Detection of virulence genes and investigation of the immunogenicity of outer membrane proteins of *Salmonella enterica*

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**ABSTRACT**

*Salmonella enterica* is one of the most important food-borne pathogens, causing a variety of diseases in humans and animals. This study aimed to detect the virulence genes in 33 *S. enterica* strains isolated from patients and to investigate the immunogenicity of the outer membrane proteins (OMPs) of *S. enterica* serovar Typhimurium. The aggregative fimbiae (*agfA*) gene was detected in all *S. enterica* isolates except one strain, *Salmonella* Paratyphi C strain SA7. In addition, 81.8% of the isolates harbored the *sefC* gene (fimbrial protein). However, all of the tested *S. enterica* isolates possessed the *fimA, hilA, invA, stn*, and *misL* virulence genes, regardless of serovar. The predominant OMPs of *S. enterica* Typhimurium SA3 identified by 12% sodium dodecyl sulfate-Polyacrylamide Gel Electrophoresis (PAGE) were used as eliciting antigens in the experimental mice. The results of the protection studies indicated that the selected OMPs conferred varying degrees of protection. However, the highest protection was observed using the 38-kDa OMP, which provided 100% protection to mice challenged with 50× LD₉₀ of *Salmonella* Typhimurium SA3 and 75% protection to mice subjected to an even higher bacterial challenge of 100× LD₉₀. The humoral response in mice caused by the 38-kDa OMP was confirmed using an immunodiffusion assay. This 38-kDa OMP is a promising candidate for the vaccine development against *S. enterica* Typhimurium. Further research on the protein structure was recommended.

**1. INTRODUCTION**

Food-borne infections caused by *Salmonella enterica* subsp. enterica (salmonellosis) represent a major public health challenge worldwide. Salmonellosis causes substantial morbidity and mortality throughout the world. Approximately 200,000 people die of typhoid globally each year, and most of these mortalities occur in the developing countries [1,2]. In addition, there has been a dramatic increase in the number of reported cases of salmonellosis in the developed countries, due in part to travelers visiting endemic regions [3]. More than 2,500 *S. enterica* strains have been reported worldwide based on their antigenic properties, and most of these strains cause either typhoidal or non-typhoidal salmonellosis [4,5]. Typhoidal salmonellosis is caused by *S. enterica* serotypes Typhi and Paratyphi A, B, and C, causing enteric fever, gastroenteritis, and bacteraemia, whereas non-typhoidal salmonellosis is caused by *S. enterica* serotypes Enteritidis and Typhimurium, causing a variety of diseases ranging from mild/severe gastroenteritis to systemic infection in humans and animals [6-8].

The wide range of *Salmonella* infections is attributed to the presence of numerous virulence factors that play essential roles in *Salmonella* pathogenesis, including in host cell adhesion, invasion, and intracellular replication [9]. Most of the genes encoding virulence factors are located in the regions known as *Salmonella* pathogenicity islands (SPIs) in the *Salmonella* chromosome, two of which (SPI-1 and SPI-2) encode a type three secretion system (T3SS) that is involved in the invasion of host intestinal cells and intercellular survival. The two main regulatory proteins for T3SS function are H1A and InvA, which play important roles in the invasion of epithelial cells and macrophage apoptosis [10]. In addition, fimbiae, encoded by several putative fimbral operons, such as *agf, fim,* and *sef,* play a significant role in *Salmonella* pathogenicity by binding to specific receptors on the host cells, leading to bacterial colonization, and/or toxin delivery [11,12]. In addition, *Salmonella* is one of the causative agents of diarrhea due

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to its ability to produce enterotoxin via a process mediated by the stn gene [2,12]. Additionally, the misL gene that encodes MisL, an auto-transporter protein, is an extracellular matrix adhesin that is involved in the invasion process and intestinal colonization, leading to systemic infection [2,12].

