RESEARCH ARTICLE

A *Salmonella typhimurium* ghost vaccine induces cytokine expression *in vitro* and immune responses *in vivo* and protects rats against homologous and heterologous challenges

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Abstract

*Salmonella enteritidis* and *Salmonella typhimurium* are important food-borne bacterial pathogens, which are responsible for diarrhea and gastroenteritis in humans and animals. In this study, *S. typhimurium* bacterial ghost (STG) was generated based on minimum inhibitory concentration (MIC) of sodium hydroxide (NaOH). Experimental studies performed using *in vitro* and *in vivo* experimental model systems to characterize effects of STG as a vaccine candidate. When compared with murine macrophages (RAW 264.7) exposed to PBS buffer (98.1%), the macrophages exposed to formalin-killed inactivated cells (FKC), live wild-type bacterial cells and NaOH-induced STG at $1 \times 10^8$ CFU/mL showed 85.6%, 66.5% and 84.6% cell viability, respectively. It suggests that STG significantly reduces the cytotoxic effect of wild-type bacterial cells. Furthermore, STG is an excellent inducer for mRNAs of pro-inflammatory cytokine (TNF-α, IL-1β) and factor (iNOS), anti-inflammatory cytokine (IL-10) and dual activities (IL-6) in the stimulated macrophage cells. *In vivo*, STG vaccine induced humoral and cellular immune responses and protection against homologous and heterologous challenges in rats. Furthermore, the immunogenicity and protective efficacy of STG vaccine were compared with those of FKC and non-vaccinated PBS control groups. The vaccinated rats from STG group exhibited higher levels of serum IgG antibody responses, serum bactericidal antibodies, and CD4⁺ and CD8⁺ T-cell populations than those of the FKC and PBS control groups. Most importantly, after challenge with homologous and heterologous strains, the bacterial loads in the STG group were markedly lower than the FKC and PBS control groups. In conclusion, these findings suggest that the STG vaccine induces protective immunity against homologous and heterologous challenges.
Introduction

Salmonella are Gram-negative foodborne zoonotic bacteria that cause salmonellosis worldwide [1, 2]. S. typhimurium and S. enteritidis are mainly transmitted to humans by consumption of contaminated eggs and poultry meat products, and represent a global public health burden [3]. Salmonella species are composed of several serotypes that can cause infections in homologous vaccinated animals. Moreover, S. typhimurium strains are often resistant to multiple antibiotics, such as aminopenicillins, gentamicin, tetracycline, chloramphenicol, and sulfonamides [4–7]. Therefore, there is a need to develop Non-Typhoidal Salmonellae (NTS) vaccine that can induce both homologous and heterologous protective immunity against Salmonella strains. Meanwhile, several efforts with live attenuated vaccines have been made to induce cross-protection against NTS infections. Previous studies demonstrated that live attenuated Salmonella aroA [8–10], capable of reducing invasion and colonization of the gastrointestinal tract and protected animals from homologous challenge, but they lack the ability to induce heterologous protective immunity against a virulent challenge. Furthermore, the administration of attenuated aroA mutants from S. enteritidis [11] or S. typhimurium [12] has been shown to reduce fecal shedding against homologous challenge. However, vaccination with Salmonella adenylate cyclase (cya) and C-reactive protein (crp) mutant protected chickens from Salmonella serotypes challenges [13].

Bacterial ghosts (BG) are structurally intact empty bacterial cell envelopes prepared from various Gram-negative bacteria [14]. In BG preparation, bacterial cell lysis is induced by the controlled expression of the bacteriophage PhiX174 lysis gene E [15, 16] or minimum inhibitory concentration (MIC) of sodium hydroxide (NaOH) [17–21]. Based on this lysis, protein E or MIC of NaOH produces transmembrane tunnel structures on the cell surface that result in empty non-living cell envelopes. However, there are some differences in two methods. BG produced by the former method maintains their original structure, preserves all the cell surface antigens and provides efficient protection against specific infections [15–17]. Especially, functional and antigenic determinants of the envelope components are not denatured during lysis. However, there are some disadvantages of the method such as limitation to Gram-negative bacteria only, potential risks because of difficulty to reach 100% lysis rate of BG strain in a short time [22], and cost expensive and time consuming, multi-step process. Alternatively, BG produced by the latter method has no limitation to both Gram-positive [18] and Gram-negative bacteria [17,19]. This method needs a short time to generate BG without any potential risks, cheap cost and time-saving, simple process. Despite some surface structures in BG may be modified or lost by NaOH [19], the NaOH-induced BG provides efficient protection against specific infections [17,18].

