Lipopolysaccharide Analogs Improve Efficacy of Acellular Pertussis Vaccine and Reduce Type I Hypersensitivity in Mice


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Pertussis is an infectious disease of the respiratory tract that is caused by the gram-negative bacterium *Bordetella pertussis*. Although acellular pertussis (aP) vaccines are safe, they are not fully effective and thus require improvement. In contrast to whole-cell pertussis (wP) vaccines, aP vaccines do not contain lipopolysaccharide (LPS). Monophosphoryl lipid A (MPL) and *Neisseria meningitidis* LpxL2 LPS have been shown to display immune-stimulating activity while exerting little endotoxin activity. Therefore, we evaluated whether these LPS analogs could increase the efficacy of the aP vaccine. Mice were vaccinated with diptheria-tetanus-aP vaccine with aluminum, MPL, or LpxL2 LPS adjuvant before intranasal challenge with *B. pertussis*. Compared to vaccination with the aluminum adjuvant, vaccination with either LPS analog resulted in lower colonization and a higher pertussis toxin-specific serum immunoglobulin G level, indicating increased efficacy. Vaccination with either LPS analog resulted in reduced lung eosinophilia, reduced eosinophil numbers in the bronchoalveolar lavage fluid, and the ex vivo production of interleukin-4 (IL-4) by bronchial lymph node cells and IL-5 by spleen cells, suggesting reduced type I hypersensitivity. Vaccination with either LPS analog increased serum IL-6 levels, although these levels remained well below the level induced by wP, suggesting that supplementation with LPS analogs may induce some reactogenicity but reactogenicity considerably less than that induced by the wP vaccine. In conclusion, these results indicate that supplementation with LPS analogs forms a promising strategy that can be used to improve aP vaccines.

Pertussis is caused by *Bordetella pertussis* infection of the respiratory tract and is among the 10 infectious diseases with the highest rates of morbidity and mortality worldwide. After introduction of whole-cell pertussis (wP) vaccines in the 1950s, the incidence of pertussis has decreased significantly. Although they are efficacious, wP vaccines were found to be reactogenic, leading to concerns about their safety in the 1970s. Therefore, acellular pertussis (aP) vaccines that comprise purified *B. pertussis* proteins have been developed. In many countries, pertussis has recently reemerged, despite the high rates of vaccine coverage (13). Several approaches to reducing disease incidence and severity have been suggested, one of them being the introduction of combination vaccines that contain aP vaccine instead of wP vaccine (28). Thus, while the reactogenicity of LPS precludes its use, its adjuvanticity is, regretfully, missed.

We and others have shown that LPS is an essential component of wP vaccines in mice, as wP-vaccinated C3H/HeJ mice that have a point mutation in the *Tlr4* gene, which results in defective signal transduction, failed to clear a *B. pertussis* challenge (21; H. A. Banus et al., unpublished data). This result underlines the important role of LPS in generating a productive immune response, at least in mice. Additionally, we have shown that a functional polymorphism in TLR4 was associated with reduced pertussis toxin (Ptx)-specific immunoglobulin G (IgG) titers in wP-vaccinated children 1 year of age (H. A. Banus et al., submitted for publication). Together, these findings strongly suggest an important role of LPS in wP vaccines.

To make use of this role of LPS, the development and use of LPS derivatives and novel LPS species have been investigated. The nontoxic LPS derivative monophosphoryl lipid A (MPL) engages TLR4 (17, 33), inducing Th1 adaptive immunity and changing Th2-directed responses to Th1-directed responses (3, 34, 38, 48). MPL combined with aluminum (denoted AS04) is registered for clinical use as an adjuvant in viral vaccines, such as the hepatitis B virus vaccine (6) and the human papillomavirus vaccine (18), while MPL combined with L-tyrosine is registered for clinical use as an adjuvant in allergy therapy (2, 22, 29, 39). Consequently, concerns have been raised with respect to the relative efficacies of aP vaccines compared with those of wP vaccines as well as those of simultaneously administered vaccines, such as diphtheria, tetanus, and *Hae- nomophilus influenzae* type b (Hib) vaccines. In fact, this concern has been substantiated by an increase in the incidence of invasive Hib disease in the United Kingdom that coincided with the distribution of combination vaccines that contain aP vaccine instead of wP vaccine (28). Thus, while the reactogenicity of LPS precludes its use, its adjuvanticity is, regretfully, missed.