In the recent years, there has been a continuous emergence of multidrug-resistant Salmonella strains; therefore, the development of an effective vaccine for systemic Salmonella infections has become of a great importance for the control of these global food-borne pathogens [13,14]. Vaccination is one of the most effective approaches for infectious disease control and can successfully protect humans against various diseases. Currently, there are two approved vaccines against typhoid Salmonella, namely, the Vi polysaccharide vaccine and the live attenuated Salmonella Typhi Ty21a strain vaccine. However, there is no licensed vaccine against non-typhoid Salmonella [15,16]. Among the components of Salmonella that may be possible candidates for vaccine development are the cell outer membrane proteins (OMPs), which play an essential role in adaptation to environmental conditions, motility, adherence, and host cell colonization. Furthermore, OMPs play a significant role in the injection of toxins and cellular proteases, as well as the formation of channels for the removal of antibiotics (efflux pumps) [5,14,17]. The aim of the current work included determination of various virulence genes in S. enterica strains (n = 33) isolated from patients in Saudi Arabia [18] and investigation of the immunogenicity of the OMPs of S. enterica serovar Typhimurium as a vaccine candidate against typhoid in experimental animals.

2. MATERIALS AND METHODS

2.1. Microorganisms and Growth Culture

The various S. enterica strains studied (n = 33) were previously isolated from clinical and environmental sources (Riyadh, Saudi Arabia) and assigned the accession numbers KU843835–KU843866 [18]. Bacterial growth was carried out on the media recommended for Salmonella spp. including xylose lysine deoxycholate agar (Oxoid, UK) and deoxycholate citrate agar (Oxoid) for 24 hours at 37°C.

2.2. PCR-Based Detection of Virulence Genes

The isolates of S. enterica (n = 33) were screened for seven virulence genes by Polymerase Chain Reaction (PCR) according to the methods reported previously with some modification using the set of specific primers shown in Table 1 [2]. The tested virulence genes included aggregative fimbiae (agfA), fimbrial proteins (sefC, fimA), Salmonella pathogenicity island-3 (misL), Salmonella pathogenicity island-1 (hilA), and Salmonella specific invasive (invA) and enterotoxic (stn) genes. The genomic DNA of the S. enterica isolates (n = 33) was extracted from overnight cultures using the DNeasy Blood and Tissue Kit (Qiagen, UK) following the manufacturer’s instructions. The PCRs were performed in a final volume of 25 µl, containing 12.5 µl of GoTaq® Green Master Mix (Promega), 2 µl of DNA (50 µg/ml), 2 µl of forward and reverse primers, and 6.5 µl of nuclease-free water (Promega, UK). The PCR was carried out under the following conditions: initial denaturation for 5 minutes at 95°C; 35 cycles of denaturation at 95°C for 30 seconds, annealing at a specific annealing temperature (Table 1) for 30 seconds, and extension at 72°C for 1.5 minutes; and a final extension step at 72°C for 5 minutes. The PCR products were analyzed by 1.5% (w/v) agarose gel electrophoresis in 1× TAE (40 mM Tris, 20 mM acetic acid, and 1 mM Ethylenediaminetetraacetic acid (EDTA), pH 8.0) at 90 V for 45 minutes using a 100-bp DNA ladder (Qiagen, UK) as the molecular size standard. DNA from Escherichia coli was used as a negative control.

2.3. Preparation of Outer Membrane Proteins

2.3.1. Extraction of outer membrane proteins

The OMPs of S. enterica serovar Typhimurium strain SA3 (accession no. KU843837) were extracted according to a previously reported