In the last two decades, BG is representing to be the attractive vaccine candidate, because it can elicit both humoral and cellular immune responses against specific infections in experimental animals [23,24]. In previous studies, an effective S. enteritidis and S. gallinarum ghost vaccines were produced to protect animals from Salmonella infections [25,26]. In the present study, non-living S. typhimurium bacterial ghosts (STG) was successfully produced by using the chemically-induced method. The main purpose of this investigation was to assess in vitro cytotoxicity of STG and the influence of STG on mRNA induction of cytokine and nitric oxide synthase in murine macrophages and in vivo protection of STG against the homologous and heterologous challenge in a rat model. Moreover, immunogenicity and protective efficacy were compared between the STG vaccine and formalin-killed inactivated cells (FKC) vaccine. This is the first study on the cross-protective efficacy of the STG vaccine in rats.
Experimental procedures

Bacterial strain and culture condition

S. typhimurium KCCM40253 was kindly provided by Prof. Ki-Sung Lee, Department of Biology and Medicinal Science, Pai Chai University, Daejeon, Korea. S. enteritidis was provided from an animal health product manufacturing company (KBNP, Inc., Korea). The bacteria were grown at 37˚C in Luria-Bertani (LB) broth or on LB agar. Incubation temperature for growth and lysis was 37˚C in a shaking incubator at 200 rpm.

Production of S. typhimurium ghosts

STG were produced by using the MIC of NaOH as described previously [17–19]. Briefly, the MIC of NaOH against S. typhimurium was determined by the two-fold broth dilution method and 72 h cultured S. typhimurium cells were harvested by centrifugation for 10 min at 10,000 × g. After washing three times with PBS (pH 7.2), the bacterial suspension was adjusted to 1 × 10^8 CFU/mL. One mL of the MIC of NaOH was added to 2 mL of the bacterial suspension and incubated at 37˚C for 1 h. After incubation, STG were collected by centrifugation and washed three times with PBS. Samples were prepared and analyzed by scanning electron microscopy (SEM) as described previously [17,18]. For FKC vaccine preparation, 0.5% formalin was added to the bacterial suspension and incubated for 24 h at 37˚C. After incubation, the bacterial suspension was washed three times with PBS, and the bacterial suspension was plated on LB agar plates to confirm the total loss of viability. FKC was stored at 4˚C until use.

Assessment of macrophage-mediated cytotoxicity

Murine macrophage (KCLB:40071, RAW 264.7) cells were purchased from Korean Cell Line Bank (Seoul, Korea) and cultured in 96-well plates (BD Falcon; BD Bioscience Discovery Labware, Bedford, MA, USA) for 24 h at 37˚C, in humidified 5% CO₂, 95% air. The cells (1.0 × 10^4 cells/well) were then treated with 1.0 × 10^8 CFU/mL of the FKC, wild-type bacterial cells and NaOH-induced STG, respectively, in culture medium, and incubated for a further 24 h. PBS treated- and LPS-treated macrophages were used as negative and positive controls, respectively. LPS was extracted from S. typhimurium (1 × 10^8 CFU/mL) using an LPS extraction kit (iNtRON Biotechnology Inc., Seongnam-si, Gyeonggi-do, Korea) according to the manufacturer’s protocol. Briefly, 5 mL culture was centrifuged at 13,000 rpm for 5 min. The bacterial pellet was treated with 1 mL of supplied lysis buffer and vortexed vigorously. After adding 200 µL of chloroform, the mixture was centrifuged at 13,000 rpm for 10 min at 4˚C. Then, the supernatant (400 µL) was mixed well with a supplied purification buffer and incubated for 10 min at –20˚C. After centrifuging the mixture solution at 13,000 rpm for 15 min at 4˚C, the upper layer was removed to obtain the LPS pellet. The pellet was washed with 70% ethanol and centrifuged at 13,000 rpm for 3 min at 4˚C. After discarding the resulting upper layer, the pellet was dried at room temperature and dissolved in 10 mM Tris-HCl (pH 8.0) by boiling for 2 min. To get the pure LPS, proteinase K (2.5 µg/ LPS 1 µg) as added to the dissolved LPS and the mixture was incubated at 4˚C for 30 min. The cell density was then assessed by using Cell Counting Kit-8 (CCK-8, Sigma-Aldrich, St. Louis, MO, USA). Absorbance was measured at 450 nm and all experiments were performed in triplicate. Cytotoxic activity is expressed as the percentage of cell viability by the following formula: % Cytotoxicity = (1 – A450nm of target cells/A450nm of control cells) × 100.
Quantitative analysis of cytokine mRNA by reverse transcription (RT)-qPCR

