Identification of immunogenic outer membrane proteins and evaluation of their protective efficacy against *Stenotrophomonas maltophilia*

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**Abstract**

**Background:** *Stenotrophomonas maltophilia* (*S. maltophilia*) is an emerging global multiple-drug-resistant organism. It becomes increasingly challenging to treat *S. maltophilia* infection effectively. Novel therapeutic and preventive approaches targeting *S. maltophilia* infection are still lacking. This study aims to isolate outer membrane proteins (Omps) from *S. maltophilia* and use immunoproteomic technology to identify potential vaccine candidates of Omps against *S. maltophilia* infections.

**Methods:** Omps from *S. maltophilia* culture were separated by two-dimensional electrophoresis and identified by matrix-assisted laser desorption/ionization time of flight mass spectrometry and nano liquid chromatography coupled fourier transform ion cyclotron resonance tandem mass spectrometry. Recombinant Omps were prepared and used to immunize mice, and the potency of mouse anti-Omp serum was tested in opsonophagocytic killing assay (OPKA). The effects of immunization with recombinant Omp on blood and tissue bacterial loads in a mouse model of *S. maltophilia*-induced infection were analyzed.

**Results:** Outer membrane protein A (OmpA) and Smlt4123 were identified by mass spectrometry. Mouse anti-Smlt4123 serum significantly reduced the bacterial counts in healthy individuals’ blood in OPKA (*P* < 0.05) but mouse anti-OmpA serum did not. Enzyme-linked immunosorbent assay revealed that the antibody subtype of mouse anti-Smlt4123 antibody was IgG1. Eight hours after an intraperitoneal challenge with *S. maltophilia*, the bacterial loads in mouse blood were significantly lower in the mice receiving immunization with recombinant Smlt4123 than in the control mice receiving no immunization (*P* < 0.05), whereas the bacterial loads in other organs, such as the liver, spleen, lung, and kidney were similar in the two groups.

**Conclusions:** The results revealed that the immunoproteomic approach was an efficient way to screen the immunogenic protein of *Stenotrophomonas maltophilia*. Moreover, the recombinant Smlt4123 had potential to protect mice from bacteremia caused by *S. maltophilia* in the early stages.

**Keywords:** Immunoproteomics, Smlt4123, *Stenotrophomonas maltophilia*, Vaccine candidate

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Background

Stenotrophomonas maltophilia (S. maltophilia), a newly emerging global opportunistic Gram-negative pathogen, exists widely in nature and hospitals [1]. It can be isolated from soil, plant roots, and aqueous-associated sources, such as water treatment and distribution systems, tap water, and bottled water [2]. S. maltophilia strains of both clinical and environmental origin have been reported to adhere to abiotic surfaces and form bacterial biofilm [3, 4]. Besides, S. maltophilia can adhere to mouse tracheal mucus and thus cause respiratory tract infections [5]. As the third most frequently isolated nonfermentative Gram-negative bacilli following Pseudomonas aeruginosa and Acinetobacter species [6], S. maltophilia is also a newly emerging multiple-drug-resistant organism [2]. S. maltophilia can cause infections in multiple organs and tissues, such as the respiratory tract, biliary and urinary tract, skin, bone and joint, heart, brain, eyes, and soft tissues [7–9]. Risk factors for S. maltophilia-associated infections include immunosuppression, cancer, indwelling devices, mechanical ventilation, and broad spectrum antimicrobial therapy [10, 11]. The mortality rate of S. maltophilia-associated infection ranges from 14 to 69% in patients with bacteremia [12]. S. maltophilia has been found to show resistance to a broad array of antibiotics, such as trimethoprim-sulfamethoxazole (TMP-SMX), β-lactam antibiotics, macrolides, fluoroquinolones, cephalosporins, aminoglycosides, chloramphenicol, carbapenems, tetracyclines, and polymyxins [13–15]. Thus, it is becoming increasingly challenging for physicians to use conventional therapies to treat S. maltophilia-associated infections effectively. Novel therapeutic or preventive approaches targeting S. maltophilia-associated infections are greatly needed. Currently, vaccines or antibody-based treatments have not been developed for S. maltophilia-associated infections.