| Gene | Primer Sequence (5'-3') | Amplicon size (bp) | Annealing temp. (°C) |
|------|-------------------------|--------------------|---------------------|
| agfA | F-TCCGCGCAGGACTCAACG     | 261                | 63                  |
|      | R-CAGCGCCGGGGTTATACCG    |                    |                     |
| sefC | F-TGGGAGAAATATCCAGTCG    | 1100               | 58                  |
|      | R-CTATTGGCCTCTGCTGCA     |                    |                     |
| fimA | F-CCTTTCTTCCATCCTGCTGAA  | 85                 | 58                  |
|      | R-TGGGTATATCTGCTGCA      |                    |                     |
| misL | F-GACGTTGATATGTCCTGATCCAG| 986                | 58                  |
|      | R-CCAATGCCGGCCAGTTCCGTCGC|                    |                     |
| hilA | F-GGAGATGTGATGAAAAACACC  | 413                | 63.5                |
|      | R-CTGCCGGAGGATATAATACCG  |                    |                     |
| invA | F-CCTGCTTCTTCATCCTGCAAGTGTCGTCG | 495 | 57 |
|      | R-GCATCAATAATACCGAGCTTC |                    |                     |
| Stn  | F-CTTGTGTCGAAAATAGGCAG   | 260                | 55                  |
|      | R-TGCCCCAGCACAGGATTC     |                    |                     |
method with some modifications [19]. Briefly, 1 g (wt weight) of *Salmonella* Typhimurium cell biomass was re-suspended in 20 ml of extraction buffer [10 mM Tris-HCl (pH 7.5), 10 mM EDTA disodium salt and 6 M urea], and stirred for 1 hour at 4°C. The cell extract was dialyzed against distilled water for 48 hours at 4°C with frequent changing of the water (every 6 hours). The dialyzed extract was centrifuged at 6,000 rpm for 1 hour at 4°C, and the supernatant was collected and lyophilized (LABCONCO, Czech Republic). The crude OMPs were stored at −75°C until use.

2.4. Estimation of Protein Concentration
The protein concentration was determined according to the method described by Bradford [20]. One milliliter of Bradford reagent was added to 50 µl of sample, and the optical density was measured after 5 minutes at 595 nm. Bovine serum albumin (50–1,000 µg/ml) was used as the protein standard.

2.5. Polyacrylamide Gel Electrophoresis
The extracted OMPs were separated by sodium dodecyl sulfate (SDS)-Polyacrylamide Gel Electrophoresis (PAGE) according to the Laemmli [21] method using a mini gel electrophoresis system (Biometra, UK). The protein samples were solubilized in sample loading buffer [0.625 M Tris-HCl (pH 6.8), 2% SDS, 10% glycerol, and 5% β-mercaptoethanol] at 100°C for 5 minutes. SDS-PAGE was carried out on a 12% separating gel with a discontinuous buffer system. Thereafter, the gels were stained by Coomassie blue staining solution for 1–2 hours with continuous shaking. The gel was transferred to destaining solution and subjected to shaking with continuous changing of the solution until distinct protein bands were visible.

2.6. Purification of OMPs
Protein-acrylamide (PA) complexes were prepared according to the method of Hamid and Jain [19]. The major OMPs extracted from *S. enterica* Typhimurium strain SA3 were separated by 12% SDS-PAGE, and six dominant protein bands with molecular sizes of 27, 29, 38, 48, 64, and 70 kDa were selected and used as eliciting antigens. The bands corresponding to the selected proteins were excised from the gels using a sterile blade, transferred to a 50-ml centrifuge tube, washed twice with distilled water, and pulverized using a small mortar and pestle. The pulverized materials containing proteins and acrylamide were lyophilized to form a fine powder and kept at −75°C until further use.

2.7. Mouse Husbandry and Immunization
BALB/c female mice, 6–8 weeks old (20–25 g), were obtained from the Central Animal House at the College of Pharmacy (King Saud University). The mice were housed in clean, well-aerated polypropylene cages at 25°C and fed a pellet diet and water. The mouse immunization schedule was performed according to Hamid and Jain [19] and Yang et al. [16]. The experiment was carried out using a total of 56 mice that were divided into 14 groups (n = 4 mice/group). Twelve groups were immunized subcutaneously with the six selected proteins (two mouse groups for each OMP), and two groups of animals served as controls (Supplementary Data, S1). Prior to immunization, the mice were bled from the tail veins to obtain pre-immune sera. The prepared PA complex powder was suspended in phosphate-buffered saline (PBS) and emulsified with an equal volume of complete Freund’s adjuvant to obtain a final protein concentration of 25 µg protein/0.5 ml of emulsion. Two mice from each group received four subcutaneous injections of the PA complex (25 µg of protein/dose) on days 0, 7, 21, and 28 and a final booster dose on day 40. Two mice in each group acted as controls and received lyophilized polyacrylamide (without protein) dissolved in PBS under the same immunization schedule. In addition, two mouse groups (n = 8) did not receive any injection and acted as healthy control B. Thereafter, the mice were bled on day 50 to obtain immune sera and kept at −75°C until use.

