The Mincle-Activating Adjuvant TDB Induces MyD88-Dependent Th1 and Th17 Responses through IL-1R Signaling

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Abstract

Successful vaccination against intracellular pathogens requires the generation of cellular immune responses. Trehalose-6,6-dibehenate (TDB), the synthetic analog of the mycobacterial cord factor trehalose-6,6-dimycolate (TDM), is a potent adjuvant inducing strong Th1 and Th17 immune responses. We previously identified the C-type lectin Mincle as receptor for these glycolipids that triggers the FcRγ-Syk-Card9 pathway for APC activation and adjuvanticity. Interestingly, in vivo data revealed that the adjuvant effect was not solely Mincle-dependent but also required MyD88. Therefore, we dissected which MyD88-dependent pathways are essential for successful immunization with a tuberculosis subunit vaccine. We show here that antigen-specific Th1/Th17 immune responses required IL-1 receptor-mediated signals independent of IL-18 and IL-33-signaling. ASC-deficient mice had impaired IL-17 but intact IFNγ responses, indicating partial independence of TDB adjuvanticity from inflammasome activation. Our data suggest that the glycolipid adjuvant TDB triggers Mincle-dependent IL-1 production to induce MyD88-dependent Th1/Th17 responses in vivo.

Introduction

Recombinant subunit vaccines are cheap and safe, but only weakly immunogenic unless adjuvants are used. The most commonly used human adjuvant Aluminium hydroxide (Alum) potently induces antibody responses but does not induce Th1 cellular immunity. Thus, new adjuvants are urgently needed to potentiate cell-mediated immune (CMI) responses crucial for protection against intracellular bacteria, e.g. Mycobacterium tuberculosis. A prerequisite for an efficient adjuvant is the activation of antigen presenting cells (APCs) by ligands for pattern recognition receptors. The choice of adjuvant(s) critically determines the type of memory response elicited, depending on the receptors and pathways triggered in APC via generation of cytokine milieu directing Th cell differentiation. E.g., the TLR9 ligand CpG ODN drives strong Th1 responses, whereas cationic dimethylidodecylammonium (DDA) liposomes containing the glycolipid trehalose-6,6-dibehenate (TDB), the synthetic analog of the mycobacterial cord factor trehalose-6,6-dimycolate, potently induce a strong Th17 in addition to Th1 immune response [1,2]. DDA/TDB (also known as CAF01) is a next generation adjuvant and has entered clinical studies for vaccination with the recombinant Mycobacterium tuberculosis fusion protein Ag85B-ESAT-6 (H1) [3,4]. We and others [5] identified the C-type lectin (CLR) Mincle as receptor for these glycolipids that triggers the FcRγ-Syk-Card9 pathway for APC activation and adjuvanticity [6]. In vitro, APC activation was solely dependent on recognition of TDB by Mincle, whilst MyD88 was dispensable. Surprisingly, TDB-immunized MyD88−/− mice failed to mount antigen-specific Th1 immune responses [7]. Since this was unexpected and contradictory to our in vivo results, we investigated in vivo requirements of known MyD88-utilizing signaling events in immunization experiments utilizing CAF01 and H1. Development of antigen-specific Th1 and Th17 immune responses was dependent on IL-1/IL-1R-mediated signals. Interestingly, inflammasome activation via ASC only partially accounted for CMI induction upon immunization with the glycolipid-containing adjuvant DDA/TDB.
Results