Bacterial outer membrane proteins (Omps) play key roles in bacterial survival and multiplication in host systems. Omps of Aeromonas salmonicida have been found to regulate the adherence and serum and bile salt resistance of the bacteria [16]. Because many bacterial Omps can be easily recognized by the host immune system as foreign substances so to initiate host immune defense mechanism against the bacterial infection [17–20], they could be potential vaccine candidates against the infection. Immunoproteomic technology is now allowing screening immunogenic candidate proteins rapidly and effectively. In the current study, we used immunoproteomic technology to identify immunogenic Omps of S. maltophilia, developed antibodies against the Omps, and tested the effects of immunization with recombinant Omps on tissue bacterial loads in a mouse model of S. maltophilia infection.

Methods

Bacterial strain and human blood samples

The bacterial strain S. maltophilia K279a was used in this study. We chose this particular strain because the whole genome sequence of the bacterial strain is available and S. maltophilia K279a was considered as a representative genome sequence strain [21]. The bacteria were grown in lysogeny broth (LB) or on LB agar plates. Escherichia coli (E. coli) BL21 (DE3) were used for recombinant protein production. Blood samples from healthy individuals were obtained from the healthy volunteers, Affiliated Hospital of Military Medical Science in China.

Animals and ethics

New Zealand White rabbits weighing approximately 2 kg and Female BALB/c mice (5–6 weeks old) were housed at the animal care center of the Academy of Military Medical Sciences (AMMS), China. All animals were maintained under specific pathogen-free conditions and were kept in a climate-controlled room (temperature of 22 ± 1 °C and humidity 55 ± 5%) with a 12 h light/dark cycle. Animals were sacrificed by euthanasia, and all experiments were handled according to protocols approved by Institutional Animal Care and Use Committee of AMMS. All efforts were made to minimize animal suffering.

Rabbit anti-S. maltophilia serum preparation

S. maltophilia K279a were grown in LB at 37 °C shaker to reach an optical density at 600 nm (OD 600) of 1.0 (approximately 4.0 × 10^8 CFU/mL), and then inactivated by treatment with 0.15% (v/v) methanol at 37 °C for 24 h. The inactivated S. maltophilia was injected into two female New Zealand white rabbits (2 kg) (4.8 × 10^7 CFU/time) with a two-week interval between injections. The first injection was emulsified with Freund’s complete adjuvant (Sigma), and the remaining two injections were emulsified with Freund’s incomplete adjuvant (Sigma). Freund’s complete adjuvant is considered to be the gold standard adjuvant for immunization [22]. The Freund’s complete adjuvant has been commonly used to induce antibody production in animal models [23]. Blood samples from the rabbits were collected 7 days before the first injection (pre-immune serum) and 10 days after the last injection (post-immune serum). An indirect enzyme-linked immunosorbent assay (ELISA) was used to measure antibody levels in the serum samples. Briefly, ELISA plates were coated with 10 μg/mL S. maltophilia total protein in coating buffer [0.05 M NaHCO_3 (pH 9.6)] at 4 °C overnight and then blocked with 3% BSA in phosphate-buffered saline containing 0.1% Tween-20 (PBST) buffer for 1 h at 37 °C. The pre-immune serum (1:100 dilution in PBST) and post-immune serum (1:10240 dilution in PBST) were added to the wells. Horse radish peroxidase (HRP) conjugated anti-rabbit
IgG secondary antibody was added. The color was developed by using the 3,3',5,5'-Tetramethylbenzidine (TMB) single-component substrate solution (Beijing Solarbio Science & Technology Co., Ltd. China). The color development reaction was stopped by 100 μL 1 M H2SO4. The optical density at 450 nm (OD450) was determined in an ELISA plate reader.