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performed according to national and international guidelines. The diet consisted of ground standard laboratory chow (RMH-B; Hope Farms, Woerden, The Netherlands) or from Harlan (Horst, The Netherlands). The diet obtained from the breeding colony of the Vaccine Institute (Bilthoven, The Netherlands), 40 g filamentous hemagglutinin, and 8 µg pertactin; GlaxoSmithKline, Rixensart, Belgium) in 0.5 ml saline (1 human dose [HD]). The vaccine contained aluminum by MPL in a diphtheria-tetanus-aP vaccine would beneficially affect the vaccine in a mouse model system. In a second series of experiments, LpxL2 LPS was also included as an alternative adjuvant. The clearance of a B. pertussis challenge, the Ptx-specific serum IgG levels, the parameters of type 1 hypersensitivity, and serum interleukin-6 (IL-6) levels (elevated IL-6 levels suggest reactogenicity) were determined.

MATERIALS AND METHODS

Vaccines and adjuvants. (i) Series 1. The acellular (DTaP) vaccine was a combined vaccine consisting of diphtheria toxoid, tetanus toxoid, and a component aP vaccine (25 µg formaldehyde- and glutaraldehyde-detoxified Ptx, 25 µg filamentous hemagglutinin, and 8 µg pertactin; GlaxoSmithKline, Rixensart, Belgium) in 0.5 ml saline (1 human dose [HD]). The vaccine contained aluminum hydroxide as an adjuvant (<0.625 mg aluminum/HD).

The DTaP vaccine was supplemented with two adjuvants, aluminum adjuvant [2% Al(OH)_3 gel; Serva, Heidelberg, Germany] or MPL. To 1 HD DTaP vaccine, 2 ml aluminum-phosphate-buffered saline (PBS; 1/3 [vol/vol]) was added. MPL (from Salmonella enterica serotype Minnesota Re 595) was from Sigma-Aldrich, Zwijndrecht, The Netherlands. To 1 HD DTaP vaccine, 2 ml of 100 µg/ml MPL-PBS was added. The amount of MPL administered is less than the amount that showed no toxicological effects in a single-dose toxicity study (2).

(ii) Series 2. The same DTaP vaccine used for series 1 was used for series 2. The Pw vaccine was prepared from B. pertussis strain B213, a streptomycin-resistant derivative of strain Tohama I (23). The bacteria were grown in synthetic THU5 medium (43) for 68 h at 35°C while the mixture was shaken (175 rpm). The bacterial cell suspensions were heat inactivated for 10 min at 56°C in the presence of 8 mM formaldehyde, after which the cells were collected by centrifugation at 16,100 g for 10 min and resuspended in PBS to an A_490 of 2.5, i.e., 50 international opacity units per ml (1 HD/ml). The suspensions were stored at 4°C.

Before immunization, the DTaP and Pw vaccines were diluted in PBS to 1/10 HD, after which 3 mg/ml aluminum phosphate (Brenntag, Dordrecht, The Netherlands), 40 µg/ml MPL (Sigma-Aldrich), or 40 µg/ml N. meningitidis LpxL2 LPS (46) was added as an adjuvant.

Animals. Female BALB/c mice were used at 6 to 8 weeks of age. They were obtained from the breeding colony of the Vaccine Institute (Bilthoven, The Netherlands) or from Harlan (Horst, The Netherlands). The diet consisted of ground standard laboratory chow (RMH-B; Hope Farms, Woerden, The Netherlands). Food and water were given ad libitum. All animal experiments were performed according to national and international guidelines.