2.8. Determination of Bacterial Lethal Dose (LD₅₀)
The 50% lethal dose (LD₅₀) of *S. enterica* serovar Typhimurium strain SA3 was determined according to the previously reported methods [22,23]. Serial dilutions of overnight cultures of strain SA3 were prepared, and cell suspensions with cell counts ranging from 2 × 10⁴ to 2 × 10⁶ CFU/ml were orally administered to six groups of mice (8 mice/group). The animals were monitored for 14 days, and the lethal effect was recorded. The LD₅₀, which was defined as the bacterial dose that led to the death of 50% of the mice, was estimated based on the number of surviving mice on day 14.

2.9. Animal Challenge with *Salmonella* Typhimurium SA-3
After 4 weeks of immunization with the PA complexes, the immunized and control mice were injected intraperitoneally with two different doses of *S. enterica* Typhimurium SA3, including 50× and 100× LD₅₀. In addition, five mice were injected with PBS as a control. After the bacterial challenge, the animals were monitored for 14 days, and their clinical symptoms and mortality rate were recorded. Bacterial infection was confirmed by re-isolation of *S. enterica* Typhimurium SA3 from the infected mice. Briefly, livers and spleens were collected from the infected and control mice under aseptic conditions on day 2 post-infection. Then, the livers and spleens were homogenized in sterile phosphate buffer, and aliquots were inoculated into selenite F broth and incubated for 18 hours at 37°C. Thereafter, a loopful of each culture was transferred to nutrient agar and incubated under the same conditions. The obtained bacterial colonies were identified by a Vitek® 2-C15 automated system (BioMerieux Inc., France) and 16S rDNA sequencing analysis as previously described [18].

2.10. Histopathological Examination
Histopathological changes in the mouse organs caused by *Salmonella* Typhimurium SA3 were investigated. The spleens and livers were removed from the infected and control mice, fixed in formalin solution (10%), and kept in paraffin blocks. Transverse sections (5 µm) were prepared using a microtome, stained with eosin and haematoxylin (H/E), and examined by microscopy [16].
2.11. Detection of OMP-Specific Antibodies

The humoral immune response elicited by OMPs in the immunized animals was detected using immunodiffusion [24]. Briefly, 1% (w/v) melted agarose solution, prepared in sterile PBS, was poured into sterile Petri dishes (=2-mm thick). After gel solidification, wells were punched in the gels using a sterile cork borer. Then, 50 µl of the crude extracts containing Salmonella Typhimurium SA3 OMPs were loaded in the central wells, while the surrounding wells were loaded with 50 µl of different sera obtained from mice immunized with different OMPs (27, 29, 38, 48, 64, and 70 kDa OMP). Thereafter, the plates were incubated at 4°C for 24–48 hours and monitored for agglutination.

3. RESULTS AND DISCUSSION

The emergence of multidrug resistance in S. enterica, the causative agent of salmonellosis, has led to further investigation of the virulence factors of this organism and to the search for an effective control method [25,26]. Consequently, this study aimed to identify the various virulence factors of S. enterica strains isolated in Saudi Arabia, in addition to preliminary identification of vaccine candidates against S. enterica serovar Paratyphi Typhimurium based on the OMP subunits.