Mincle and MyD88 are Required for Induction of Th1 and Th17 Responses in vivo

We have previously shown that the cationic liposome formulation DDA/TDB induces not only a strong Th1 immune response but also a Th17 response [2]. Vaccination with DDA/TDB induces stable, long-lived multifunctional CD4 memory T cells [6,9]. We also found IFNγ and IL-17 secreted predominantly by CD4 T cells upon re-stimulation, while contribution of innate cells to IL-17 release was negligible (Fig. S1). This adjuvant effect depended on recognition of the synthetic glycolipid TDB by the CLR Mincle, when analyzed 7 days after a single subcutaneous (s.c.) immunization at the base of tail [6]. Here, we performed s.c. footpad prime-boost immunizations in order to utilize local swelling as an additional readout. Substantial swelling, peaking 7 days post immunization, was observed and significantly reduced in the absence of Mincle (Fig. 1A). This was in line with dramatically reduced secretion of IFNγ and IL-17 upon antigen-specific restimulation with H1 protein (Fig. 1B, C). Formation of H1-specific IgG2a antibodies, associated with Th1 responses, was also reduced whereas Th2 polarized IgG1 antibodies developed independently of Mincle (Fig. 1D, E). Footpad swelling was also significantly reduced in the absence of MyD88 (Fig. 1F), as well as IFNγ (Fig. 1G) and IL-17 (Fig. 1H) secretion. In addition, H1-specific IgG2a antibody generation was strongly impaired in the absence of MyD88, whereas the reduction of H1-specific IgG1 was not significant (Fig. 1I, J). Thus, in contrast to in vitro APC activation which solely required Mincle, in vivo adjuvanticity also strongly depends on MyD88 signaling.

Expression of Mincle by macrophages is inducible in vitro by TLR stimuli [10]. Therefore, it was possible that MyD88−/− mice have reduced Mincle levels due to a lack of responsiveness to TLR signals, derived e.g. from the gut flora, which could account for abrogated inflammatory and immune responses observed in MyD88−/− mice after immunization. To test this possibility, we first measured Mincle expression by qRT-PCR in FACS-sorted monocytes, neutrophils and T cells from naive C57BL/6 and MyD88-deficient mice (Fig. 2A). Expression was much higher in neutrophils than in monocytes, whereas T cells expressed very little Mincle mRNA. Of note, expression in monocytes and neutrophils was equally high in MyD88−/− as in control cells. Expression of Mincle was also determined at the site of injection (Fig. 2B). In naive C57BL/6 and MyD88−/− mice, Mincle mRNA was nearly undetectable but increased more than 3 orders of magnitude in both genotypes, most likely reflecting the infiltration by inflammatory leukocytes. Thus, absence of the adjuvant effect in MyD88−/− mice cannot be attributed to a lack of Mincle expression.

Since MyD88 is the common adaptor molecule in TLR-mediated signaling, we analyzed TLR2, 3, 4, 7 quadruple knockout of MyD88 also conveys signaling of the receptors for IL-1, IL-18 and IL-33 [11–14]. TDB induces the expression and secretion of IL-1β from macrophages in vitro [2]. We therefore analyzed IL-1 expression at the site of injection and detected increased mRNA levels of Il1b and to a lesser extent also Il1a in the feet 3 days post immunization (Fig. 3A). These changes at the mRNA level were at least in part due to leukocyte infiltration, as neutrophils constitutively express Il1a and Il1b (Fig. 3B). Importantly, cells isolated from footpads 7 days post immunization secreted IL-1β after over night incubation in culture medium (Fig. 3C). This was strongly dependent on Mincle and MyD88.

To assess the contribution of IL-1 receptor signals to adjuvanticity of TDB, we next investigated whether blockade of IL-1R1 with the soluble IL-1 receptor antagonist Anakinra influences the outcome of immunization. Anakinra treatment did not affect footpad swelling (Fig. 4A), but significantly reduced IFNγ and IL-17 secretion (Fig. 4B, C). Even though the serum concentration of Anakinra was consistently high in all mice treated (Fig. 4D), we do not know whether Anakinra...
treatment completely inhibited IL-1-mediated signaling. Therefore, we immunized IL-1R1−/− mice with DDA/TDB/H1. Due to differences in animal protection regulations in the different animal facilities, IL-1R1−/− mice were vaccinated s.c. at the tail base. Requirement for IL-1-mediated signals was more pronounced in these experiments as compared to short term Anakinra treatment. Cytokine secretion was significantly reduced (Fig. 4E, F); albeit not as strongly as seen in MyD88−/− mice. Interestingly, even though antibody generation was dependent on MyD88 signaling, absence of IL-1R1 did not alter IgG2a and only marginally affected IgG1 antibody formation (Fig. 4G, H). Thus, IL-1/IL-1R1 signaling contributes to the development of a Th1/Th17 immune response upon immunization with a CLR triggering adjuvant without strong effects on B cell responses or isotype switching.