Vaccination. (i) Series 1. Mice received a subcutaneous (s.c.) injection of 1/5 HD DTaP vaccine in 0.5 ml aluminum, 1/5 HD DTaP vaccine in 0.5 ml MPL, 0.5 ml aluminum alone, or 0.5 ml MPL alone and 14 days before infection. In one experiment, mice received an s.c. injection of 1/5, 1/25, or 1/125 HD DTaP vaccine in 0.5 ml aluminum, 1/5, 1/25, or 1/125 HD DTaP vaccine in 0.5 ml MPL, 0.5 ml aluminum alone; or 0.5 ml MPL alone.

(ii) Series 2. Mice received an s.c. injection with 1/10 HD Pw vaccine in 0.5 ml aluminum, 1/10 HD DTaP vaccine in 0.5 ml aluminum, 1/10 HD in 0.5 ml LpxL2 LPS, 1/10 HD in 0.5 ml MPL, or 0.5 ml PBS 28 and 14 days before infection.

Bacterial strain and growth conditions. B. pertussis Tohama strain B213 was used. The Tohama strain has been shown to multiply in the lungs of mice (8, 16, 19). The bacteria were grown on Bordet-Gengou (BG) agar plates supplemented with 30 µg/ml streptomycin (Tritium, Veldhoven, The Netherlands) at 35°C for 3 days. Subsequently, the bacteria were plated on BG agar plates without antibiotics, cultured for 3 days, resuspended in Verwey medium (NVI, Bihofhoven, The Netherlands), and used for infection.

Infection of mice and autopsy. Intranasal infection was performed as described previously (47). Briefly, the mice were lightly anesthetized and a single drop of a 40-µl inoculum containing 2 × 10^8 B. pertussis cells was carefully placed on the top of the nose and allowed to be inhaled.

The mice were killed at 3, 5, or 7 days after infection. The animals were anesthetized with ketamine, xylazine (Rompun), and atropine, and blood was collected from the orbital plexus. Perfusion of the right ventricle was performed with 2 ml PBS supplemented with 3.5% heat-inactivated fetal calf serum (PAA, Linz, Austria). The lungs were excised and used either to obtain bronchial lymph nodes (LNs) and lungs for enumeration of the bacteria and for histological examinations or to obtain bronchoalveolar lavage fluid (BALF) cells.

Lung lobes, CFU determination, and histological examination. A ligature was made around the right bronchus, after which the right lobes were removed for the enumeration of the bacteria. The lobes were homogenized in 900 µl of Verwey medium by using a tissue homogenizer (Pro-200; ProScientific, Monroe, CT) at maximum speed for 10 s. The homogenates were diluted in Verwey medium 10- and 100-fold for the immunized mice and 1,000-fold for the control mice, and 100-µl aliquots of the dilutions were plated on BG agar plates supplemented with streptomycin and incubated at 35°C for 5 days. The remaining left lung lobes were fixed intratracheally by using 4% formalin for 24 h. After dehydrating overnight, they were embedded in paraffin. Five-micrometer sections were cut and stained with hematoxylin-eosin. Histological lesions were semiquantitatively scored as absent (0), minimal (1), slight (2), moderate (3), strong (4), or severe (5), respectively. This score incorporates the frequency as well as the severity of the lesions.

Ptx-specific IgG. Ptx-specific IgG was measured essentially as described previously for the analysis of human sera (14). Briefly, 96-well plates (Immunno Plate; Nunc, Roskilde, Denmark) were precoated with Ptx (NVI). Positive control sera was a pool of sera obtained from a previous experiment in which mice were vaccinated with the DTaP vaccine plus either aluminum or MPL and challenged with B. pertussis. The concentration of the positive control serum was arbitrarily set at 1,000 U. Dilution series of test sera and positive control sera were prepared in blocking buffer (0.5% bovine serum albumin [Sigma-Aldrich, Axxel, The Netherlands] and 0.01% Tween 20 [Meerk, Amsterdam, The Netherlands]) in PBS. The plates were incubated with 2,000-fold-diluted peroxidase-labeled rabbit anti-mouse IgG (Dako, Glostrup, Denmark) in blocking buffer for 1 h at 37°C and washed. Finally, the plates were incubated with substrate solution (10% sodium acetate, 1.66% tetramethylbenzidine [Sigma-Aldrich] and 0.02% H_2O_2) for 5 min and read at 450 nm.