RAW 264.7 cells (1.0 × 10⁴ cells/well) were cultured in 24-well flat-bottom plates and treated with the FKC, wild-type bacterial cells and NaOH-induced STG, respectively, at a concentration of 1.0 × 10⁸ CFU/mL. After 1–12 h stimulation, total RNA was isolated using RNAiso (Takara Bio, Shiga, Japan), according to the manufacturer’s instructions. Tumor necrosis factor (TNF)-α, interleukin (IL)-1β, IL-6, IL-10 and inducible nitric oxide synthase (iNOS) mRNA levels were quantified by RT-qPCR amplification. Sequences for the primers of target genes are listed in S1 Table. RT reaction was performed in a 20 μL reaction mixture containing 300 ng of total RNA, 50 mM Tris-HCl (pH 8.3), 75 mM KCl, 8 mM MgCl₂, 10 mM DTT, 0.1% NP-40, 40 mM dNTP, 2 pM of respective primer set, 20 U of RNase inhibitor (Takara Bio), and 200 U PrimeScript Reverse Transcriptase (Takara Bio). The thermal cycler was programmed for 1 RT cycle at 50°C for 30 min and 70°C for 15 min. cDNA was amplified in a 20 μL reaction mixture containing 10 μL 2X SYBR® Premix Ex Taq™ II (Tli RNaseH Plus, Takara Bio, Shiga, Japan), 0.2 μL ROX reference dye II, 0.4 μL of 10 μM of both forward and reverse primer and 1 ng of cDNA, using Stratagene Mx3005P cycler (1 cycle at 95°C for 30 s, 30 cycles of denaturation at 95°C for 5 s, and primer annealing and extension at 60°C for 34 s). Each gene was amplified in triplicate and cDNA concentration differences were normalized to glyceraldehyde 3-phosphate dehydrogenase (GAPDH).

Experimental animals

Ten-week-old male Sprague-Dawley rats were housed under standard conditions of temperature and relative humidity with a 12 h light/dark cycle. Food and water were available ad libitum. All animal experiments were approved by institutional animal care and use committee at Pai Chai University.

Vaccination and challenge protocol

Thirty-nine rats were equally divided into three groups: A, B, and C. The rats in group A (n = 13) were injected subcutaneously with sterile PBS as a non-vaccinated control. The rats in groups B (n = 13) and C (n = 13) were vaccinated subcutaneously with FKC (1 × 10⁸ cells/mL) and STG (1 × 10⁸ cells/mL), respectively. Rats from all groups were vaccinated three times at two-week intervals. Two weeks after the final vaccination (week 7), all rats were challenged orally with S. typhimurium (1 × 10⁸ CFU/mL) [27] or S. enteritidis (2 × 10⁶ CFU/mL) in PBS [17,28,29].