3.1. Detection of Virulence Genes of S. enterica Isolates

The degree of pathogenicity of S. enterica depends on several virulence factors that are encoded by various genes, including the fimbriae operon, virulence plasmid, SPIs, and enterotoxin gene. However, the association of some of these factors with pathogenicity remains controversial [27]. In the present study, the presence of putative virulence genes (n = 7) responsible for the pathogenicity of S. enterica (n = 33) isolated from patients in Saudi Arabia was detected by PCR using a set of primers specific to various virulence genes. The aggregative fimbriae (agfA) gene was detected, with the expected PCR product size of approximately 261 bp (Fig. 1), in most S. enterica strains (n = 32/33), being absent in only S. enterica serovar Paratyphi C strain SA7. As shown in Figure 2, the sefC (fimbrial protein) gene was amplified with the correct amplicon size of 1,100 bp in 81.8% of the isolates (27/33). Fimbriae play a role in Salmonella pathogenicity by binding to specific receptors of host cells, leading to bacterial colonization, and/or toxin delivery, which is reflected in the appearance of infection [28,29]. The high frequency of agfA was similar to the results of previous studies on different Salmonella serovars [25,28]. In addition, fimbrial protein SEF14, which encodes the sef operon, was detected in only Salmonella Enteritidis and Salmonella Paratyphi B isolates. This finding is consistent with the results of previous studies, providing a useful marker to discriminate S. Enteritidis from Salmonella Typhi as well as Salmonella Paratyphi B from Salmonella Typhimurium isolates [30]. Salmonella strains express type-1 fimbriae that enable the bacteria to bind to eukaryotic cells. Fimbrial proteins are encoded by the fim gene. In this study, all the isolates (n = 33) were found to harbor the fimA gene (S1), which consistent with previous reports that studied the prevalence of the fimA operon in various Enterobacteriaceae strains [28,30]. On the other hand, all the tested S. enterica isolates (n = 33) were found to harbor hilA (Salmonella pathogenicity island-1), invA (Salmonella specific invasive), stn (enterotoxic), and misL (Salmonella pathogenicity island-3), regardless of serovar and the source of the isolate, yielding PCR amplicons of the expected sizes of 413, 495, 260, and 986 bp, respectively (Supplementary Data S2–S5). All these genes exhibited a high degree of conservation among Salmonella strains, but are absent among other closely related enteric bacteria [2,6,11] (Fabián et al., 2012).

3.2. Immunogenicity of OMPs

The OMPs of Gram-negative bacteria can elicit a significant antibody response from the host defense system. However,
there is relatively limited information regarding the potential of OMPs to confer protection against Salmonella infections [14,31]. Moreover, while there are two approved vaccines against typhoid Salmonella, including the Vi polysaccharide vaccine and a live attenuated Salmonella Typhi Ty21a strain vaccine, there is no licensed vaccine against non-typhoid Salmonella [15,30].

3.3. Preparation of OMPs of Salmonella Typhimurium SA3

The OMPs of Salmonella Typhimurium SA3 were extracted using the urea-EDTA extraction method. The urea extraction method has been proven to be effective for solubilization of the surface proteins of a number of bacterial species and has yielded favorable results [19]. The SDS-PAGE analysis pattern showed approximately 20 proteins with molecular masses ranging from approximately 15–95 kDa (Fig. 3). Six of the most dominant OMPs of S. enterica Typhimurium SA3 were selected for the immunization studies, including proteins with molecular masses of 27, 29, 38, 48, 64, and 70 kDa.

3.4. Mouse Protection Studies

The roles of the selected OMPs in providing protection against Salmonella infection in experimental mice were investigated. The selected OMP bands were excised from the PAGE gel, processed to obtain pulverized PA complexes, and then used to immunize mice. Thereafter, the mice immunized with various OMPs were challenged with two different doses of S. enterica Typhimurium SA3, including 50× and 100× LD₅₀, and monitored for the appearance of infection symptoms and/or mortality in comparison with the control mice. For confirmation of infection of the mice with Salmonella Typhimurium SA3, the bacteria were re-isolated from livers and spleens collected from infected and control mice. The obtained bacteria were identified by biochemical tests and 16S rDNA sequencing.