Total serum IgE. Blood was allowed to clot at 4°C overnight and was centrifuged at 13,000 × g for 2 min. Total serum IgE was measured as described previously (44).

BALF cells. A cannula was placed intratracheally and was fixed with a suture. The lungs were placed in a 50-ml tube filled with PBS. One milliliter PBS was brought into the lung and sucked up. This was repeated twice. BALF cells were pelleted by centrifugation, resuspended in PBS, counted with a Coulter Counter (Coulter Electronics, Latun, United Kingdom), and visually differentiated after they were stained with Giemsa stain.

Cell culture. The cell culture medium used was RPMI 1640 (Gibco, Grand Island, NY) supplemented with 10% fetal calf serum, 100 µg/ml streptomycin, and 100 IU/ml penicillin. Cell suspensions were made by pressing the LNs or spleens through a cell strainer (Falcon, Franklin Lakes, NJ). The cells were counted with a Coulter Counter. LN cell suspensions were cultured at 10^6 cells per ml culture medium with 5 µg/ml concanavalin A (ConA; MP Biomedicals, Irvine, CA) in flat-bottom 12-well culture plates (Costar, Cambridge, MA) at 37°C in a humidified atmosphere containing 5% CO_2 for 24 h. Splen cell suspensions were cultured at 10^6 cells per ml culture medium with 5 µg/ml ConA or B. pertussis (10^8 heat-inactivated bacteria per well) in 96-well tissue culture plates (Nunc) at 37°C in a humidified atmosphere containing 5% CO_2 for 72 h. The bacteria were heat inactivated at 56°C for 30 min.

Cytokine measurements. The cytokine concentrations in the culture supernatants were measured by using a five-plex panel containing beads for mouse IL-4, IL-5, IL-10, IL-13, and gamma interferon (IFN-γ) or an eight-plex panel containing beads for mouse IL-2, IL-4, IL-5, IL-10, IL-12p70, granulocyte-macrophage colony-stimulating factor (GM-CSF), IFN-γ, and tumor necrosis factor alpha.
RESULTS

*B. pertussis* colonization of the lungs. To analyze whether supplementation of the DTaP vaccine with MPL improved the efficacy of the vaccine in comparison with that achieved with supplementation with aluminum, mice were vaccinated twice with 1/5, 1/25, or 1/125 HD DTaP vaccine supplemented with either MPL or aluminum as the adjuvant and were then challenged intranasally with *B. pertussis*. At 3 and 7 days after infection the lungs were excised and the number of viable *B. pertussis* organisms in the right lung lobes was determined. Each symbol represents the number of bacteria in the lung of an individual mouse; horizontal lines represent the group average. Nonboxed numbers show *P* values when the different vaccine doses were compared for the same adjuvant and day of killing. Boxed numbers show *P* values when the different adjuvants are compared for the same vaccine dose and day of killing. ANOVA was followed by *t* test. The results of a single representative experiment of three are shown.

**FIG. 1.** Colonization of the lungs by *B. pertussis*. Mice were subcutaneously injected with (H) 1/5, (M) 1/25, or (L) 1/125 HD DTaP vaccine plus aluminum (alum) or MPL or (C) with the adjuvants only twice before intranasal *B. pertussis* infection. At 3 and 7 days after infection the lungs were excised and the number of viable *B. pertussis* organisms in the right lung lobes was determined. Each symbol represents the number of bacteria in the lung of an individual mouse; horizontal lines represent the group average. Nonboxed numbers show *P* values when the different vaccine doses were compared for the same adjuvant and day of killing. Boxed numbers show *P* values when the different adjuvants are compared for the same vaccine dose and day of killing. ANOVA was followed by *t* test. The results of a single representative experiment of three are shown.
Only twice before intranasal DTaP vaccine plus aluminum (alum) or MPL or with the adjuvants only twice before intranasal B. pertussis infection, the mice were euthanized. Serum was taken, and serial dilutions of test and positive control sera were tested for Ptx-specific IgG. The data are indicated as the means ± standard errors of the means (n = 5). ANOVA was followed by t test. The results of a single representative experiment of two are shown.