IL-18 and IL-33 are Dispensable for Adjuvanticity

We next immunized IL-18−/− mice. Footpad swelling was significantly reduced (Fig. 5A), however IL-18 signaling was not required for Th1/Th17 response induction (Fig. 5B). Development of humoral immune responses remained also unchanged in the absence of IL-18 (Fig. 5C). Thus, contribution of IL-18/IL-18R signaling was rather negligible although it does mediate local inflammation at the injection site. Finally, in mice deficient in the IL-33 receptor component ST2, footpad swelling and secretion of IFNγ and IL-17 were not impaired after immunization (Fig. 5D, E).}

Canonical Inflammasome Activation Contributes to Th17 but not Th1 Responses

IL-1β and IL-18 have to be cleaved in order to exert their proinflammatory effects. This is mediated by caspase-1 in a process called inflammasome activation [15,16]. Since IL-1-mediated signaling contributing to DDA/TDB adjuvanticity, we asked whether inflammasome activation was also a requirement for the adjuvant effect. An essential adapter protein for inflammasome activation is ASC. To address the control of IL-1 production in response to the glycolipid adjuvant, we first stimulated bone marrow-derived DC from ASC−/− as well as Mincle−/− and MyD88−/− mice in vitro. Stimulation of DC with TDB induced expression of Il1a, Il1b and of Csf3 (encoding G-CSF) with complete dependence on Mincle, but independent of MyD88 and ASC (Fig. S3A). At the protein level, IL-1β was detected in the supernatant in a Mincle- and ASC-dependent, albeit MyD88-independent manner, whereas IL-1α protein was absent in Mincle−/− DC cultures, but independent of ASC (Fig. S3B). Of note, while Il1a and Il1b mRNA induction was triggered by immobilized TDB as well as by TDB in suspension, both IL-1 proteins were secreted only when DC were stimulated with the particulate TDB in suspension (Figure S3A, B), indicating that phagocytosis of glycolipids is a prerequisite for ASC-dependent release of IL-1β as well as the ASC-independent secretion of IL-1α.

ASC was shown to fulfill an important inflammasome-independent function in post-transcriptional regulation of cytoskeletal rearrangements and lack of ASC impaired antigen uptake and priming capacity of bone marrow-derived DC as well as chemotaxis of lymphocytes; all due to absent Dock2-signals [17]. Recently, the same group reported that not all of the different ASC−/− mouse lines available show defective Dock2 expression [18]; likely resulting in differences regarding the antigen presentation and migratory capacity of lymphoid and myeloid cells. We have used the ASC−/− mice originally generated at Genentech [19] and detected comparable Dock2 expression by qRT-PCR (data not shown). We tested ASC−/− bone marrow-derived macrophages in phagocytosis and spreading assays and found both unimpaired (Figure S4), arguing against an inflammasome-independent requirement for ASC in TDB-mediated adjuvanticity.

We next investigated the impact of ASC-deficiency on adjuvanticity in vivo (Fig. 6). Following footpad immunization of ASC−/− mice, local swelling and induction of IL-17 production was significantly reduced, whereas IFNγ induction was unaltered (Fig. 6A). In a set of experiments employing base of tail immunization, a significant effect of ASC-deficiency was again observed for IL-17 induction but not for IFNγ secretion (Fig. 6B). Antibody responses were not altered in ASC−/− mice. Thus, the lack of ASC has less severe consequences for induction of Th responses by DDA/TDB than the loss of IL-1R signaling, indicating that inflammasome activation contributes to but does not completely account for generation of IL-1-dependent signaling through MyD88 in vivo.

