse T cell memory cell subsets using the combination of three S. aureus recombinant proteins and GM-CSF DNA vaccination when stimulated with S. aureus. Further, we must highlight the increased production of IL-17A by the TCRγδ4+ cell population when stimulated with S. aureus. In the present study, we demonstrated for the first time that the combination of S. aureus CTS, SDD, and SAS recombinant proteins associated with a GM-CSF DNA plasmid DNA vaccine foster type 3 cell-mediated immunity that turns these outcomes into a promising added value.

We should emphasize the strategy to identify three recombinant proteins used in the present study using the serum immunoproteomics approach in the previous study [2]. In this latter study, we searched the antibodies produced by dairy cows exposed to S. aureus but remained healthy, hypothesizing that the antibodies repertoire directed against these specific S. aureus proteins contributed to mastitis resistance. The first S. aureus protein, so-called SAS protein, plays a pivotal role in dictating biofilm growth and structure [33], enhances tolerance of S. aureus towards some host antimicrobial peptides of the innate immunity [34], inhibits pro-inflammatory cytokines production by monocyte-derived macrophages and myeloid-derived suppressor cells and renders S. aureus more resistant to the bactericidal activity of monocyte-derived macrophages [33]. The second S. aureus SDD protein is critical for bacteria survival and proliferation. It is involved in protein synthesis and constructing the peptidoglycans of the bacterial wall [35]. Lastly, S. aureus CTS is an aminocly-tRNA enzyme that catalyzes the binding of a specific amino acid to tRNA, attaches the amino acid to other molecules, and is therefore involved in protein synthesis [35]. Altogether, although molecular functions for these proteins are not exactly known, we consider it reasonable to speculate that stimulating the host immune response against these proteins is the precise strategy to protect animals from S. aureus infections.

In this scenario, although most available vaccines’ success relies on antibody-mediated immunity, it is conceivable that humoral immunity alone may not be sufficient to protect against S. aureus infections [5,6] fully. By the way, protective immunity against recurrent S. aureus skin and soft tissue infection in a human was associated with both antibody and T-cell mediated (i.e., IL-17) immunities, suggesting that a multimechanistic approach targeting both humoral and T cell-mediated immunities may be a key strategy to provide protective immunity and prevent new S. aureus infections [5]. In this line, we identified that animals vaccinated with these S. aureus proteins associated with a GM-CSF DNA vaccine trigger type 3 cell-mediated immunity reinforcing their potential use to prevent new IMIs by this pathogen.

It is notorious that T cells make an essential contribution toward S. aureus control [36-38]. In fact, T-cell deficiencies increase the susceptibility to S. aureus infections [6,37]. Astonishingly, the T cell responses in developing an effective vaccine for bovine S. aureus mastitis have not received much attention, especially the unconventional T cells neglected. In this setting, γδ T cells represent a largely ignored target for vaccine design. However, it comprises up to 60% of circulating lymphocytes in young cattle and up to 30% of blood mononuclear cells in adult animals [39]. The critical role of this T cell population in cattle
could be envisaged, seen that WC1⁺γδ T cells are the first to arrive at the sites of mycobacterial purified protein derivate (PPD) injection and increase in the frequency in blood following mycobacterial vaccination [39,40]. Actually, γδ T cells present cytotoxic activity in cows, and a decrease of this cell population is associated with periods of increased susceptibility to IMIs [41].

Therefore, there is an increasing interest in studying the T-cell response to S. aureus infections. Different subpopulations of T cells are likely to contribute to anti-staphylococcal immune defense. Among them, recent evidence has highlighted the importance of type 3 immunity for protective immunity against infections by S. aureus. Therefore, they emphasize their importance in developing new vaccines [10,12,42]. Type 3 immunity can be characterized by cells that encode IL-17A, IL-17F, IL-22 genes and the Rorγt and Rora transcription factors and their gene targets. The cells responsible for type 3 immunity are diverse, including γδ T cells, T CD4⁺ helper (Th17), T CD8⁺ (Tc17), and innate lymphoid cells 3 [7].

Among T-cells producing IL-17, the nontraditional γδ T cells have been identified as a potent source of innate IL-17 and implicated in host protection in murine models of S. aureus infection. These findings reveal that γδ T cells are an important source of IL-17 in adaptive immunity and indicate that targeting the induction of nontraditional lymphocytes as specific γδ T cell subsets that secrete IL-17 represent a potentially important and novel target for the rational design of future vaccines against S. aureus [6,13]. The γδ T cells are known to substantially contribute to protective immunity against S. aureus infections involving innate and adaptive immune responses, as they act as the first line of defense and control the innate response by neutrophil-mediated regulation [41,49]. It has also been proposed that γδ T-lymphocytes play a crucial role in antibacterial immunity and may provide a unique barrier for mucosal microenvironments against bacterial pathogens [50]. Furthermore, it was recently shown that an increase in the subpopulation of γδ T cells (i.e., TCRVγ4⁺ cells) is essential for protection against subsequent infections by S. aureus [13,51,52]. Therefore, this T γδ lymphocyte subpopulation represents an important participant in the protective vaccine against S. aureus infections [12,13,51,53]. In fact, a central finding of the present study is the higher percentage of IL-17A⁺ TCRVγ4⁺ cells upon stimulation with S. aureus, which strongly indicates a pivotal protective role of our vaccination approach against S. aureus challenge infections. Further studies in vaccinated animals under intramammary challenge with S. aureus are certainly needed to confirm this assumption.