**Evaluation of histological changes.** We have previously shown that mice vaccinated with DTaP vaccine with aluminum as the adjuvant and subsequently challenged with B. pertussis revealed increased lung pathology compared to that in mice that were treated only with the adjuvant before challenge (44). To address whether the adjuvant in the vaccine influenced lung pathology, the lung histology was scored 3 days after infection. For both types of adjuvant, the vaccinated animals showed increased perivasculitis (P = 0.009 compared to the results obtained with aluminum only and P = 0.015 compared to the results obtained with MPL only). The level of eosinophilia was lower in mice vaccinated with the DTaP vaccine with MPL as the adjuvant than in mice that received MPL only and than in mice that received the DTaP vaccine with aluminum as the adjuvant (Fig. 3). Vaccination resulted in minor increases in peribroncholitis, hypertrophy of the bronchiolar mucus cells, and alveolitis (data not shown). In conclusion, the MPL-supplemented vaccine induced lower levels of eosinophilia than the aluminum-supplemented one, which is suggestive of a lower type I hypersensitivity.

**Total serum IgE levels.** Another parameter of type I hypersensitivity is an increased serum IgE level. We have previously shown that mice vaccinated with the DTaP vaccine with aluminum as the adjuvant and subsequently challenged with B. pertussis displayed a large increase in serum IgE levels compared to the levels in mice that were treated only with adjuvant before challenge (44). To investigate whether supplementation of the DTaP vaccine with MPL instead of aluminum resulted in lower total serum IgE levels, serum was taken 3 days after infection and analyzed. The total IgE level was higher in both vaccinated groups, and no significant differences that depended on the type of adjuvant in the vaccine were seen (Fig. 4). Thus, the type of adjuvant used in the vaccine did not influence total serum IgE levels.

**Cytokine production by bronchial LN cells.** Next to lung eosinophilia, the increased production of the Th2 cytokines IL-4, IL-5, IL-10, and IL-13 was suggestive of a type I hypersensitivity response in previous vaccination experiments with the DTaP vaccine with aluminum as the adjuvant (44). As MPL can redirect Th2 to Th1 responses, we reasoned that the hypersensitivity response might be lower after vaccination with the MPL-supplemented vaccine. To examine this possibility further, bronchial LN cells were cultured in the presence of ConA for 24 h and the supernatants were analyzed for their cytokine contents by using an eight-plex assay that measured three Th2 cytokines (IL-4, IL-5, and IL-10), one Th1 cytokine (IFN-γ), and four additional cytokines involved in immune regulation (IL-2, IL-12p70, GM-CSF, and TNF-α). Bronchial LN cells from mice that were vaccinated with the DTaP vaccine with aluminum as the adjuvant showed higher levels of IL-4 production than those from mice that received aluminum only or mice that were vaccinated with the DTaP vaccine with MPL as the adjuvant (Fig. 5A). No treatment-related differences were seen for IL-2 and IFN-γ. The levels of production of IL-5, IL-10, IL-12p70, GM-CSF, and TNF-α by the bronchial LN cells from mice that received aluminum only were below the detection limit. Bronchial LN cells from mice that were vaccinated with DTaP vaccine in MPL showed higher...
levels of TNF-α production than those of mice that received MPL only ($P = 0.011$). No further differences were seen when the production of these cytokines by the bronchial LN cells of mice vaccinated with the DTaP vaccine in MPL was compared with those of mice vaccinated with the DTaP vaccine in aluminum or mice that received MPL only (data not shown). In conclusion, vaccination with the vaccine with MPL as the adjuvant resulted in lower levels of ex vivo ConA-induced IL-4 production by bronchial LN cells compared to those achieved after vaccination with the aluminum-supplemented vaccine, indicating that the immune response was skewed more toward a Th1-type response.