Detection of antibody levels by ELISA

Serum Immunoglobulin (Ig) G antibodies were determined by indirect enzyme-linked immunosorbent assay (ELISA) as previously described [18,19]. Microtiter plates were coated with 100 μL of S. typhimurium or S. enteritidis (1 × 10⁸ cells/mL) in coating buffer (pH 9.6). ELISA was carried out with sera from non-vaccinated and vaccinated rats. Goat anti-rat IgG conjugated alkaline phosphatase (1:30,000; Sigma-Aldrich) was used as the secondary antibody. The plates were developed with 100 μL of the substrate p-nitrophenyl phosphate (Sigma-Aldrich, St. Louis, MO) and 100 μL of 3N NaOH was added to stop the reaction. The optical density was measured at 405 nm with a microplate reader (Bio-Rad, USA).
Analysis of T-cells populations by FACS

To analyze T cell populations, blood samples were collected from non-vaccinated and vaccinated rats at one week after the final vaccination. The peripheral blood mononuclear cells were prepared using Histopaque-1077 (Sigma-Aldrich, St. Louis, MO) according to the manufacturer’s protocol. The cells \(1.0 \times 10^6\) were washed three times in PBS and incubated either fluorescein isothiocyanate (FITC) labeled anti-CD4 or CD8 monoclonal antibodies at 4˚C for 30 min in the dark. After the incubation, all the samples were washed twice with PBS, and analyzed on a BD FACSCanto II flow cytometer (BD Biosciences, NJ, USA). Data analysis was done on BD FACSDiva software (BD Biosciences).

Determination of serum bactericidal activity

The bactericidal activity of the sera was tested as described previously [30]. Twenty-five μL of serum were mixed with 100 μL of S. typhimurium \((1 \times 10^6 \text{ CFU/mL})\) or S. enteritidis \((1 \times 10^6 \text{ CFU/mL})\) and incubated for 1 h at room temperature. After incubation, samples were plated on selective media and incubated for 24–48 h at 37˚C. Instead of serum, PBS was served as a control. The percentage of serum bactericidal activity was determined by using the following formula: 

\[ \text{SBA} = \left(1 - \frac{\text{the number of viable bacteria after serum treatment}}{\text{the number of viable bacteria after PBS treatment}}\right) \times 100\%. \]

Western blot analysis

S. typhimurium or S. enteritidis envelope proteins were extracted as previously described [31,32]. Twenty μg of protein were loaded and subjected to 12\% Sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE). Western blots were probed with vaccinated or non-vaccinated sera (1:1,000) in non-fat dry milk as primary antibodies. Goat anti-rat IgG conjugated alkaline phosphatase (1:7,000; Sigma-Aldrich) was used as a secondary antibody. Detection was developed using nitroblue tetrazolium and 5-bromo-4-chloro-3-indolylphosphate (MBI Fermentas, Canada).

Bacterial clearance analysis

To evaluate the STG vaccine-induced protective immunity against homologous and heterologous challenges, the rats were sacrificed on 2 weeks post-challenge. The liver, lung, spleen, and kidney of each was harvested aseptically and homogenized in PBS using a tissue homogenizer (Sun MI Technology, Korea). Serial dilution of homogenates was then plated on selective media and incubated at 37˚C for 24–48 h. After incubation, CFUs were determined.

Statistical analysis

All statistical analyses were performed with one-way ANOVA using SPSS software (version 14.0) and data were expressed as mean ± standard error of the mean. Differences were considered statistically significant at \(P\) values of < 0.001, < 0.01 and < 0.05.

Results

Production and characterization of S. typhimurium ghosts

In order to produce STG, we first determined the MIC of NaOH (3.13 mg/mL) against S. typhimurium by using the two-fold broth dilution method. STG was successfully produced by using the specific concentration of NaOH. At the end of lysis process, no viable colonies were detected in the MIC of NaOH-induced STG. This indicates that the efficiency of chemically-
induced STG was 100%. Formation of the transmembrane tunnels in STG was examined by SEM (Fig 1A and 1B). Electron microscopic analysis revealed the presence of transmembrane lysis tunnel structures in STG (Fig 1B) as compared to the unlysed wild-type S. typhimurium cells (Fig 1A). Although the cellular morphology doesn’t seem to be affected by the MIC of NaOH, some of the cell surface structures seem to be slightly affected.

