**Bordetella bronchiseptica** is a potent and safe adjuvant that enhances the antigen-presenting capability of dendritic cells

You-Jeong Lee¹, Yong Han¹, and Hong-Gu Joo¹,²,*

¹College of Veterinary Medicine, ²Veterinary Medical Research Institute, Jeju National University, Jeju 63243, Korea

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*Correspondence  
Hong-Gu Joo  
E-mail: jooh@jejunu.ac.kr

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**ABSTRACT**  
We previously demonstrated that *Bordetella bronchiseptica* (B. bronchiseptica) antigen (Ag) enhances the *Mycoplasma hyopneumoniae* Ag-specific immune response. The focus of this study was whether acellular bacterin of *B. bronchiseptica* could be used as an adjuvant to increase antigen-presenting capability of dendritic cells (DCs) by increasing the level of activation. The metabolic activity of DCs was increased by *B. bronchiseptica*, similar to lipopolysaccharide (LPS). Flow cytometry analysis revealed that *B. bronchiseptica* increases the expression of major histocompatibility complex class-2, cluster of differentiation (CD)40, CD54, and CD86 which are closely related to DC-mediated immune responses. *B. bronchiseptica* enhanced the production of cytokines related to adaptive immune responses. Furthermore, the survival rate of *B. bronchiseptica*-injected groups was 100% at 15 and 20 mg/kg doses, whereas that of LPS-injected groups was only 20%, 0% at 15 and 20 mg/kg doses respectively, and so *B. bronchiseptica* is likely to be safer than LPS. Taken together, these results indicate that *B. bronchiseptica* can be used as an adjuvant to enhance the antigen-presenting capability of DCs. *B. bronchiseptica* is a candidate for producing vaccines, especially in case of DC-mediating efficacy and safety demands. This study provides researchers and clinicians with valuable information regarding the usage of *B. bronchiseptica* as a safe bacteria-derived immunostimulating agent for developing efficient vaccines.

**INTRODUCTION**

Vaccines are used to obtain an acquired immunity against certain pathogens [1]. A vaccine contains antigen (Ag) and immunostimulating agent. Vaccine Ags include disease-causing weakened or killed microorganisms, its toxins, or one of its surface proteins [1]. However, some vaccine Ags do not have enough intrinsic capability to stimulate immune system, and thus the vaccines need to contain the immunostimulating agents, called adjuvants [2]. Vaccine adjuvant is added to enhance the efficacy of vaccine at a variety of steps. It extends the presence of Ag at injection sites and helps antigen-presenting cells (APCs) to uptake Ag, and also stimulates immune responses by the increase of cytokine production and immune-related molecules on the surface of APCs [3]. Vaccine adjuvants have been divided into inorganic substances including alum, cytokines, and bacterial products according to their source types [4]. *Bordetella pertussis* (*B. pertussis*) and *Mycobacterium bovis* (*M. bovis*) have been used as bacteria-derived immunostimulating agents, partly vaccine adjuvants [5,6]. *M. bovis* is the causative agent of tuberculosis in cattle and *Bacillus Calmette-Guerin* (BCG) is prepared from an attenuated strain of *M. bovis*. BCG has been used as a vaccine adjuvant, but its side effects include fibrosis and granuloma, and thus it is not used currently [6-8]. *B. pertussis* is the causative agent of pertussis or whooping cough, and its virulence factors include pertussis toxin, filamentous hae...
magglutinin, pertactin, and tracheal cytotoxin. Pertussis toxin is a major toxin of *B. pertussis* and has an adjuvant effect such as antibody response [5,9], but can generate the sensitization leading to anaphylaxis [10].

*Bordetella bronchiseptica* (*B. bronchiseptica*) is a similar pathogen to *B. pertussis* and causes respiratory disease [11]. Whereas *B. pertussis* is relatively limited to human, *B. bronchiseptica* affects several animals including dogs, cats, and pigs with different strains [12]. Infection of *B. bronchiseptica* alone does not seriously damage to respiratory system of hosts, but the combined infection with other pathogens can cause significant economic loss [13,14]. *B. bronchiseptica* modulates the phenotypes of macrophages, leading to the inhibition of CD4⁺ T cell proliferation [15].