**Cytokine production by splenocytes.** To evaluate whether the adjuvant used in the vaccine affected ex vivo cytokine production by splenocytes, these cells from vaccinated and control mice were isolated and cultured in the presence of heat-killed *B. pertussis* for 72 h and the supernatants were analyzed for their cytokine contents. (B) IL-5 production by splenocytes stimulated ex vivo with heat-killed *B. pertussis*. The splenocytes were cultured with heat-killed *B. pertussis* for 72 h, and the supernatants were analyzed for their cytokine contents. The data are indicated as the means ± standard errors of the means ($n = 6$). ANOVA was followed by use of the Bonferroni correction. The results of a single representative experiment of two are shown.

**Vaccine reactogenicity.** While both LPS analogs, in comparison to aluminum, improve the efficacy of the DTaP vaccine and reduce type I hypersensitivity, they should not increase the reactogenicity of the DTaP vaccine. To address this issue, the concentration of the proinflammatory cytokine IL-6 was analyzed in serum samples taken 4 h after primary or booster immunization. A group of mice immunized with the DTaP vaccine plus aluminum showed a significantly higher number of eosinophils than all other groups (Fig. 7). In conclusion, the use of LpxL2 LPS and MPL as the adjuvants provided significantly better protection than the vaccine with aluminum as the adjuvant. Thus, with respect to efficacy, LpxL2, like MPL, compared favorably to aluminum as the adjuvant in the vaccine.

**BALF cells.** In order to determine whether the various adjuvants in the vaccine affected the cell-type distribution differently, lung lavage was performed 5 days after infection, and the BALF cells were counted and visually differentiated. The percentages and numbers of macrophages, neutrophils, and lymphocytes were similar in all groups; and the total number of BALF cells was also not differentially affected. However, the group immunized with the DTaP vaccine plus aluminum showed a significantly higher number of eosinophils than all other groups (Fig. 6). In conclusion, the use of LpxL2 LPS, like the use of MPL, instead of aluminum as the adjuvant in the DTaP vaccine resulted in a lower percentage and a lower number of eosinophils in the BALF, which is indicative of a lower type I hypersensitivity

**FIG. 5.** (A) IL-4 production by ex vivo ConA-stimulated bronchial LN cells. Mice were injected s.c. with 1/5 HD DTaP vaccine plus aluminum (alum) or MPL or with the adjuvants only twice before intranasal *B. pertussis* infection. Three days after infection, the bronchial LNs were excised, the cells were cultured with ConA for 24 h, and the supernatants were analyzed for their cytokine contents. (B) IL-5 production by splenocytes stimulated ex vivo with heat-killed *B. pertussis*. The splenocytes were cultured with heat-killed *B. pertussis* for 72 h, and the supernatants were analyzed for their cytokine contents. The data are indicated as the means ± standard errors of the means ($n = 6$). ANOVA was followed by use of the Bonferroni correction. The results of a single representative experiment of two are shown.
vaccine with LpxL2 LPS as the adjuvant ($P = 0.007$) but not with the DTaP vaccine with MPL as the adjuvant. In conclusion, the higher IL-6 levels obtained when LpxL2 LPS or MPL was used as the adjuvant in the DTaP vaccine may suggest some reactogenicity. The considerably lower IL-6 level induced by the DTaP vaccine than the level induced by the wP vaccine suggests that the DTaP vaccine supplemented with LPS analogs has a lower reactogenicity than the wP vaccine.