Discussion

Here we show that CMI induction upon immunization with the cationic adjuvant formulation DDA/TDB essentially requires MyD88 signaling. Our evidence for a TLR-independent role of MyD88 signaling in response to TDB stimulation is in line with data from Geisel et al. showing that cell recruitment and cytokine production induced by trehalose dimycolate was severely reduced in MyD88−/− mice but unaffected in TLR2−/− and TLR4−/−
mice [20]. The strong MyD88-dependence observed here contrasts to the results obtained by Gavin et al. claiming that MyD88 is dispensable for generation of T cell dependent antibody responses upon vaccination with a range of adjuvants [21]. This discrepancy may be due to differences in the antigen (whole protein vs. hapten antigen) or the readout used (Th cell polarization vs. antibody responses).

We identified here IL-1/IL-1R1 as the TLR-independent MyD88 pathway significantly contributing to induction of Th1 and Th17 cellular immune responses upon vaccination with DDA/TDB. IL-1 has been linked to induction of pathogenic Th17 cells in an EAE model [22] but was also shown to induce protective Th17 immune responses using *Escherichia coli* heat-labile enterotoxin as adjuvant [23]. Furthermore IL-1β and IL-18 can promote IL-17 and IFNγ production in CD4 and γδ T cells [24], in line with our data showing that both Th1 and Th17 induction is reduced in the absence of IL-1R1. Of note, CMI generation was reduced in IL-1R1−/− mice to a lesser extent than in MyD88−/− mice, raising the question which other factors explain the strong MyD88-dependence. One possible explanation could be synergistic effects of IL-1 and IL-18, which promotes Th1 T cell differentiation together with IL-12 [25]. Lalor et al. also suggest redundancy for IL-1β and IL-18 in Th17 induction [24]. We did not detect reduced CMI induction in IL-18−/− mice where IL-1/IL-1R1 is still functional; however, local inflammation in the footpad was reduced in IL-18−/− as in MyD88−/− mice. Although IL-1R1 signaling was required for robust IFNγ production by T cells, the Th1-associated IgG2a antibody response was unimpaired in IL-1R1−/− mice (Fig. 4). The basis for IL-1R1-independent, MyD88-dependent antigen-specific B cell responses is currently unknown; in future experiments, we plan to use conditional MyD88 transgenic mice to investigate the requirement for MyD88 in T and B lymphocytes, as well as myeloid cells, for efficient isotype switching.
MyD88 Is Required for Th1 Adjuvanticity In Vivo

A. Footpad swelling

B. Treatment

C. IL-17 secretion

D. Treatment

E. IFNγ secretion

F. IL-17 secretion

G. IgG1

H. IgG2a
In addition, the inducibility of Mincle by TLR ligands [10] raised the possibility that MyD88<sup>−/−</sup> mice may be less responsive to TDB as adjuvant because of reduced receptor expression. However, our results showing that Mincle expression is also increased in MyD88<sup>−/−</sup> mice upon immunization and that basal expression of Mincle in sorted monocytes, neutrophils and T cells is comparable between MyD88<sup>−/−</sup> and C57BL/6 control mice (Fig. 2) allow us to discard this possibility. Instead, impaired responses to other cytokines and chemokines, e.g. IFNγ [26] might contribute to the observed lack of responsiveness.

Unexpectedly, the adjuvant effect of DDA/TDB, even though dependent on IL-1/IL-1R1, was less strongly reliant on the inflammasome adaptor molecule ASC. Here again, DDA/TDB differed in the requirements for induction of immune responses compared to other known adjuvants, i.e. Alum which induces Th2 responses via Nlrp3-mediated inflammasome activation [27–29]. Schweneker et al. recently showed that TDB activates the Nlrp3 inflammasome in an ASC-dependent manner [30]. IL-1β secretion from bone marrow-derived DC upon stimulation with TDB was dependent on Caspase-1, Nlrp3 and ASC. It was further demonstrated that phagocytosis of TDB was a prerequisite for

![Figure 4. IL-1R1 signaling contributes to DDA/TDB adjuvanticity.](Image)

*Figure 4. IL-1R1 signaling contributes to DDA/TDB adjuvanticity.* Footpad swelling (A), IFNγ, IL-17 (B, C) secretion of cells isolated from the draining lymph nodes and Anakinra serum concentration (D) 7 days post immunization. Daily injection of 100 µg/g Anakinra or corresponding volume of PBS i.p.; first treatment 3–6 hours prior to immunization. Data presented as mean ± SD pooled from 5 independent experiments with 4–5 mice/group; except (D), IFNγ, IL-17 secretion of splenocytes (E, F) and H1-specific antibodies (G, H) in IL-1R1<sup>−/−</sup> and C57BL/6 controls; base of tail immunization. Data presented as mean ± SD pooled from 2 independent experiments with 5–6 mice/group.