In a previous study, we demonstrated that *B. bronchiseptica* Ag can enhance the production of *Mycoplasma (M.)* hyopneumoniae Ag-specific immunoglobulin G [16]. In this study, we investigated whether formalin-fixed whole-cell and acellular vaccine of *B. bronchiseptica* can improve the Ag-presenting capability of dendritic cells (DCs) and may be a candidate of vaccine adjuvant with immunostimulatory activity.

**METHODS**

**Animals and materials**

C57BL/6 and Balb/c mice were obtained from OrientBio (Seongnam, Korea) and maintained at our animal facility. 8 to 12-week old mice were used for all experiments. Animal experiments in this study were performed in accordance with the Institutional Guideline for Animal Use and Care of Jeju National University (2018-0011, 2018-0045). Lipopolysaccharide (LPS; Sigma-Aldrich, St Louis, MO, USA) was used to as a positive control of immunostimulatory activity.

In order to minimize the interference of lymphocytes and granulocytes in DCs, the floating cells were removed on the 2nd, 4th day and DC precursor cells attached to the bottom were cultured. After 6 day culture, the floating cells were harvested and used as DCs.

**Measurement of DC metabolic activity**

To measure the metabolic activity of DCs, the cells were seeded in 96-well plates and treated with LPS or *B. bronchiseptica* for 3 days. Then, 5 µl/well Cell counting-kit-8 (CCK-8; Dojindo Molecular Tech., Kumamoto, Japan) solution was added and incubated at 37°C for 4 h. The optical density was measured at 450 nm with a microplate reader (Multiskan FC; Thermo Fisher Scientific, Waltham, MA, USA).

**Measurement of cytokines**

DCs were cultured in 96-well plates to analyze the production of cytokines such as tumor necrosis factor (TNF)-alpha, interleukin (IL)-10, and IL-12. The cells were treated with LPS or *B. bronchiseptica* for 3 days. The supernatants of treated cells were collected and used for Enzyme-linked immunosorbent assay (ELISA). The amounts of TNF-alpha, IL-10, and IL-12 were determined by CytoSet kits (Thermo Fisher Scientific) according to the instructions. The optical density was measured at 405 nm.

**Mixed lymphocyte reaction (MLR)**

To evaluate the adjuvanticity of *B. bronchiseptica*, the cultured DCs were treated with LPS, *B. bronchiseptica* for 2 days, and then co-cultured with allogeneic spleen cells. Allogeneic spleen cells were harvested from Balb/c mice. The treated DCs were incubated with 50 µg/ml mitomycin C for 30 min before the co-culture setup. The DCs and allogeneic spleen cells were cultured in 96-well plates at a concentration of 1 × 10⁶ cells/ml and 2 × 10⁶ cells/ml, respectively. As a control, only allogeneic spleen cells were cultured. After 5 days, 10 µl/well of CCK-8 solution was added and incubated at 37°C for 4 h. The optical density was measured...
at 450 nm with a microplate reader.

**Injection of LPS or B. bronchiseptica into mice**

7-week old Balb/c mice were divided into 6 groups. B. bronchiseptica or LPS was injected at 10, 15 and 20 mg/kg doses respectively, by a single intraperitoneal injection. The mice were monitored for one week to check mortality rate and weight change.

**Statistical analysis**

The data of CCK-8 assay and ELISA are presented as mean ± standard deviation and were analyzed with one-way analysis of variance, Tukey’s multiple comparison test (InStat; GraphPad Software, San Diego, CA, USA). A p-value of < 0.05 was considered as significant. **, *** indicate p < 0.01, 0.001 respectively, compared to control.

