Cytotoxic T Cell Adjuvant Effects of Three salmonella enterica Flagellins

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ABSTRACT

Bacterial flagellins are important virulence-associated factors and strong inducers of inflammatory responses in mammalian hosts. Flagellins have also been investigated as potential vaccine adjuvants, either for induction of humoral or cellular immune responses, to different target antigens. In this study we investigated the adjuvant properties of three Salmonella enterica flagellins types (FliC^d^, FliC^i^ and FljB) to an ovalbumin-derived CD8^+^ T cell-restricted epitope (OVA257–264). Although mice immunized with the three tested flagellins elicited antigen-specific activated CD8^+^ T cells, only animals immunized with FliC^i^ and FliC^d^ flagellins admixed with ovalbumin mounted specific in vivo cytotoxic responses to peptide-pulsed target cells. The present results indicate that Salmonella flagellins are endowed with type-specific adjuvant effects toward murine CD8^+^ T cells, a feature that may impact their use as adjuvants for prophylactic or therapeutic vaccines.

Key words: Salmonella, flagellin, cytotoxic T cells, adjuvants, vaccines.

INTRODUCTION

Flagellin, the structural subunit of flagellar filaments, contribute to the virulence of different pathogenic bacteria and promote activation of inflammatory responses in different mammalian hosts (17). The strong immunological effects of bacterial flagellins are mediated mainly by the specific binding to the toll-like receptor 5 (TLR 5) leading to signal transduction cascades that culminate with activation of transcription factors, production of pro-inflammatory mediators and activation of different cell types including antigen presenting cells (APC) such as macrophages and dendritic cells (13,20). Indeed, the immunomodulatory behavior of bacterial flagellins has also been explored as a potential vaccine adjuvant both for induction of specific humoral and cellular immune responses (4,7,9,11).

Salmonella enterica strains produce diseases ranging from mild enteritis to severe systemic infections in different animal hosts according to the specific strain serovar. Most Salmonella strains, such as S. enterica serovar Typhimurium (S. Typhimurium), are characterized by the phase variation phenomenon in which controlled expression of two or more flagellin-encoding genes results in the expression of alternate flagellar antigens, initially named as phase 1 (H1) and phase 2 (H2) flagellins but presently recognized as FliC and FljB flagellins, respectively (8). The variable nature of FliC and FljB flagellins together with the multiple lipopolysaccharide side-chains (O antigens), define more than 2,000 recognized Salmonella serovars (5). The natural diversity of Salmonella flagellins reflects mainly the polymorphism at the central portion of the protein, which creates sub-types such as illustrated by FliC flagellins expressed by S. Typhimurium (FliCi), S. Typhi (FliCd) and S. Muenchen (FliCd) (6).

Although several reports have investigated the adjuvant effects of specific Salmonella flagellins (4,7,11,16) a comparison of the adjuvant effects of different Salmonella flagellins types to CD8^+^ T cells have not been reported so far. Therefore, the objective of the present study was the investigation of the adjuvant effects of three Salmonella flagellins.
enterica flagellins, including two expressed by S. Typhimurium (FliC and FljB) and one originally encoded by S. Muenchen (FliCd), toward a CD8+ T cell-specific epitope derived from ovalbumin. The results indicated that, although the three tested flagellin types activated antigen-specific CD8+ T cells, in vivo cytotoxic effects were elicited only in mice immunized with the FliCi and FliCd flagellins.

MATERIALS AND METHODS

Bacterial strains and growth conditions

S. Typhimurium SL3201 derivative strains were generated following insertion of transposons in fliCi (fliC::Tn10) or fljB (fljB::MudJ) genes (19). These strains produced only FliCi (phase 1) or FljB (phase 2) flagellins and were kindly supplied by Dr. Alison D. O’Brien at the University of the Health Sciences, Bethesda, Maryland, USA. S. Dublin SL5930 expressed a plasmid-encoded FliCd flagellin, originally derived from S. Muenchen, as previously described (14). S. Typhimurium and S. Dublin strains were routinely cultivated at 37°C for 6 h in Luria-Bertani (LB) broth (Difco Laboratories) supplemented with antibiotic under aerobic conditions (200 rpm in an orbital shaker) for flagellin extractions.