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![Figure 5. IL-18 and IL-33 are dispensable for DDA/TDB adjuvanticity.](Image)

*Figure 5. IL-18 and IL-33 are dispensable for DDA/TDB adjuvanticity.* Footpad swelling (A), IFNγ, IL-17 secretion of cells isolated from the draining lymph nodes (B) and H1-specific antibodies (C) in IL-18<sup>−/−</sup> and C57BL/6 controls. Data presented as mean ± SD pooled from 3 independent experiments with 3–5 mice/group. Footpad swelling (D), IFNγ and IL-17 secretion of cells isolated from the draining lymph nodes in ST2<sup>−/−</sup> and C57BL/6 controls (E). Immunization at day 0 and day 21, mice sacrificed at day 35. Data presented as mean ± SD from 1 experiment with 4–5 mice/group.

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inflammasome activation. This is in line with our data (Fig. S3) showing also Mincle- and ASC-dependent IL-1β secretion only upon stimulation with TDB in suspension, whereas Il1b expression was increased independent of uptake and inflammasome activation. However, in the context of vaccination with DDA/TDB, dependency on ASC was less pronounced as compared to the loss of IL-1R signaling. One possible explanation could be that neutrophils utilize serine proteases to cleave pro-IL-1β [31]. Massive neutrophil influx is a hallmark of DDA/TDB adjuvanticity (data not shown) and we detected very high expression of Il1b in sorted neutrophils isolated from naive C57BL/6 and MyD88−/− mice (Fig. 3B). Alternatively, ASC-independent non-canonical inflammasome activation via Caspase-11 and subsequent IL-1α release [32] may be operating. IL-1α, in contrast to IL-1β, has entered clinical trials, further dissection of IL-1R-dependent effects on innate and adaptive immune cells will be relevant for a detailed understanding of the molecular mode of action of this adjuvant.

**Materials and Methods**

**Ethics Statement**

All procedures were discussed with and approved by the animal protection committees of regional Bavarian governments (Regierung von Mittelfranken or Oberbayern animal protocols number 54-2532.1.12/09 and 211-2531-33/05) according to German animal protection law (BGBl I S. 1206, 1313).

**Mice**

MyD88−/− mice were used with permission of Dr. S. Akira [11], Mincle−/− mice have been described [36]. IL-1R1−/− and ASC−/− mice were bred at the Technische Universität München. ASC−/− were originally generated at Genentech [19]. C57BL/6, Mincle−/−, MyD88−/−, ST2−/−, IL-18−/− and TLR9−/− mice were bred at the animal facility of the Medical Faculty in Erlangen. For some experiments, C57BL/6 mice were purchased from Harlan or Charles River.

**Immunization and IL-1 Receptor Blockade**

DDA/TDB liposomes and recombinant H1 were provided by the Statens Serum Institut. Adjuvant formulations were prepared as described [2]. Mice were immunized twice in a 14-day interval unless otherwise stated. IL-1 receptor signaling was blocked by intraperitoneal injection of 100 μg/g body weight soluble IL-1 receptor antagonist Anakinra (Kineret, Amgen) daily. Since

Taken together, we have shown here that the glycolipid adjuvant TDB relies on MyD88-dependent pathways for efficient Th1/Th17 adjuvanticity. Pharmacologic and genetic abrogation of IL-1R signaling identified IL-1 as the major MyD88-dependent factor induced by TDB through Mincle-Card9 signaling, providing new insight into the adjuvant mechanism. As DDA/TDB has entered clinical trials, further dissection of IL-1R-dependent effects on innate and adaptive immune cells will be relevant for a detailed understanding of the molecular mode of action of this adjuvant.
Anakirin had to be given daily and previous experiments showed IFNγ and IL-17 production could be detected 7 days after a single immunization [6], we chose a 7-day immunization protocol. Anakirin serum concentration was determined by sandwich ELISA using anti-human IL-1-RA antibodies (BioLegend).