**RESULTS**

**Effect of B. bronchiseptica on the metabolic activity of DCs**

In order to determine whether B. bronchiseptica affects the metabolic activity of DCs, we performed CCK-8 assays. DCs were treated with B. bronchiseptica or LPS for 3 days at a range of concentration, 0 to 300 ng/ml, and the metabolic activity of DCs was measured. B. bronchiseptica at 33.3 and 100 ng/ml significantly increased the metabolic activity of DCs compared to control (0 ng/ml) (Fig. 1). The highest increase of metabolic activity was appeared at 100 ng/ml, an optimal concentration used for DC activation in this study.

**B. bronchiseptica increases the expression of immune response-related surface markers of DCs**

To determine whether B. bronchiseptica induces the activation and maturation of DCs, we measured the expression level of surface markers on DCs. The cells were treated with 0.1 µg/ml of B. bronchiseptica or LPS for 2 days and stained with surface marker-specific Abs. B. bronchiseptica strongly increased the surface markers expression on DCs, which increase the MHC class 2 molecule by 67% compared to control, and also other 3 molecules more than 150% (Fig. 2). This data showed that LPS also had effects on the activation of DCs, but B. bronchiseptica increased the expression level of the 4 surface markers higher than LPS.

**B. bronchiseptica decreases the antigen uptake capability of DCs**

Antigen uptake assay was performed to confirm whether B. bronchiseptica induces the maturation of DCs. The cells were incubated with FITC-dextran at 37°C for 1 h. Both LPS and B. bronchiseptica showed lower levels of the uptake of FITC-dextran compared to that of control (Fig. 3). The uptake of FITC-dextran in DCs at 4°C was used as negative control.

**B. bronchiseptica enhances the cytokine production of DCs**

To determine the effect of B. bronchiseptica as an adjuvant for vaccines, we measured the production of TNF-alpha, IL-10, and IL-12 in DCs treated with B. bronchiseptica or LPS. ELISA results demonstrated that both B. bronchiseptica and LPS increased the production of TNF-alpha, IL-10, and IL-12 (Fig. 4). In the case of TNF-alpha, B. bronchiseptica produced less TNF-alpha than LPS at 0.02 µg/ml, but produced more at 0.1 and 0.5 µg/ml. Interestingly, IL-10 production was significantly elevated in B. bronchiseptica-treated DCs compared to LPS-treated DCs at 0.5 µg/ml. B. bronchiseptica enhanced the IL-12 production of DCs higher than LPS in all concentrations.

**Effect of B. bronchiseptica on the antigen-presenting capability of DCs**

To evaluate whether B. bronchiseptica can increase the antigen-presenting capability of DCs, we co-cultured the treated DCs and allogeneic spleen cells. DCs were pre-treated with B. bronchiseptica or LPS at 0.1 µg/ml for 2 days, and then co-cultured at a ratio of 1:20 with allogeneic spleen cells harvested from Balb/c mice. Mitomycin C was used to limit the metabolic activity of DCs itself and inhibit further maturation during the co-cultures. Both B. bronchiseptica- or LPS-treated DCs significantly increased the metabolic activity and proliferation of allogeneic spleen cells compared to control DCs (Fig. 5). This result demonstrated that B. bronchiseptica...
B. bronchiseptica can increase the antigen-presenting capability of DCs.

**B. bronchiseptica is safe in vivo**

To investigate if B. bronchiseptica is safe in vivo, Balb/c mice were injected with 10, 15, and 20 mg/kg of B. bronchiseptica or LPS. After 7 days, the survival rate and weight change were examined. All mice were survived in both B. bronchiseptica- or LPS-injected groups at 10 mg/kg dose. B. bronchiseptica-injected groups showed 100% survival rate at 15 and 20 mg/kg dose, whereas LPS-injected groups showed only 20%, 0% survival rate at 15 and 20 mg/kg dose respectively (Fig. 6). Interestingly, the body weight change of B. bronchiseptica-injected mice was less than 3% decrease. This result indicates that B. bronchiseptica is safer in vivo than LPS.