Purification of Salmonella flagellins

Flagellins expressed by S. Typhimurium (FljB or FliCi) and S. Muenchen (FliCd) were routinely extracted from 50 ml cultures. Cells were collected by centrifugation, suspended in 2 mL of phosphate-buffered saline (pH 7.4) and sheared for 2 min in bench mixer (Genie 2) at maximal speed. Cells were kept in ice bath for 1 min and the procedure was repeated 3 times. The cell suspensions were then centrifuged at 10,000 X g for 10 min to remove the bacterial cells. The culture supernatants containing the sheared flagellins were precipitated with acetone, the resulting pellet suspended in PBS and, finally, submitted to heat treatment (65°C for 30 min) to depolymerize the filaments into flagellin monomers. Protein concentration was determined using BCA assay (Pierce) and protein purity checked by SDS-PAGE. Removal of contaminating lipopolysaccharide was accomplished with Detoxi-Gel column in accordance to the manufacturer’s instructions (Pierce). Endotoxin levels were determined using the Chromogenic Limulus Amebocyte Lysate assay (Cambrex Bio Science) as directed by the manufacturer. The determined endotoxin concentrations in the purified flagellin preparations were always bellow 0.125 EU/µg.

SDS-PAGE and immunoblot analyses

SDS-PAGE was performed following standard procedures using a Mini Protean II vertical electrophoresis unit (Bio-Rad). Proteins sorted in 12% polyacrylamide gels were stained with Comassie Blue or transferred to nitrocellulose sheets (0.45 µm pore size) at 200 mA for 1 h. After overnight blocking with 1% (w/v) bovine serum albumin (BSA) in phosphate-buffered saline (PBS) at 4°C, nitrocellulose sheets were developed for reactive protein bands with different antibodies, according to previously published procedures (18).

Immunization regimens

Female C57BL/6 mice at 8 to 12 weeks of age were supplied by the Isogenic Mouse Breeding Facility of the Department of Parasitology, Institute of Biomedical Sciences, São Paulo University (USP). All procedures were carried out in accordance with the principles of the Brazilian code for the use of laboratory animals and were approved by the Ethics Committee on Use of Laboratory Animals from the Institute of Biomedical Sciences, USP. Animals were immunized with sterile phosphate buffered saline (PBS), ovalbumin (OVA) (12 µg) (Sigma-Aldrich) alone or with each purified FliCd, FliCi or FljB flagellins (5 µg). All immunizations were administered subcutaneously (s.c.) in a 100 µl volume on days 0 and 21. Antigen-specific T cell responses were evaluated seven days after immunization.

Antigen-specific ELISPOT assays

The ELISPOT assay for quantification interferon-gamma (IFN-γ) producing cells was performed essentially as described earlier (7, 10, 12). CD8+ T cells (105) recovered from spleens immunized animals were added to microplate wells (MultiScreen, Millipore), previously coated with 10 µg/mL anti-IFNα capture antibody (BD Pharmingen) diluted in PBS. CD8+ T cells were enriched (>97%) using magnetic beads on an AutoMacs cell separator (Miltenyi Biotec.). T cells were then incubated overnight with naive antigen presenting cells (106 cells/well) in the absence or presence of the OVA-derived H-2Kk restricted synthetic nonapeptide OVA254-267 (SIINFEKL), purchased at PSL (Heidelberg, Germany) and employed at a final concentration of 0.5 µg/ml. Anti-CD3 (BD Pharmingen) was used as a positive control at a final concentration of 0.25 µg/ml. Plates were incubated overnight at 37 ºC in the presence of 5% CO2. After incubation, the cells were washed and incubated with 2 µg/mL of biotinylated anti-mouse IFN-γ (BD Pharmingen) diluted in PBS and 0.05% Tween 20 (PBS-T). Plates were washed and incubated with streptavidin peroxidase conjugate (Sigma) diluted 1:800 in PBS. Spot development was performed as previously described (12). Antigen-specific response was performed in duplicate and quantified using an automated ImmunoScan reader (Cellular Technology Ltd.). Data is represented as the number of IFN-γ secreting cells/106 cells.