Interleukin-17 is associated with neutrophil mobilization, immune response modulation, and antigen-specific inflammation [13,43] and mediating the communication between the immune system and mammary epithelial cells [44]. For example, epithelial cells may express receptors for IL-17 and IL-22. The exposure to IL-17 leads to the release of chemokines and granulocyte-macrophage colony-stimulating factor (GM-CSF), leading to neutrophil recruitment and activation leading to the death of S. aureus [10,42,45]. The GM-CSF, in its turn, activates the antigen-presenting cells, such as macrophages and dendritic cells, increasing major histocompatibility complex (MHC) expression, increasing its antigen-presenting capacity, and amplifying the primary antibody response. GM-CSF also stimulates the maturation of dendritic cells and their consequent migration towards regional lymph nodes. GM-CSF may therefore induce differentiation, proliferation, and activation of several cell types, facilitating the development of both humoral and cellular immunity [46]. We may thus conclude that GM-CSF is an important mediator of the immune response in the mammary gland of dairy cows [15,44,47,48], corroborating the importance of our findings and the use of GM-CSF DNA vaccination to improve vaccine efficacy against S. aureus.

The importance of γδ T cells is because they are the primary source of interleukin-17 (IL-17) while other non-γδ T cells just provide a small contribution for the production of IL-17 during S. aureus infections [13,29,51,52,54,55]. In the same way, our outcomes go in the same direction with γδ T cells, including its subpopulation TCRVγ4⁺ cells having a critical role in IL-17A production, while αβ T cells just marginally producing IL-17A in
the regional draining lymph nodes. Besides γδ T cells that a pivotal contribution for IL-17A production by splenocytes with or without S. aureus stimulus in cell culture, Tc17 contributed to type 3 cell-mediated immunity. Furthermore, γδ T cells, particularly TCRVγ4+ cells, robustly support IFN-γ production in the studies draining lymph nodes. In contrast, both γδ T cells and Tc1 (T CD8+) cells markedly contribute to the IFN-γ when recall immune response was evaluated. The production of IFN-γ by γδ T cells, including TCRVγ4+ cells, has been previously demonstrated [53,56,57]. Overall, these findings support the substantial and broad role of this nonconventional T cell population in immunity against S. aureus.

Furthermore, we observed that the vaccination with the three recombinant proteins associated with the GM-CSF DNA vaccine downregulates Tn2 immunity, observing decreased IL-4. Of note, a type 2 immune environment can promote S. aureus colonization and superinfection [29]. We have also found lower levels of IL-2 in the supernatant of splenocytes cell culture in animals that received the combination of S. aureus proteins together with the GM-CSF. Activation of T cells with IL-2 induces TEM cells while T cells stimulated in vitro with IL-15 acquire a TCM phenotype [58]. However, as IL-15 was not determined, the latter remains speculative but somehow corroborates with the formation of the TCM phenotype found here. However, although the mechanism is still under debate, it should be noted immediately after pathogen clearance that the antigen-specific memory T-cell pool comprises mainly TEM cells, which gradually convert to a TCM cell phenotype over time [59]. In this perspective, TCM cells are regarded as a renewable source of T effector cells responsible for protection from acute infections and primed for a rapid effect response [60].

5. Conclusions

In the present study, the immunization with the three recombinant proteins associated with GM-CSF DNA improved T-cell memory response, especially γδ TCRγδ and its subpopulation TCRVγ4+, which are the major responsible for the production of IL-17A. Thus, as our vaccination approach fosters type 3 immunity, a protective immune response against S. aureus was expected. Therefore, although our promising outcomes, further studies in S. aureus challenged mice model and ruminants will be necessary to validate our findings regarding the protective immunity against S. aureus.

6. Patents

The authors have a patent titled “Vaccine composition, kit for diagnosis of Staphylococcus aureus infections in ruminants, method and uses”.


Funding: This work was supported by the Fundação de Amparo à Pesquisa do Estado de São Paulo – FAPESP (grant no 2018/23572-3), and by the Coordenação de Aperfeiçoamento de Pessoal de Nível Superior - Brasil (CAPES) (Finance Code 001). AMMPDL and MBH thank the National Council for Scientific and Technological Development (CNPq) for their fellowships. KRS thanks CAPES for her fellowship. FNS (Process FAPESP n. 2014/23189-4) thanks FAPESP for his fellowship.

Institutional Review Board Statement: The study was conducted according to the guidelines of the Declaration of Helsinki, and approved by the Animal Research Ethics Committee of the UNIVERSIDADE DE SÃO PAULO, Brazil (protocol number 6276100519; June, 23th, 2020).

Informed Consent Statement: Not applicable.

Data Availability Statement: The data presented in this study are available on reasonable request from the corresponding author.
Conflicts of Interest: The authors have a patent titled “Vaccine composition, kit for diagnosis of Staphylococcus aureus infections in ruminants, method and uses”. The funders had no role in the design of the study; in the collection, analyses, or interpretation of data; in the writing of the manuscript, or in the decision to publish the results.

References