**Cytotoxicity tests and induction of cytokine mRNA expression in murine macrophages exposed to STG**

The cytotoxicity was compared using the viability of RAW 264.7 murine macrophages exposed to the FKC, wild-type bacterial cells and NaOH-induced STG, respectively (Fig 2). The given concentration of S. typhimurium LPS included in this analysis showed cell viability of 93.9%. The macrophages exposed to STG showed 84.9% cell viability (PBS: \( P < 0.01 \) and LPS: \( P < 0.05 \)), which was similar to those exposed to FKC (85.6%) but significantly higher than those exposed to wild-type bacterial cells (66.5%) under the same concentration.

To determine the activation of macrophages exposed to STG to produce inflammatory cytokines, mRNA expression of pro-inflammatory cytokines and factor (TNF-\( \alpha \), IL-1\( \beta \) and iNOS), anti-inflammatory cytokine (IL-10) and dual properties of cytokine (IL-6) was investigated using RT-qPCR (Fig 2). In TNF-\( \alpha \) mRNA, the highest mRNA level was found at 6 h in the macrophages exposed to wild-type bacterial cells (PBS and LPS: \( P < 0.001 \)), which was much higher than those exposed to FKC and STG, respectively. In STG-exposed macrophages, the maximum accumulation of TNF-\( \alpha \) mRNA was found at 2 h (PBS: \( P < 0.001 \) and LPS: \( P < 0.05 \)), which was a similar quantity found in macrophages exposed to wild-type bacterial cells, and then declined in a time-dependent manner. Similarly, the induction of IL-1\( \beta \) mRNA was induced in a time-dependent manner in the macrophages exposed to STG. Its maximum level was found at 6 h, which was less than those exposed to FKC and wild-type bacterial cells. In iNOS, IL-6 and IL-10 mRNA, the respective maximum levels were found at 6 h in the macrophages exposed to wild-type bacterial cells (PBS and LPS: \( P < 0.001 \)), which was slightly higher than those exposed to STG. Taken together, our data indicated that STG can activate macrophages to secrete both pro-inflammatory and anti-inflammatory cytokines.

**Humoral immune response**

To evaluate humoral immune responses of the STG vaccine against S. typhimurium and S. enteritidis antigens, the titers of total IgG in the sera of vaccinated and non-vaccinated animals

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**Fig 1.** Scanning electron microscopic analysis of (a) S. typhimurium and (b) STG. Arrows indicate trans-membrane lysis tunnels.

https://doi.org/10.1371/journal.pone.0185488.g001
To determine cytotoxicity, murine macrophages (RAW 264.7) were exposed to PBS buffer, LPS purified from *Salmonella typhimurium*, FKC, wild-type bacterial cells (ST) and NaOH-induced STG, respectively (bars). At 24 h post-exposure, macrophages were collected for analysis of cell viability using Cell counting Kit-8. Absorbance was measured at 450 nm and all experiments were performed in triplicate. Cytotoxic activity is expressed as the percentage of cell viability by the formula described in Materials and Methods. At given time post-exposure with PBS buffer, LPS purified from *Salmonella typhimurium*, FKC, ST and STG, respectively (bars), macrophages were collected for analysis of mRNA expression for pro- and anti-inflammatory cytokines and factor (TNF-α, IL-1β, iNOS, IL-6 and IL-10) using RT-qPCR. Data are representative of triplicate experiments with each sample run in triplicate. All statistical analyses were performed with one-way ANOVA using SPSS software (version 14.0) and data were expressed as mean ± standard error of the mean. *** = P < 0.001; ** = P < 0.01; * = P < 0.05 (significant difference from PBS) and ### = P < 0.001; ## = P < 0.01; # = P < 0.05 (significant difference from LPS).
were determined by indirect ELISA (Fig 3). Rats from FKC and STG groups elicited higher serum IgG antibody responses as compared to the PBS control group (Fig 3A). However, there were significant differences in the levels of serum IgG antibodies between FKC and STG at weeks 2 and 6. At week 9 post-challenge, rats vaccinated with either STG or FKC showed significantly higher IgG antibody responses than PBS control rats ($P < 0.01$). Most importantly, serum IgG antibodies in STG group rats increased robustly as compared to PBS group ($P < 0.01$) and FKC group ($P < 0.05$) throughout the vaccination and *S. enteritidis* challenge period (Fig 3B).