In vivo cytotoxicity assays

Splenocytes from naive mice were stained with 1 µM or 10 µM of carboxyfluorescein diacetate succinimidyl ester (CFSE) (Invitrogen). The cells labeled with 10 µM CFSE were pulsed
for 40 min at 37°C with 1 µM of the OVA254-267 peptide. Another cell aliquot was labeled with 1 µM CFSE but was not pulsed with the target peptide. Both cell populations were transferred i.v. (2x10^7 cells/each) to immunized mice. One day later, spleens were isolated and the two cell populations were monitored by flow cytometry in a multivariant FACScan analyser (FACSCalibur, BD Biosciences). The percentages of target specific cell killing were determined as previously described (2,21).

Cytokine ELISA
Aliquots containing 10^7 spleens cells were cultured overnight with or without the OVA254-267 peptide. The culture supernatants were collected after 48 h and stored at -80°C until testing. IFN-γ concentration were determined by sandwich ELISA using duplicate samples and a specific kit, according to instructions of the manufacturer (BD Pharmingen).

Statistic analysis
The data were analyzed by one-way ANOVA test. Values of p < 0.05 were considered to be statistically significant.

RESULTS
The adjuvant effects of the three tested Salmonella flagellins (FliCd, FliCi and FljB) were evaluated with purified proteins extracted from two recombinant monophasic S. Typhimurium strains and one recombinant S. Dublin strain expressing a plasmid-encoded S. Muenchen FliCd flagellin (Fig. 1). Each purified flagellin was admixed with OVA and s.c. delivered in two doses to C57BL/6 mice. The OVA-specific T-cell responses were measured in ELISPOT assays using the H-2Kb-restricted CD8+ epitope (OV A257-264) and purified CD8+ T cells derived from spleens of vaccinated mice. The results indicated that the three tested Salmonella flagellins activated OVA-specific T-cells responses, such as measured by the number of antigen-specific IFN-γ-secreting CD8+ T cells (Fig. 2). The numbers of activated CD8+ T cells in mice immunized with OVA admixed with one of the three tested flagellins were not statistically significant among them. These results indicated that the three tested Salmonella flagellins are endowed with similar adjuvant effects toward immune responses mediated by activated antigen-specific CD8+ T cells, as measured by the number of IFN-γ secreting cells in ELISPOT assays.

We also measured the secretion of IFN-γ by spleen cells recovered from mice immunized with OVA and one of the three tested Salmonella flagellins. As indicated in Fig. 3, the amount of IFN-γ produced by cells derived from mice immunized with FliCd and stimulated with the OVA254-267 peptide was higher than...
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those detected in cells derived from mice immunized with FliC or FljB flagellins. The in vivo activities antigen-specific CD8+ T cell response were determined following inoculation of CFSE-labeled OVA254-267 pulsed cells into in mice immunized with OVA admixed with different Salmonella flagellins. The results presented in Fig. 4 shows that mice immunized with OVA and FliCd or FliCi flagellin elicited similar activation of antigen-specific CD8+ T cell-dependent cytotoxic responses. In contrast, mice immunized with OVA and FljB did not induce any significant specific cellular lysis, as measured in vivo cytotoxicity assay (Fig. 4). This result indicates that the tested Salmonella flagellins expressed differential adjuvant effects toward cytotoxic responses mediated by CD8+ T cells.

DISCUSSION

In spite of the enormous impact of vaccines in the control, and eventually eradication, of infectious disease vaccines are still not available for many of infectious diseases that still scourge millions of people around the world. Diseases associated with invasive pathogens require activation of strong T cell responses for the complete and sustained eradication of the pathogen, a feature not usually achieved by the presently available subunit vaccine formulations. The discovery of mammalian Toll-like receptors (TLR) and their corresponding microbial agonists, has opened up renewed perspectives for the development of alternative adjuvants modulating the desired type of immune responses including more effective activation of CD8+ T cell-dependent responses. In the present study, we investigated the differential adjuvant effects of TLR5-specific ligands, three Salmonella enterica flagellins, with regard to activation of specific CD8+ T cell responses. The present data indicate that, although all tested flagellins can elicited CD8+ T cell responses in vaccinated mice, only mice immunized with FliC flagellins elicited significant in vivo cytotoxic responses.