**DISCUSSION**

Adjuvants increase the effects of vaccines by enhancing antigen-specific immune responses, such as maximal T cell-dependent immune responses [3]. This effect of vaccine adjuvant can be achieved through several critical events including the activation of antigen-presenting cells such as DCs [17]. After uptaking antigens, DCs migrate to lymph nodes and present antigens to lymphocytes for initiating adaptive immune responses [18]. In previous studies, we demonstrated that B. bronchiseptica antigens
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increase the production of M. hyopneumoniae-specific IgG [16]. In this study, we investigated whether B. bronchiseptica can be used as an adjuvant to increase the antigen-presenting capability of DCs in several aspects.

This study demonstrated that B. bronchiseptica up-regulated the functions of DCs including metabolic activity. CCK-8 assay showed that B. bronchiseptica increased the metabolic activity of DCs at a level similar to that of LPS. Flow cytometry analysis showed that B. bronchiseptica stimulated the expression of MHC class II, an essential antigen-presenting molecule, on the surface of DCs and increased the expression of DC activation markers such as CD40, CD54, and CD86. In addition, the antigen uptake of B. bronchiseptica-treated DCs was consistently low. The matured DCs have much lower antigen uptake capability than immature DCs. Our data collectively indicates that B. bronchiseptica can increase the degree of activation and maturation of DC, similar to LPS.

TNF-alpha, IL-10 and IL-12 production were measured to determine how much B. bronchiseptica can activate DCs. ELISA showed that B. bronchiseptica strongly increased the production of TNF-alpha, IL-10, and IL-12. Our data indicate that B. bronchiseptica is effective as an adjuvant to activate DCs. Interestingly IL-10 production was significantly elevated in the DCs treated with B. bronchiseptica compared to LPS at 0.5 µg/ml. IL-10 is a representative anti-inflammatory cytokine and its production is closely connected to the production of inflammatory cytokine including TNF-alpha [19,20]. This result suggests that B. bronchiseptica may have a unique regulatory role in the production of inflammatory and anti-inflammatory cytokines.

The immune enhancing effect of B. bronchiseptica is similar to or higher than that of LPS. B. bronchiseptica does have less cytotoxicity than LPS on DCs. B. bronchiseptica is a Gram-negative bacterium containing LPS and is likely to act as an immune stimulating factor. It is also suggested that the LPS of B. bronchiseptica may have a different structure or content compared to
that of E. coli. In the future, we will study which components of B. bronchiseptica including LPS have the immunostimulatory effect.

In addition, we measured the effect of B. bronchiseptica on the antigen-presenting capability of DCs. MLRs measured by CCK-8 assay showed that B. bronchiseptica- and LPS-treated DCs increased the metabolic activity and proliferation of allogeneic lymphocytes. Our data indicate that B. bronchiseptica can enhance the antigen-presenting capability of DCs.

To confirm if B. bronchiseptica is safe in vivo, mice were injected with the same dose of B. bronchiseptica or LPS. Interestingly, B. bronchiseptica-injected mice did not die at all doses (10, 15, 20 mg/kg), whereas LPS-injected mice seriously died at 15 and 20 mg/kg (survival rate only 20%, 0% respectively). This result suggests that B. bronchiseptica is much safer in vivo than LPS.

Taken together, our data demonstrate that B. bronchiseptica enhances the antigen-presenting capability of DCs and has immunostimulatory activity as an adjuvant. In addition, the safety of B. bronchiseptica is better than LPS which is already used as a kind of adjuvant (Supplementary Fig. 1). This study shows the availability of B. bronchiseptica as a vaccine adjuvant based on its activity on DCs.

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CONFLICTS OF INTEREST

The authors declare no conflicts of interest.

SUPPLEMENTARY MATERIALS

Supplementary data including one figure can be found with this article online at http://pdf.medrang.co.kr/paper/pdf/Kjpp/Kjpp2020-24-01-06-s001.pdf.

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