Preparation of outer membrane proteins (Omps)

*S. maltophilia* Omps were prepared according to the previous description [24]. In brief, *S. maltophilia* were grown in LB at 37 °C shaker to reach OD600 = 1.0 and then collected by centrifugation. The bacterial pellets were washed and re-suspended in ice cold TS buffer (150 mM NaCl, 10 mM Tris-HCl, pH 7.4). The bacterial suspension was kept in ice, sonicated for 20 min, and centrifuged at 6000 rpm/min for 10 min at 4 °C. The supernatant was transferred to a 10 mL centrifuge tube and further centrifuged at 12800 rpm/min for 30 min at 4 °C. The supernatant was discarded and the cell membrane pellets were re-suspended in 10 mL TS buffer by repeated pipetting. An equal volume of 2% Sarkosyl was added to the cell membrane suspension to dissolve the cytoplasmic membranes. The mixture was incubated at room temperature for 30 min with intermittent mixing, and then centrifuged at 12800 rpm/min for 30 min at 4 °C to collect Omps. The supernatant was discarded. The Omp pellets were re-suspended in 5 mL PBS. Protein concentration of the Omp suspension was determined by the Plus One 2-D Quant Kit (GE Healthcare).

Two-dimensional polyacrylamide gel electrophoresis (2D PAGE) and western blot

Proteins in the Omp suspension were separated by 2D 10% sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE) isoelectric focusing (IEF) electrophoresis. Two hundred μg Omps in 350 μL rehydration buffer [7 M Urea, 2 M Thiourea, 4% (w/v) CHAPS, 50 mM DTT] were loaded in pH 4–7 IPG strips (18 cm, GE Healthcare, USA). The loaded strips were rehydrated overnight and underwent electrophoresis at room temperature at 300 V for 1 h, 600 V for 1 h, and 1000 V for 1 h, and then 8000 V for 8 h, to reach a total of 64 kVh. The proteins were separated and transferred to a polyvinylidene fluoride (PVDF) membrane (Bio-Rad, USA). The membrane was blocked with 5% skim milk in PBS overnight and then probed with the rabbit pre-immune or post-immune serum samples. The membrane was then washed and incubated with the secondary HRP-conjugated goat anti-rabbit IgG. The signals were developed by using the Super Signal West Dura Extended Duration Substrate (Pierce, USA), and the signals were detected using Image Scanner (GE Healthcare, USA).

Matrix-assisted laser desorption/ionization time of flight mass spectrometry (MALDI-TOF-MS) and nano liquid chromatography coupled fourier transform ion cyclotron resonance tandem mass spectrometry (nanoLC-FT ICR MS/MS)

MALDI-TOF MS was performed on a Bruker Reflex III MALDI-TOF-MS (Bruker Daltonics, Germany) operating in the reflectron mode with 20 kV accelerating voltage and 23 kV Reflecting voltage [25]. A saturated solution of cyano-4-hydroxy-cinnamic acid in 50% acetonitrile and 0.1% trifluoroacetic acid was used as the matrix. Sample preparation for the mass spectrometry followed the protocol reported by Geng and colleagues [25]. One μL of the matrix solution and one μL sample solution were added in the Score 384 target well. The mass spectrometry analysis was performed by a specialist in the Instrument Application Center of the Academy of Military Medical Sciences following the previously published protocol [25]. In brief, mass accuracy for peptide mass finger-prints (PMF) analysis was calibrated with a 0.1–0.2 Da external standard, and internal calibration was carried out with enzyme autolysis peaks, at a resolution of 12,000. The nanoLC-FT ICR MS was performed on an APEX-Q FT-ICR tandem mass spectrometer (Bruker Daltonics, Germany) equipped with a 9.4 T superconducting magnet (Magnex Scientific, UK) and an infinity cell. The trypsin digested peptides were sequenced by auto MS mode with MS/MS boost function. The FT-ICR mass spectra were processed using Data Analysis 3.4 software (Bruker Daltonics GmbH, Germany) as a gateway to set up database searches.

Expression and purification of recombinant Omps

To express OmpA and Smlt4123 proteins in *E. coli*, the Omp genes were first amplified by PCR and then cloned into the pET-30a (+) vector (Takara, Japan). The PCR primers were synthesized by Sangon Biotech (Shanghai, China) Co., Ltd. The primer sequences and cloning sites in pET-30a (+) are displayed in Additional file 1: Table S1. The recombinant vector was transformed into *E. coli* BL21 (DE3). The gene insertion in the recombinant vector was confirmed by DNA sequencing. The 6X-His tagged proteins expressed by the transformed *E. coli* BL21 were purified using Ni-agarose affinity chromatography according to the manufacturer instructions (GE Healthcare, USA). Endotoxin was removed from the Omp elution by using Triton-114 [26]. The Omp elution was analyzed by MALDI-TOF-MS (Bruker Daltonics, Germany) operating in the reflectron mode with 20 kV accelerating voltage and 23 kV Reflecting voltage [25].