**DISCUSSION**

Here, we have shown that the replacement of aluminum by either one of two LPS analogs, MPL or LpxL2 LPS, as the adjuvant in the DTaP vaccine improves the vaccine in two ways; first, it enhances its efficacy, as shown by the reduced colonization of the lungs after challenge and (in the case of MPL) increased the Ptx-specific IgG titer; second, it skews the response more toward a Th1-type response, as indicated by the lower levels of Th2 cytokine production, resulting in a decrease in the parameters indicative of type I hypersensitivity, i.e., lung eosinophilia and eosinophil numbers. The higher IL-6 levels induced by these supplemented DTaP vaccines compared to

**FIG. 6.** Colonization of the lungs by *B. pertussis*. Mice were injected s.c. with PBS or 1/10 HD DTaP vaccine plus aluminum, LpxL2 LPS, or MPL twice before intranasal *B. pertussis* infection. At 5 days after infection the lungs were excised, and the number of viable *B. pertussis* organisms was determined. Each symbol represents the number of bacteria in the lung of an individual mouse; horizontal lines represent the group average. ANOVA was followed by $t$ test. The results of a single representative experiment of two are shown.

**FIG. 7.** BALF eosinophil numbers. Mice were injected s.c. with 1/10 HD DTaP vaccine plus aluminum, LpxL2 LPS, or MPL or with PBS twice before intranasal *B. pertussis* infection. At 5 days after infection, lung lavage was performed and the BALF cells were counted and visually differentiated. The data are indicated as the means ± standard errors of the means ($n = 6$). ANOVA was followed by use of the Bonferroni correction. The results of a single representative experiment of two are shown.

**FIG. 8.** Serum IL-6. Mice were injected s.c. with 1/10 HD wP vaccine; 1/10 HD DTaP vaccine plus aluminum, LpxL2 LPS, or MPL; or PBS. Serum IL-6 concentrations were determined 4 h postimmunization. The data are indicated as the means ± standard errors of the means ($n = 7$). ANOVA was followed by use of the Bonferroni correction. The results of a single representative experiment of two are shown.
those induced by the DTaP vaccine with aluminum as the adjuvant may suggest some reactogenicity, although the reactogenicity is well below that of the wP vaccine.

We have investigated whether MPL is able to enhance vaccine efficacy by performing a dose-response analysis. While vaccination with 1/5 HD DTaP vaccine with MPL as the adjuvant resulted in a significantly decreased colonization compared to that achieved with the DTaP vaccine with aluminum as the adjuvant, this was not observed with 1/25 and 1/125 HD, suggesting that MPL is more effective than aluminum only at a relatively high DTaP vaccine dosage. When LpxL2 LPS and MPL were compared as adjuvants, both LPS analogs seemed to improve the efficacy of the DTaP vaccine to similar extents when 1/10 HD DTaP vaccine was used.

There is controversy regarding the correlation between Ptx-specific IgG levels and protective immunity. Ptx-specific IgG levels have been shown to correlate with protection in both humans (11, 41, 42) and mice (7). Cell-mediated immunity, however, also critically contributes to protection in both humans (1, 9, 24) and mice (30). The latter notion may be an explanation of the various degrees of association between the IgG levels against B. pertussis antigens in serum and protective immunity. The association between Ptx-specific IgG levels and protective immunity is most apparent early after vaccination (10). We measured the Ptx-specific IgG titers 17 and 21 days after the second vaccination, that is, relatively early after vaccination, making it plausible that in our study Ptx-specific IgG levels indeed correlated with protection.

The use of MPL as the adjuvant resulted in higher Ptx-specific IgG levels than those achieved when aluminum was used, suggesting improved protection. Thus, both the improved clearance of B. pertussis and the higher Ptx-specific IgG levels suggest a more efficacious vaccine than when aluminum was used as the adjuvant. MPL induces much higher Ptx-specific IgG levels than wP vaccines. However, in the present study we compared the Ptx-specific IgG levels between aP-vaccinated mice only. It may be suggested that the Ptx-specific IgG levels that are induced by vaccination are affected by bacterial challenge. Prechallenge levels are, however, similar to the levels detected at 3 and 7 days postchallenge (R. M. Stenger and R. J. Vandebriel, unpublished observations).