Antigen-specific Restimulation and Detection of H1-specific Antibodies
Mice were sacrificed two weeks after the second immunization unless stated otherwise. Pooled draining inguinal and popliteal lymph nodes or spleens were meshed through a 100 μm nylon sieve and 5×10^6 cells were restimulated with 10 μg/ml H1 protein for 96 hours. Supernatants were analyzed for IFN secretion by ELISA (R&D Systems). Background (unstimulated cells) was subtracted and cytokine release expressed as fold change relative to the mean response of restimulated cells from immunized control mice. We chose to present summarized data as fold change in order to account for inter-experimental variation in overall cytokine production because experiments were conducted in different laboratories and with mice from different animal houses. Serum was analyzed in tenfold dilutions (starting from 1/1,000) for H1-specific antibodies with rabbit anti-mouse IgG1 and IgG2a (BD Biosciences). Pooled serum from immunized mice was included on each ELISA plate as a positive inter-assay control.

Intracellular Cytokine Staining
2×10^6 cells isolated from draining lymph nodes 7 days after the second immunization were re-stimulated with a mixture of 10 μg/ml H1 protein and 5 μg/ml of the peptides: Ag65B CD8 epitope (p1–19), FSRLPVLYLQVPSPSMG, Ag65B CD4 epitope (p241–255), QDAYANA-AGGHNAVFN, Ag65B CD4 epitope (p261–280), THSWEYWGALNNKMGDLQS and ESAT-6 (p1–15), MTEEQWNFAGIEAA. After 1 h 10 μg/ml Brefeldin A was added and incubation continued for 23 h. PMA (50 ng/ml) was used as a positive and culture medium as negative controls. Intracellular cytokine staining was performed according to standard protocols with antibodies against CD3, CD4, CD8, NK1.1, γδ TCR, CD11b, IFNγ and IL-17 (eBioscience or BioLegend) and data recorded on a FACS Canto II (BD Biosciences).

Generation and Stimulation of Bone Marrow-derived DC
Bone marrow cells were cultured on Petri dishes for 8 days in cRPMI containing 10% X63-cell conditioned medium. 2.5×10^5 cells/ml (cytokine secretion) or 5×10^6 cells/ml (qRT-PCR) were stimulated as indicated with plate coated TDB [2] or TDB in suspension (TDB suspended in DMEM by vortexing, heating to stimulated as indicated with plate coated TDB [2] or TDB in suspension). Supernatants were analyzed for IFN secretion by ELISA using anti-human IL-1-RA antibodies (BioLegend). Anakinra serum concentration was determined by sandwich ELISA (R&D Systems). Background (unstimulated cells) was subtracted and cytokine release expressed as fold change in order to account for inter-experimental variation in overall cytokine production because experiments were conducted in different laboratories and with mice from different animal houses. Serum was analyzed in tenfold dilutions (starting from 1/1,000) for H1-specific antibodies with rabbit anti-mouse IgG1 and IgG2a (BD Biosciences). Pooled serum from immunized mice was included on each ELISA plate as a positive inter-assay control.

Single Cell Isolation from Footpad
The injection site was excised using a scalpel and snap frozen in liquid nitrogen for qRT-PCR analysis. RNA was isolated using TriFast (peqLab) according to the manufacturer’s protocol. For IL-1β release single cell suspensions were obtained using gentleMACS (Miltenyi Biotec) according to the manufacturer’s protocol and 5×10^6 cells cultured in 200 μl cDMEM over night.

Quantitative RT-PCR
Expression levels of the housekeeping gene Hprt as well as Il1α, Il1β, Csf3 and Mincle were analyzed using primer/probe combinations selected from the Roche Universal Probe Library. Fold changes were calculated with the 2-ΔΔCT method using calibrators as indicated.