The results revealed the isolation of Salmonella Typhimurium SA3 from the infected and control mice, indicating the success of the challenge experiment. The animals were monitored for up to 14 days post-infection, and their clinical symptoms and mortality rate were recorded. The results indicated that after 1 day of infection, the control mice (n = 4) showed symptoms of salmonellosis, including skin pustules, sluggishness, ruffled hair, slow or no responsiveness, and loss of appetite, and these mice died within 3–5 days (Supplementary Data S7). However, immunization of the mice with various OMPs led to the protection of the mice to varying degrees (Table 2). Immunization of the mice with the 38-kDa OMP resulted in survival of 100% and 75% of the mice challenged with doses of 50× LD₅₀ and 100× LD₅₀ of Salmonella Typhimurium strain SA3, respectively. In addition, immunization with both the 64- and 70-kDa OMPs resulted in the protection of 66.7% of the mice challenged with a dose of 50× LD₅₀. The 48-kDa protein provided 75% and 50% protection under similar conditions, whereas the 27- and 29-kDa proteins exhibited 75% and 66.7% protection, of mice injected with doses of 50× and 100× LD₅₀, respectively. The lowest protection (33.3% of the mice) was observed in animals immunized with the 64-kDa OMP.

This strong protection of the immunized mice by the 38-kDa OMP, in comparison to the control, against Salmonella Typhimurium infection can be attributed to elicitation of specific and/or nonspecific immune responses [14]. In comparison to the results of other studies, the 38-kDa OMP provided effective and superior protection against Salmonella Typhimurium infection. A 43-kDa truncated OmpC pore protein showed 50%–75% protection against Salmonella Typhimurium infection in birds [32]. Recently, Toobak et al. [33] identified three porin OMPs from S. enterica serovar Typhi with similar molecular sizes of approximately 37.62 kDa (OmpA), 40.7 kDa (OmpC), and 39.05 kDa (OmpF); however, immunization of experimental mice with these proteins did not lead to protection against Salmonella infection [33]. However, a non-porin 19-kDa OMP investigated by Hamid and Jain [19] showed the increased protection (100%) of mice challenged with 100× LD₅₀ of Salmonella Typhimurium. Immunization with the OMPs of Salmonella is considered to be a potential approach for conferring protection against typhoid. The OMPs of Gram-negative bacteria include a family of por-forming proteins that form water-filled channels that allow small hydrophilic solutes to pass through the pore, in addition to other

Table 2: Results of mice immunization with various outer membrane proteins of S. enterica serovar Typhimurium SA3.

| Proteins MW (kDa) | Number of mice that survival at 14 days/Total number |
|------------------|-----------------------------------------------------|
|                  | 50 × LD₅₀                                           | 100 × LD₅₀                                    |
| 27               | 3/4 (75%)                                            | 2/3 (66.7)                                   |
| 29               | 3/4 (75%)                                            | 2/3 (66.7)                                   |
| 38               | 4/4 (100%)                                           | 3/4 (75%)                                   |
| 48               | 3/4 (75%)                                            | 2/4 (50%)                                   |
| 64               | 2/3 (66.7%)                                          | 1/3 (33.3%)                                  |
| 70               | 2/3 (66.7%)                                          | 0/3 (0%)                                    |
| Control          | 0/4 (0%)                                             | 0/4 (0%)                                     |
non-porin proteins that play significant roles in the pathogenicity of the organism [34,35].

3.5. Detection of OMP-Specific Antibodies

The sera collected prior to and after the mice immunizations were evaluated for the humoral immune response elicited by OMPs using immunodiffusion (Fig. 4). The results indicated that the OMP with a molecular weight of 38 kDa stimulated a significant humoral response in mice. Sera from animals immunized with the 38-kDa OMP could be precipitated with the OMPs of S. enterica Typhimurium SA3. In addition, the antiserum of the OMP with a molecular weight of 29 kDa exhibited a faint precipitation line, whereas no reaction was observed with antisera obtained from other OMP groups. The serum from unimmunized animals did not show any precipitation with the extracted Salmonella Typhimurium OMPs.

3.6. Histopathological Studies

The protection efficiency of the 38-kDa OMP against Salmonella Typhimurium challenge in immunized mice was further evaluated based on the morphological analysis of the histopathological changes. The livers and spleens that were collected from the unimmunized infected control showed clear hepatomegaly and splenomegaly compared with those of uninfected mice (Fig. 5). The pathological study indicated that the H/E-stained liver and spleen sections of the control mice showed clinical signs of Salmonella infection, including focal necrosis of the liver, distinct splenorrhagia, and vacuolar degeneration of splenic reticular cells. In addition, the H/E-stained sections showed the presence of inflammatory cells in both the liver and spleen as well as abnormal structures such as vacuoles and hydropic degeneration (Figs. 6 and 7). On the other hand, the immunized mice showed standard healthy structures [16].