Preparation of mouse anti-Omp serum

Omp elution was injected into female BALB/c mice (6-week old) (5 μg/time) with a 2-week interval between injections. The first injection was emulsified with Freund’s complete adjuvant (Sigma), and the remaining two injections were emulsified with Freund’s incomplete adjuvant.
μ37 °C, 350 post-immune serum. After incubation for 30 min at pre-immune serum was added in place of the mouse post-immune serum, for the control reaction, mouse peritoneal inoculation of 10^7 CFU immunization. In our pilot study, we found that intraperitoneal injection of S. maltophilia a mixture of PBS and adjuvant. To establish a mouse infection of S. maltophilia genes are displayed in Additional file 1: Table S3. Female BALB/c mice (6-week old) were randomized into two groups (n = 10 in each group): Smlt4123-immuned group and control group. Mice in the Smlt4123-immuned group were received a subcutaneous immunization of 5 μg Smlt4123 in PBS and was emulsified with the same volume of Freund’s complete adjuvant (CFA, total volume 100 μL/mouse) for primary immunization and Freund’s incomplete adjuvant (IFA) for boosting on days 14 and 28. Mice in the control group were immunized similarly with incomplete adjuvant (IFA) for boosting on days 14 and 28. 100 μL/mouse) for primary immunization and Freund’s incomplete adjuvant (IFA) for boosting on days 14 and 28. Mice in the control group were immunized similarly with incomplete adjuvant (IFA) for boosting on days 14 and 28. Microbiological examination of tissue and blood samples from healthy individuals appear to be powerful immunogens. In addition, we further identified the antibody subtype in the anti-Smlt4123 serum collected 2 weeks after the final immunization with S. maltophilia were grown in LB at 37 °C shaker to reach OD_{600} = 1.0. Cells were collected, washed twice with PBS, and re-suspended in PBS at the 2.0 × 10^8 CFU/mL for the intraperitoneal injection. Mice in the Smlt4123-immuned group and the control group were injected intraperitoneally with 200 μL S. maltophilia suspension (4.0 × 10^8 CFU) [27]. Eight hours after the injection, the mice were sacrificed by cervical dislocation. Mouse blood, liver, spleen, lung, and kidney were harvested and the tissues were homogenized in sterile PBS. Blood samples and homogenized tissue samples were cultured to determine bacterial burden.

Statistical analyses
Continuous variables are presented as mean ± standard deviation (SD). Two-group comparison was analyzed using Student’s t-test. Wilcoxon signed rank test was used for paired comparisons and Mann Whitney test was used for unpaired comparisons. P value was 2-sided and P < 0.05 was considered statistically significant.

Results
Identification of Omps
Two dimensional (2D) gel electrophoresis of the Omp preparation showed several protein spots (Fig. 1a). All of the protein spots were analyzed by MALDI-TOF-MS and nanoLC-FT ICR MS/MS. The mass spectrometry analysis identified 5 proteins from 10 protein spots (Fig. 1a and Additional file 1: Table S2). The 5 proteins are OmpA (protein spot 1 and 2), outer membrane Omp family protein (Smlt4123) (protein spot 3 and 4), Omp TolC (protein spot 5), TonB dependent receptor (protein spot 6–9), and fatty acid transport system membrane protein (protein spot 10). OmpA and Smlt4123 genes were amplified by PCR and sequenced. The DNA sequences of the OmpA and Smlt4123 genes are displayed in Additional file 1: Table S3. Western blot analysis of the 2D gel revealed that OmpA had the strongest reaction to the rabbit anti-S. maltophilia serum whereas Smlt4123 reacted with the rabbit serum weakly (Fig. 1b). The rabbit pre-immune serum did not react with the protein spots on the 2D gel.