Cell Sorting
Single cell suspensions from bone marrow and spleen were stained with antibodies against CD11b, Ly6C, Ly6G (bone marrow) or CD3, CD19 (spleen); all antibodies from eBioscience. Monocytes (CD11b+Ly6C)Ly6G−, neutrophils (CD11b+Ly6C)T cell (CD3+CD19−) were sorted on a MoFlo (Beckman Coulter). 5×10^5-1×10^6 sorted cells were lysed in TriFast for RNA isolation and qRT-PCR.

Phagocytosis and Spreading Assay
Spreading assay of bone marrow-derived macrophages (BMM) was performed as described [37]. For phagocytosis assays BMM were incubated with green fluorescent latex beads (1 μm; Flouresbrite Microparticles, Polyscience, Inc.) with an MOI of 20 and incubated for 1 h or 4 h at 37°C. Cells were washed twice with ice cold PBS, detached from the 24 well plate by using a cell scraper and resuspended in PBS containing 0.2% BSA. Percentages of living cells which had phagocytosed beads were determined using a FACSCanto II (BD Biosciences).

Statistical Analysis
Statistical analysis was performed using Prism 5 from GraphPad Software, Inc. Significance was determined by 2-way ANOVA with Bonferroni correction for footpad swelling and H1-specific antibodies. For cytokine secretion normal distribution was tested by Shapiro-Wilk followed by unpaired student’s t-test for Gaussian or Mann-Whitney for non-Gaussian distribution. *p<0.05, **p<0.01 and ***p<0.001.

Supporting Information
Figure S1 IFNγ and IL-17 are secreted by bona fide Th1 and Th17 cells. Number of IFNγ (A) and IL-17 (B) secreting cells determined by intracellular cytokine staining. Cells isolated from draining popliteal lymph nodes 7 days after the second immunization (footpad). 2×10^6 cells per well, 24 h incubation with 10 μg/ml H1, PMA/Ionomycin or medium control in the presence of Brefeldin A. IFNγ and IL-17 (eBioscience or BioLegend) and data recorded on a FACSCanto II (BD Biosciences).

Figure S2 TLR2,3,4,7 and 9 seem dispensable for DDA/TDB adjuvanticity In Vivo. Footpad swelling, IFNγ and IL-17 secretion in TLR2,3,4,7−/− and C57BL/6 controls (A). Data presented as mean ± SD from 1 experiment with 3 mice/group. Footpad swelling, IFNγ and IL-17 secretion in TLR9−/− and C57BL/6 controls (B). Data presented as mean ± SD from 3 independent experiments with 3–5 mice/group. Cytokine production of cells isolated from the draining lymph nodes. (TIF)

Figure S3 TDB-induced expression and release of IL-1α and IL-1β. Expression of Il1α, Il1β and Csf3 (A), and IL-1α and IL-1β secretion (B). Bone marrow-derived DC were stimulated for 24 h with plate-coated TDB (solvent control isopropanol) or TDB in suspension. 2.5×10^5 cells/well were seeded for cytokine release (TDB concentration as indicated) and 5×10^6 cells/well for qRT-PCR (5 μg/ml TDB). Fold change calculated against DC from C57BL/6 mice in medium. One experiment performed in duplicates. (TIF)
Figure S4 ASC−/− macrophages show no defects in spreading kinetics and phagocytosis capacity. Spreading kinetics (A) of C57BL/6 (circles) and ASC−/− (squares) BMM stimulated with LPS (closed symbols) or media control (open symbols). Mean ± SEM of at least 300 cells per condition. Statistical significance refers to the comparison of LPS stimulated C57BL/6 and ASC−/− BMM. One experiment performed. Phagocytosis capacity of C57BL/6 (B) and ASC−/− (C) BMM. Cells were incubated with fluorescent latex beads (1 μm; MOI 20) for 1 h and 4 h. Percentages of cells which phagocytosed beads determined flow cytometry. One of two representative experiments shown. (TIF)

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Author Contributions

Conceived and designed the experiments: CD KW RL. Performed the experiments: CD KW MR KJ NR JW. Analyzed the data: CD KW JW. Contributed reagents/materials/analysis tools: US DG SW CK EMA CPGC. Wrote the paper: CD MR CPdC RL.

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