Fig 3. The levels of IgG antibody responses were determined by indirect ELISA. (a) *S. typhimurium* and (b) *S. enteritidis* were coated as an antigen for ELISA. Results are expressed as means ± standard errors of the means. The asterisks indicate significant differences between antibody responses of the vaccinated and non-vaccinated groups. ** = $P < 0.01$; * = $P < 0.05$.

https://doi.org/10.1371/journal.pone.0185488.g003
Cellular immune response

The CD4+ and CD8+ T-cell populations in vaccinated and non-vaccinated animals were observed by FACS (Fig 4A and 4B). As shown in Fig 4C, CD4+ T-cell populations significantly increased in all vaccinated animals (FKC: $P < 0.05$ and STG: $P < 0.01$) than the non-vaccinated control PBS group, but no significant difference detected between FKC and STG. Similarly, animals from FKC ($P < 0.05$) and STG ($P < 0.01$) had statistically higher percentage of CD8+ T-cells compared to the control PBS group (Fig 4D). Meanwhile, the statistical difference between FKC and STG was not significant.

Determination of serum bactericidal activity

Serum bactericidal activity from rats vaccinated with the STG vaccine was tested against *S. typhimurium* (Fig 5A) and *S. enteritidis* (Fig 5B). As can be seen in Fig 5, the group vaccinated with the STG vaccine showed significantly higher levels of bactericidal antibodies compared to FKC and PBS control groups ($P < 0.01$). However, at weeks 4 and 6, bactericidal activity was not significantly different between STG and FKC against homologous strain. Interestingly,
FKC vaccinated rats showed significant bactericidal antibodies against heterologous strain at week 6 ($P < 0.05$). Overall, the results suggest that STG vaccinated group increased bactericidal activities throughout the vaccination period (weeks 2, 4, and 6).

**Immunoblotting examination of the antibody response**

Specific anti-*S. typhimurium* and anti-*S. enteritidis* antibody responses in sera of vaccinated and non-vaccinated control animals were further analyzed by immunoblot using *S. typhimurium* or *S. enteritidis* envelope proteins as antigens. Serum from STG vaccinated animals were capable of recognizing envelope protein antigens from *S. typhimurium* and showed stronger...
protein bands including OMPs and envelope proteins than other groups (PBS and FKC). The molecular masses of envelope protein antigens were approximately 82, 62, 59, 57, 50, 40, 38, 37, 36, 27, 25, 21, 19, 17, 15, 12, 11 and 10 kDa (Fig 6A). Similarly, envelope proteins from \textit{S. enteritidis} were able to cross-recognize by STG vaccinated serum. The molecular masses of \textit{S. enteritidis} envelope protein antigens were found to be approximately 82, 62, 55, 47, 40, 38, 37, 36, 27, 21, 16, 15, 13 and 10 kDa (Fig 6B). Analysis of sera from FKC vaccinated animals showed weaker bands against \textit{S. enteritidis} protein antigens.

Homologous and heterologous protections of \textit{S. typhimurium} ghost vaccine

To examine the ability of the STG vaccine against \textit{S. typhimurium} challenge, rats were challenged orally with \textit{S. typhimurium} two weeks after the final booster vaccination. The bacterial loads in the liver, lung, spleen, and kidney were significantly lower in FKC ($P < 0.05$) and STG ($P < 0.01$) than in PBS (Fig 7A–7D). The protection against \textit{S. typhimurium} afforded by the
Fig 7. Protection of STG against homologous and heterologous challenges. Bacterial loads in liver, lung, spleen and kidney homogenates after (a-d) a homologous challenge with *S. typhimurium* or (e-h) a heterologous challenge with *S. enteritidis*. Results are expressed as means ± standard errors of the means. The asterisks indicate significant differences between the bacterial clearance of the vaccinated and non-vaccinated groups.