Titer evaluation of mouse anti-Omp serum
Recombinant OmpA and Smlt4123 proteins produced from bacteria were used as antigen to immunize mice. ELISA showed that compared with the serum from control mice that were injected with the elution buffer, the mouse serum collected 2 weeks after the final immunization with the recombinant OmpA and Smlt4123 contained significantly increased levels of anti-OmpA and anti-Smlt4123 antibodies (P < 0.05, Fig. 2a). The serum was diluted at 1:102400 for ELISA, representing a titer greater than 1 × 10^5 and also suggesting that OmpA and Smlt4123 appear to be powerful immunogens. In addition, we further identified the antibody subtype in the anti-Smlt4123...
serum. Compared with the levels of IgG1 and IgG2a in the pre-immune serum, the IgG1 levels were significantly higher ($P < 0.05$), whereas the IgG2a levels were similar in the post-immune serum (Fig. 2b), indicating that anti-Smlt4123 antibody subtype may be IgG1 and recombinant Smlt4123 may induce Th2 immune response in the mice. We also identified the antibody subtype in the anti-OmpA serum. The result showed that compared with those in the pre-immune serum, the levels of both IgG1 and IgG2a were increased significantly after immunization with recombinant OmpA (Additional file 1: Figure S1).

**Potency evaluation of mouse anti-Omp serum**

OPKA was performed to evaluate the potency of the mouse anti-Omp serum in vitro. The mouse anti-Smlt4123 serum significantly reduced *S. maltophilia* counts in healthy individuals’ blood ($P < 0.05$, Fig. 3a). In contrast to the anti-Smlt4123 serum, the mouse anti-OmpA serum did not affect *S. maltophilia* counts in the blood of healthy individuals (Fig. 3b). These data suggest that mouse anti-Smlt4123 antibody may have some protective effects against *S. maltophilia* infection. We used the Smlt4123 protein sequence as a query and the Basic Local Alignment Search Tool to search genome sequences of 11 other *S. maltophilia* strains.
maltophilia strains. Of the 11 strains, 4 have a protein with 100% amino acid sequence similarity to the Smlt4123 sequence; 6 have a protein showing >90% amino acid sequence similarity to the Smlt4123 sequence; one showed no homologue. These results suggest that Smlt4123 may be highly conserved among S. maltophilia strains.

Assessment of protective effects of immunization with recombinant Smlt4123 against S. maltophilia infection

We then tested the protective effects of immunization with recombinant Smlt4123 on bacterial loads in a mouse model of S. maltophilia-induce infection. The bacterial loads in the blood, liver, spleen, lung, and kidney were compared in the control mice versus the mice that were immunized with recombinant S. maltophilia Smlt4123. Eight hours after infection with $4.0 \times 10^8$ CFU S. maltophilia, bacterial loads in the blood were significantly lower in the vaccinated mice than in the control mice, whereas bacterial loads in the other organs were similar between the two groups (Fig. 4).

Discussion

Omps are highly abundant proteins of Gram-negative pathogens and can interact with the host immune system directly. In the current study, we used mass spectrometry and immunoproteomic technology to screen immunogenic Omps of S. maltophilia and successfully identified Smlt4123 and OmpA as immunogenic proteins.

Previous studies have demonstrated that bacterial Omps can be promising vaccine candidates against bacterial infection. Hamid and Jain have identified a strong immunogenic Omp from Salmonella enterica serovar Typhimurium and demonstrated that the Omp can induce humoral and cell-mediated immune responses in rats [28]. They have also found that the vaccination of the Omp can protect rats against typhoid [28]. McConnell and colleagues have developed an Omp complex vaccine containing multiple bacterial membrane antigens of Acinetobacter baumannii [18]. They have tested the Omp complex vaccine in a murine sepsis model and found that the vaccine can evoke humoral and cellular responses, decrease post-infection bacterial loads and serum levels of pro-inflammatory cytokines, and protect mice from Acinetobacter baumannii infection [18]. Recently, Olsen and colleagues have developed a vaccine construct based on the VD4s region from the Chlamydia trachomatis major outer membrane protein and demonstrated that the vaccine construct can protect mice against Chlamydia trachomatis challenge, reduce bacterial loads in mouse vagina, and prevent pathogenesis in mouse upper genital tract [29]. Wang and colleagues have shown that Omps are involved in inducing host defense against S. maltophilia infection in channel catfish [30].