We have previously shown that pertussis vaccination, especially with the DTaP vaccine, resulted in type I hypersensitivity. IL-4-knockout mice that showed a reduced hypersensitivity response showed an unaffected clearance, suggesting that the hypersensitivity is not beneficial and is possibly detrimental to the host (44). As MPL can redirect Th2 to Th1 responses (3, 34, 38, 48), we reasoned that the hypersensitivity response might be decreased by adding MPL to the DTaP vaccine. Indeed, the level of lung eosinophilia, lung eosinophil numbers, and the level of Th2 cytokine production were all decreased when MPL was added to the DTaP vaccine. The increase in the total IgE level in serum was, however, unaffected by the addition of MPL. Possibly, the immune-modulating capacity of MPL is too small to affect this response or the underlying mechanism(s) of this response is (partly) different from that for Th2 to Th1 redirection.

LpxL2 LPS harbors strongly reduced endotoxic activity while still exhibiting some adjuvant activity compared to that of wild-type N. meningitidis LPS (46). We speculated that it might also be effective in redirecting Th2 to Th1 responses. Consistently, lung eosinophil numbers were reduced when the DTaP vaccine was supplemented with LpxL2 LPS.

The levels of the proinflammatory cytokine IL-6 in serum samples taken 4 h after the primary and booster immunizations were higher when either of the LPS analogs was used as the adjuvant than when aluminum was used as the adjuvant. As expected, the IL-6 levels induced by the wP vaccine were significantly higher than those induced by the DTaP vaccine with any of the three adjuvants. Thus, the higher IL-6 levels induced by the DTaP vaccine supplemented with either LPS may suggest some reactogenicity, albeit a reactogenicity considerably lower than that of the wP vaccine. Unexpectedly, vaccine with aluminum as the adjuvant evoked significantly higher IL-6 levels during booster immunization than during primary immunization (P = 0.004). This suggests that booster immunization, also in the absence of strong immune stimulatory molecules such as LPS, may elicit a stronger IL-6 response than primary immunization. A similar effect was observed for vaccination with the DTaP vaccine with LpxL2 LPS as the adjuvant (P = 0.007) but not with the DTaP vaccine with MPL as the adjuvant. The latter finding may suggest that LpxL2 LPS and MPL differ in their mechanism(s) of action.

We have chosen to measure IL-6 as the parameter for reactogenicity, as this cytokine had the most sensitive response to several pyrogens in an in vitro system based on a human monocytic cell line and the ex vivo human whole-blood culture test system. The latter test represents the rabbit pyrogen test (31). While low or moderate IL-6 levels form an essential part of the immune response, excessive levels may be detrimental. A level that can be taken as a threshold for reactogenicity has not been established, however, and we therefore interpret increased serum IL-6 levels as being suggestive of reactogenicity.

Although the effects of MPL are believed to mimic the effects of LPS, albeit with a considerably lower toxicity, differences in cytokine induction between these two molecules have been reported, with MPL inducing IL-10 and IL-12 and LPS inducing only IL-12 (40). This finding may be explained by later studies that have shown that MPL engages both TLR2 and TLR4, whereas LPS acts only on TLR4 (26), with TLR2 agonists inducing IL-10 and TLR4 agonists inducing IL-12 (35, 36). The lack of induction of IL-1β and caspase 1 may also be an expression of the reduced toxicity of MPL compared to that of LPS (32). Knowledge of the mechanisms of action of LpxL2 regarding receptor specificity and downstream effects are currently lacking.

The present study has shown that supplementation of the DTaP vaccine with LPS analogs improves the efficacy of the vaccine and reduces type I hypersensitivity. Follow-up studies are, however, required. In such studies, the aP vaccine with aluminum as the adjuvant should be compared to the aP vaccine with LPS analogs as adjuvants in the total absence of aluminum, a reference B. pertussis strain should be used for challenge, and pre- and postchallenge T-cell responses against the individual vaccine components should be measured.

In conclusion, our results demonstrate that use of the DTaP vaccine with the LPS analogs MPL or LpxL2 LPS as the adjuvant improves vaccine efficacy and redirects the immune response from a Th2- to a Th1-type response, thereby reducing type I hypersensitivity.
REFERENCES