** = $P < 0.01$; * = $P < 0.05$.

https://doi.org/10.1371/journal.pone.0185488.g007
STG vaccine was significantly greater than that afforded by the FKC vaccine ($P < 0.05$). In order to determine the cross-protective efficacy of the STG vaccine, rats were vaccinated with STG, FKC or PBS and challenged with a virulent *S. enteritidis* strain. The results of the bacterial loads in liver, lung, spleen, and kidney are shown in Fig 7E–7H. The bacterial count in liver, lung, spleen, and kidney were significantly lower in STG than those in PBS and FKC ($P < 0.05$). These results suggest that the vaccination with STG vaccine provided protection against *S. enteritidis* strain, but FKC vaccinated rats failed to protect against *S. enteritidis* strain.

**Discussion**

The development of safe and effective vaccines against different *Salmonella* serotypes is an urgent need for the control and prevention of *Salmonella* infections. It is known that macrophages and dendritic cells are the key cells to be infected by *S. typhimurium*, and these antigen-presenting cells transport the bacteria to the liver and spleen, the major organs of bacterial replication [33,34]. Both cells are critical components to the immune response to *S. typhimurium* infection, with macrophages controlling bacterial growth in the early phase of infection and dendritic cells initiating the T cell response [35–37].

In this study, the higher cell viability in the macrophages exposed STG than that of wild-type bacterial cells, suggesting that the MIC of NaOH reduces the cytotoxic effect of wild-type bacterial cells. Previously, alkaline treatment to a Gram-negative bacterial LPS resulted in reduced toxicity and decylated LPS. The resultant LPS showed intact amide-linked fatty acids and removal of ester-linked fatty acids [38], but it was antigenically poor [38,39] or deficient [40]. However, our previous studies demonstrated that immunization with the NaOH-induced BG from a Gram-negative bacterium or a Gram-positive bacterium induces effective immune responses and provides a good protection against virulent challenge [17,18]. The present study also supports previous studies and suggests that the NaOH treatment does not affect the immunogenicity of BG or its potential as a vaccine candidate. Furthermore, our previous results showed that LPS of BG is modified or party lost by NaOH treatment and suggested that immunogenicity of the NaOH-induced BG could be derived from other cell envelope components rather than LPS [19].

Murine macrophages respond with secreting a variety of cytokines (TNF-α, IL-1, IL-6, IL-10 and IL-12) [37,41], chemokines, cell surface receptors, signaling molecules and transcriptional activators [42]. Simultaneously, co-incubation *S. typhimurium* with murine macrophages enhanced production of oxygen-dependent antimicrobial molecules such as superoxide, hydrogen peroxide and nitric oxide (NO) to kill intracellular *Salmonella* bacteria [43–45]. Like wild-type *S. typhimurium*, the NaOH-induced STG effectively activated the mRNA expression of TNF-α, IL-1β, iNOS, IL-6 and IL-10 in this study.

Furthermore, we demonstrate that the NaOH-induced STG vaccine has the ability to induce both humoral and cellular immune responses and protects rats from *S. typhimurium* and *S. enteritidis* challenges. The cellular morphology and cell surface structures were unaffected by the MIC of NaOH during lysis process. This result indicates that the MIC of NaOH successfully induced non-living STG vaccine. Previously, we produced the chemically induced BG from a Gram-positive bacterium *Staphylococcus aureus* and a Gram-negative bacteria *S. enteritidis* and *Vibrio parahaemolyticus* with 100% lysis efficiency [17–20].

In our previous study with chemically-induced *S. enteritidis* ghosts (SEG), the SEG vaccine has shown the usefulness of detecting serum IgG antibody response after challenge and provided the protective immunity that might be mediated by induction of humoral immune responses [17]. Peng et al [46] also demonstrated that humoral immune responses play an important role in preventing *Salmonella* infections. In this regard, serum IgG antibodies in
STG vaccinated rats after challenge with *S. typhimurium* and *S. enteritidis* strains were evaluated. Compared with the levels of antibody response in PBS and FKC control groups, STG vaccinated animals had higher serum IgG antibody response throughout the vaccination as well as the challenge period. Similar results were shown in *S. enteritidis* challenge studies, suggesting STG vaccine has the ability to induce protective immune responses against different *Salmonella* serotypes. In contrast, the FKC vaccine induced significant protective immunity against the *S. typhimurium* strain, but failed to induce significant antibody responses and protective immunity against the *S. enteritidis* strain.