OmpA is abundantly expressed in Edwardsiella tarda and highly immunogenic in rabbit and fish, and vaccination of common carp with recombinant OmpA protects the fish from Edwardsiella tarda infection and increases the fish survival [20]. In contrast to the report by Maiti and colleagues [20], the current study found that mouse anti-recombinant OmpA serum did not affect S. maltophilia counts in the OPKA although the recombinant OmpA appeared to be strongly immunogenic in mice. However, whether the recombinant OmpA could protect mice from S. maltophilia challenge remains unclear. Further investigation is required to clarify whether OmpA could also be a potential vaccine candidate against S. maltophilia infection.

The current study shows that vaccination with recombinant Smlt4123 significantly reduced post-infection bacterial loads in mouse blood but did not affect the bacterial loads in mouse organs 8 h after the mice were challenged with S. maltophilia. The relatively short-time exposure to the S. maltophilia challenge may explain why the vaccination with recombinant Smlt4123 only reduced
the bacterial loads in mouse blood but not in other organs. It is possible that we could observe reduced bacterial loads in the tissues of liver, spleen, lung, and kidney in mice immunized with recombinant Smlt4123 if we could extend the observation time longer than 8 h after the infection challenge. Previous studies have showed that bacterial loads in blood and organs of mice immunized with Omp were decreased significantly at post-infection 12, 14, 24, or 48 h compared with those of the mice without immunization [19, 31, 32]. Therefore, we determined bacterial loads in liver, spleen, lung, and kidney of PBS immunized and Smlt4123 immunized mice at different time points. The result showed that 4, 8, 24 h post-infection no differences in bacterial loads were observed between groups of mice without immunization [19, 31, 32].

Conclusions

Our findings indicated that the immunoproteomic approach was an efficient way to screen the immunogenic protein of S. maltophilia. Besides, we found that the mouse anti-recombinant Smlt4123 serum enhanced opsonophagocytic killing activity of blood. We also showed that mice receiving vaccination with recombinant Smlt4123 had significantly lower S. maltophilia loads in the blood than mice without the vaccination, whereas no differences in bacterial loads were observed between groups of mice in liver, spleen, lung, and kidney. This result suggested that immunization of mouse with rSmlt4123 could provide a protection from bacteremia caused by S. maltophilia in the early stages.

Additional file

Additional file 1: Table S1. Primers used for cloning. Underlining indicates the recognition sites of restriction enzymes. Table S2. Identification of protein spots on 2D gel. Proteins identified by MALDI-TOF-MS and nanoLC-FT ICR MS/MS from S. maltophilia OMPs shown in Fig. 1a. Table S3. DNA sequences of Smlt0955 and Smlt4123 of S. maltophilia. OmpA and Smlt4123 genes were amplified by PCR and sequenced. Table S4. Determination of LD50. Fifty percent lethal dose (LD50) value of S.maltophilia to BALB/c mice was tested. Figure S1. IgG1 and IgG2a levels in the pre-immune and post-immune mice injected with...
Abbreviations
2D PAGE: Two-Dimensional Polyacrylamide Gel Electrophoresis; ELISA: Enzyme-linked immunosorbent assay; LB: Lysogeny broth; MALDI-TOF-MS: Matrix-assisted laser desorption/ionization time of flight mass spectrometry; Omps: Outer membrane proteins; PMF: Peptide mass finger-prints

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Authors' contributions
Conceived and designed the experiments: YL1, JY. Performed the experiments: GX, XT. Analyzed the data: XS, YL2, JW. Wrote the paper: YL1, GX. All authors read and approved the final manuscript.

Ethics approval and consent to participate
Written informed consent was obtained from Laboratory Animal Centre of the Academy of Military Medical Sciences. Ethics approval was obtained from the Review Board of the Academy of Military Medical Sciences, and the Affiliated Hospital of Academy of Military Medical Sciences. Written informed consent was obtained from the healthy volunteers.

Consent for publication
Not applicable

Competing interests
The authors declare that they have no competing interests.

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