Figure 4: Gel immune-diffusion precipitation test showed precipitate line with OMP-38 antiserum against OMPs mixture.

Figure 5: The spleen and Splenomegaly (A); and Liver and Hepatomegaly (B) of infected and uninfected mice.

Figure 6: Spleen section of healthy (control) (A) and infected (B) spleen section with inflammatory cells.
4. CONCLUSION

PCR-based detection of the virulence genes in *S. enterica* strains (*n* = 33) isolated from patients in Saudi Arabia revealed slight variations in gene distribution among the isolates, particularly for the aggregative fimbriae gene (*agfA*) and the fimbrial protein-encoding gene *sefC*. Investigation of the immunogenicity of OMPs of *Salmonella* Typhimurium SA3 revealed that several OMPs conferred varying degrees of protection against infection. Mice immunized with the 38-kDa OMP exhibited 100% protection when challenged with 50× *LD*₅₀ of *Salmonella* Typhimurium SA3 and 75% protection against an even higher bacterial challenge of 100× *LD*₅₀. This 38-kDa OMP is a promising potential vaccine against *S. enterica* serovar Typhimurium. Further research on the protein structure is in progress.

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SUPPLEMENTARY MATERIAL

**Supplementary Figure 1:** Agraose gel electrophoresis of the amplified *fimA* gene. 15 µl of the PCR product was separated in 1% agarose containing ethidium bromide solution (1 µg/ml), and visualized using Gel documentation system. M: 100bp ladder; Neg: negative control; Other lanes: *S. enterica* strains (n=33).

**Supplementary Figure 2:** Agraose gel electrophoresis of the amplified *misL* gene. 15 µl of the PCR product was separated in 1.5% agarose containing ethidium bromide solution (1 µg/ml), and visualized using Gel documentation system. M: 100bp ladder; Neg: negative control; Other lanes: *S. enterica* strains (n=33).
Supplementary Figure 3: Agraose gel electrophoresis of the amplified hilA gene. 15 µl of the PCR product was separated in 2% agarose containing ethidium bromide solution (1 µg/ml), and visualized using Gel documentation system. M: 100bp ladder; Neg: negative control; Other lanes: S. enterica strains (n=33).

Supplementary Figure 4: Agraose gel electrophoresis of the amplified invA gene. 15 µl of the PCR product was separated in 2% agarose containing ethidium bromide solution (1 µg/ml), and visualized using Gel documentation system. M: 100bp ladder; Neg: negative control; Other lanes: S. enterica strains (n=33).
Supplementary Figure 5: Agarose gel electrophoresis of the amplified stn gene. 15 µl of the PCR product was separated in 2% agarose containing ethidium bromide solution (1 µg/ml), and visualized using Gel documentation system. M: 100bp ladder; Neg: negative control; Other lanes: S. enterica strains (n=33).

Supplementary Figure 6: Immunization of experimental mice with OMPs extracted from S. enterica Typhimurium SA-3. The experiment was carried out using a total of 56 mice that were divided into 14 groups (n=4 mice/group). Twelve groups were immunized subcutaneously with the selected six proteins, (two mice groups for each OMP), and two groups of animals were served as control. The prepared proteins-acrylamide (PA) complex powder was suspended in phosphate-buffered saline (PBS), emulsified with an equal volume of complete Freund’s adjuvant to give final protein concentration of 25 µg protein/0.5 ml emulsion. Two mice from each group received four subcutaneous injections of the PA complex (25 µg protein/dose) on days 0, 7, 21, 28 and a final booster dose on day 40. The control mice (n=2) in each group received lyophilized polyacrylamide (without protein) dissolved into PBS at the same immunization schedule. Control B mice didn't receive any injection.
Supplementary Figure 7: Shows the immunization experiments of *Salmonella* vaccine attempt (control) (A); infected mice with salmonellosis symptoms such as ruffled hair and Skin pustules (B, C & D); Dead mice (E & F).