Western blot analysis supported the difference between immune response induced by the STG vaccine and FKC vaccine. Compared with the FKC vaccine, the STG vaccinated sera showed stronger protein bands against the *S. typhimurium* and *S. enteritidis* antigens. This could be due to the loss of major surface antigenic components in the FKC vaccine. Hofstra et al [47] described that antigenic cross-reactivity of the major outer membrane proteins (30–42 kDa) was a general phenomenon in Enterobacteriaceae. The lack of heterologous antibodies revealed in the western blot analysis could be a reason for the lesser protection against the *S. enteritidis* challenge. In both the *S. typhimurium* and *S. enteritidis* strains, the serum bactericidal antibodies were significantly induced in STG vaccinated animals as compared to FKC and PBS vaccinated animals. Particularly, serum bactericidal analysis revealed that rats vaccinated with FKC vaccine produced higher levels of antibodies to the *S. typhimurium* strain. These antibodies could not induce cross-protective antibodies to the *S. enteritidis* strain. The data from the present study indicate that vaccination with the STG vaccine induces a strong protective immunity against *S. typhimurium* and *S. enteritidis* challenges.

It has been reported that BG from pathogenic bacteria has the ability to induce strong humoral and cellular immune responses and protect experimental animals from virulent challenge [48,49]. CD4⁺ MHC II- and CD8⁺ MHC I-restricted T cells play an important role in destroying intracellular bacteria, such as *Salmonella* [50–52]. Our study showed that STG vaccinated rats significantly induced CD4⁺ and CD8⁺ T cell populations compared to FKC and PBS control groups. Jawale et al [25] demonstrated that BG vaccine-induced CD4⁺ and CD8⁺ T-cells were involved in protection against *Salmonella* infections. However, recent studies have suggested that B-cell and T-cell immune responses are equally important for protective immunity against *S. typhimurium* [53,54].

Recently, genetically inactivated BG vaccines decreased bacterial load in tissue homogenates and provided protection against *Salmonella* infections [25,46]. Similarly, immunization with the chemically-induced SEG vaccine reduced bacterial clearance in liver, lung, spleen, and kidney and protected rats from *S. enteritidis* challenge [17]. In a previous study with *Salmonella* live vaccines, aromatic amino acid-dependent, streptomycin dependent and galE mutants [55–58] effectively induced protective immunity against a homologous challenge. However, these vaccines have limited potential to induce protective immunity against heterologous challenge. Effective *Salmonella* vaccination in livestock needs the induction of homologous and heterologous protective immunity. Based on these results, vaccination with the STG vaccine strongly induced protective immune responses and protected experimental animals from *S. typhimurium* and *S. enteritidis* infections.

**Conclusion**

In conclusion, vaccination with the STG vaccine induced higher levels of humoral and cell-mediated immunity than the FKC vaccine and exhibited a significant reduction in bacterial colonization against *S. typhimurium* and *S. enteritidis* strains. The potential of chemically-induced BG to heterologous protection is now being evaluated. Moreover, the results obtained
from this study suggest that the envelope antigen expression could be an important component of inducing cross-protective immune responses to the \textit{S. enteritidis} strain. Therefore, the STG vaccine could be a promising candidate for the development of inactivated vaccines against \textit{Salmonella} infections.

**Supporting information**

\textbf{S1 Table.} Primers sequences of the targeted genes in real-time PCR. (DOCX)

**Acknowledgments**

We thank Dr. Han-Oh Park (Professor in Department of Biology & Medicinal Science of Pai Chai University and CEO of Bioneer Corporation) to encourage and support this study to produce economical BG vaccine as global public good for the third world.

**Author Contributions**

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