The results based on ELISPOP assays demonstrated that the three tested flagellins types activate similar OVA-specific CD8+ T cell responses, further confirming the previously reported adjuvant effects of bacterial flagellins on either CD4+ and CD8+ T lymphocytes (4,7,9,11,18). However, detection of secreted IFN-γ by spleen cells incubated with the CD8+ restricted OVA254-267 peptide indicated that mice immunized with OVA and FliCd developed enhanced CD8+ T cell responses with regard to mice immunized with OVA admixed with FliCi or FljB as vaccine adjuvants.

The differential adjuvant effect of the tested Salmonella flagellins was further supported by the in vivo cytotoxic
responses. In contrast to the ELISPOT results, in which the three tested flagellins exerted a clear adjuvant effect to the cytotoxic responses toward target cells pulsed with the OVA\textsubscript{254-267} peptide. Based on the present results, we conclude that FliCd and FliCi flagellins express enhanced adjuvant effects to cytotoxic responses in vaccinated mice.

Since the *Salmonella* flagellins differ mainly at the central hypervariable domain, which are not specifically involved in the binding of the TLR5 receptor, the differential immunological effects of FliC and FljB may not be explained solely by the recognition of the cognate TLR receptor. Indeed the recent finding that bacterial flagellins bind to a family of Nod-like receptors, such as Naip5 and Ipaf, in which flagellin-binding sites are still unknown, suggests that amino acid residues outside the strongly conserved N- and C-terminal ends participate in modulation of inflammatory responses elicited by *Salmonella* flagellins (1,23).

The potent adjuvant effects of *Salmonella* flagellins, both for activation of antigen-specific humoral and cellular immune responses, has received considerable attention by several research groups dealing with vaccine development. Flagellins have been employed either as an epitope carrier expressed by attenuated *Salmonella* vaccine strains (14,15,22) as purified adjuvants admixed with different soluble antigens (11,16), or as a purified hybrid protein genetically fused to the target antigen (4,7). So far, the specific mechanisms by which bacterial flagellins improve the immune responses to co-delivered antigens is unclear but trafficking and processing of antigens following endocytosis by antigen presenting cells is probably an important aspect (3). Such adjuvants effects would be better reasoned for antigens genetically fused to flagellin but other mechanisms may also contribute to activation and maturation of antigen presenting cells exposed simultaneously to the target antigen and soluble flagellin monomers. The question concerning the better performance of *Salmonella* flagellins following physical coupling to the target antigen is still open but our results shows that activation of antigen-specific CD8\(^+\) T cells as well as *in vivo* cytotoxic effects can be achieved in animals immunized without the need to fuse both antigen and adjuvant either by chemical or genetic approaches.

The finding that toll-like receptors ligands stimulate strong T cell-dependent immune responses opened up renewed perspectives for the development of therapeutic vaccines against chronic infectious diseases. The present study represents a further demonstration that *Salmonella* flagellins may have an important role on the field of vaccine development not only as an adjuvant for conventional antibody-dependent prophylatic vaccines but also as a component for vaccine formulation aiming eradication of invasive pathogens requiring activation of CD8\(^+\) T cell-dependent immune responses.

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**RESUMO**

**Efeitos adjuvantes para células T citotóxicas de três flagelinas de *Salmonella enterica***

As flagelinas bacterianas são importantes fatores associados à virulência e potentes induutores de resposta inflamatória em mamíferos. Estas moléculas são também investigadas como potencial adjuvante para uso em vacinas na indução de resposta imune humoral e celular para diferentes antígenos alvo. No presente estudo investigamos as propriedades adjuvantes de três tipos de flagelinas de *Salmonella enterica* (FliCd, FliCi e FljB) para um epitope derivado da ovalbumina específico para células T CD8\(^+\). As três flagelinas testadas induziram respostas de células T CD8\(^+\) específicas em camundongos imunizados, porém, somente animais imunizados com as flagelinas FliCi e FliCd co-administradas com ovalbumina montaram resposta citotóxica específica *in vivo* para células-alvo pulsadas com peptide OVA. Os resultados apresentados indicam que flagelinas de *Salmonella* são dotadas de efeitos adjuvantes tipo-específico frente a células T CD8\(^+\) *in vivo*, uma característica que pode gerar impactos no uso dessas proteínas como adjuvantes em vacinas profiláticas ou terapêuticas.

**Palavras-chave:** *Salmonella*, flagelina, células T citóxicas, adjuvantes, vacinas.

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