Signaling Networks Responsible for the Dynamic Modulation of IL-12 in Response to Varying Dosages of LPS in Macrophages

Abstract

The innate immune response to lipopolysaccharide (LPS) has many possible outcomes depending on the dose, from acute endotoxic shock to broad immunomodulation. High doses (>10 ng/mL) evoke both pro- and anti-inflammatory mediators, while super-low doses (<1 ng/mL) result in preferential induction of pro-inflammatory genes. The mechanisms governing the switch between a preferentially pro-inflammatory or a balanced, resolving response are poorly understood. We show that in murine macrophages, the pro-inflammatory cytokine interleukin 12 (IL-12) is induced most robustly by intermediate LPS doses (1-10 ng/mL), while higher doses evoke a diminishing response. In the absence of the suppressive Toll-like receptor 4 signaling cascade members Lyn and IRAK-M, macrophages sustain robust IL-12 production in response to high doses of LPS. The transcription factor cAMP response element-binding protein (CREB) appears to play an important role in the regulation of the inflammatory response to LPS, with Lyn and IRAK-M deficient macrophages failing to activate CREB effectively in response to LPS. On the other hand, co-stimulation of these cells with LPS and the CREB agonist adenosine suppresses pro-inflammatory responses to LPS in wild-type cells, but does not robustly suppress IL-12 production in either Lyn- or IRAK-M-deficient cells. Our study reveals potential mechanisms for the dynamic modulation of innate responses by varying dosages of LPS.

Introduction

Inflammation is the first step in the natural response of the body to infection or injury. It broadly consists of an initial phase, in which damaged cells, pathogens, and debris are destroyed and cleared away, and a resolving phase, in which anti-inflammatory mediators and cell types allow for the regeneration and restoration of damaged tissue [1]. Unresolved inflammation may become persistent, and is implicated in a wide range of chronic conditions including atherosclerosis and diabetes [2-4].

Low-grade inflammation is more likely to become persistent, since the inflammatory signals involved tend not to cross the thresholds necessary to trigger resolution. We have shown that very low doses (<1 ng/mL) of lipopolysaccharide (LPS), which signals through the Toll-like receptor 4 (TLR4), can induce low-grade inflammation [5]. The signaling pathways downstream of TLR4 are complex, with different adaptor proteins triggering divergent responses [6,7]. Prominently, cell-surface TLR4 preferentially activates pro-inflammatory signaling through MyD88 and NFkB, while CD14-dependent TLR4 endocytosis leads to signaling through TRIF, resulting in the activation of phosphoinositide-3-kinase (PI3K) and anti-inflammatory compensating mechanisms [8,9]. Most existing studies of the inflammatory response to LPS have been conducted with doses between 100-1000 ng/mL, leaving the specific response to stimulation by LPS at doses lower than 1 ng/mL largely unexplored.

PI3K appears to be an important means of resolving TLR-induced inflammation, playing a predominantly anti-inflammatory role in the LPS response [10,11]. Interference with signaling molecules upstream of PI3K such as Lyn or B-cell adaptor for PI3K (BCAP) inhibits PI3K activation upon challenge with inflammatory stimuli including LPS, leading to anti-inflammatory skewing of the immune reaction due to a lack of modulatory influence [12].

The Src-family kinase Lyn is implicated in TLR signaling and B cell function, but its precise role is unclear. Lyn-/- dendritic cells (DC) have heightened sensitivity to LPS, showing increased NFkB and MAPK activity upon LPS stimulation [13], and DC-specific knockout of Lyn leads to B cell hyperactivity. B cells lacking Lyn show reduced Akt activity in response to CD19 stimulation [12]. HEK293 cells transfected with TLR4 and MD-2 display decreased association between Lyn and TLR4 after tolerization with LPS [14], and Lyn knockdown in THP-1 cells increases TRIF-dependent signaling in response to cytomegalovirus infection [15]. The autoimmunity associated with Lyn deficiency can be reversed by additional knockout of MyD88 [16,17], suggesting that Lyn is important for the suppression of MyD88-dependent TLR4 signaling, and that it acts by engaging the PI3K/Akt pathway in response to LPS.

The interleukin-receptor associated kinase M (IRAK-M) is a well-known suppressive modulator of TLR4 signaling, dampening signal transduction through the MyD88-dependent pathway [18], although the precise mechanisms of its activity are unclear. It suppresses p38 MAPK activation by stabilizing MKP-1, and IRAK-M-/- cells show altered distribution of the suppressive NFkB family member RelB upon TLR stimulation [19]. As such, it represents a promising target for study of the dynamics of the inflammatory response to LPS.

In this study, we tested the effects of varying LPS doses on the expression of IL-12 from primary murine macrophages harvested from wild type, IRAK-M deficient and Lyn deficient mice. We demonstrated differential expression profiles of IL-12 in macrophages challenged with varying LPS doses. Deletion of either Lyn or IRAK-M skew macrophages toward heightened expression of IL-12, but
addition of adenosine can nudge the macrophages toward reduced expression of IL-12. The dynamic balance of IL-12 expression may be controlled by a molecular circuit that requires Lyn- and IRAK-M.

**Materials and Methods**

**Cell culture**

Wild type (WT) mice were purchased from the Charles River laboratory. IRAK-M-deficient mice were initially supplied by Dr. Richard Flavell of Yale. All animals were housed and bred in the animal care facility of Life Sciences I in accordance with protocols approved by the Virginia Tech IACUC. Lyn-deficient mouse bones were generously provided by the laboratory of Dr. Clifford Lowell from the University of California, San Francisco. Murine BMDM were cultured as described previously [5].

**Western blots**

BMDM were cultured overnight in DMEM containing 1% fetal bovine serum, 1% penicillin/streptomycin, and 1% L-glutamine before performing experiments. Whole-cell lysates were harvested using lysis buffer consisting of 2% SDS, 5% Tris-HCl pH 6.8, and 10% glycerol, placed on ice for twenty minutes, boiled for five minutes, then centrifuged at 12,000 RPM for four minutes at room temperature for removal of intracellular debris. Protein concentration was assessed by Bradford assay. Nuclear lysates were obtained using a lysis buffer consisting of 2% SDS, 5% Tris-HCl pH 6.8, and 10% glycerol, placed on ice for twenty minutes, boiled for five minutes, then centrifuged at 12,000 RPM for four minutes at room temperature for removal of intracellular debris. Protein concentration was assessed by Bradford assay. Nuclear lysates were obtained using a lysis buffer containing 10 mM HEPES, 1.5 mM MgCl₂, 10 mM KCl, 0.5 mM DTT, and 0.05% NP40. Lysates were placed on ice for ten minutes, and then centrifuged at 3,000 RPM for ten minutes at 4 °C. The resulting pellet was then resuspended in 5 mM HEPES, 1.5 mM MgCl₂, 0.2 mM EDTA, 0.5 mM DTT, 26% glycerol by volume, and 300 mM NaCl and vortexed before being placed on ice for thirty minutes, then centrifuged at 12,000 RPM for twenty minutes at 4 °C. The supernatant was recovered and concentration was determined by Bradford assay. Protein samples were run on 10% acrylamide gels at 100 V, followed by transfer at 110 V before blocking in 5% milk in TBS-T, and CREB (Cell Signaling 9197S) and against pCREB-S133 (Cell Signaling 9191) was used at a concentration by transfer at 110 V before blocking in 5% milk in TBS-T. Antibody samples were run on 10% acrylamide gels at 100 V, followed by transfer at 110 V before blocking in 5% milk in TBS-T, and CREB (Cell Signaling 9197S) and GAPDH (Santa Cruz sc-25778) at 1:1000. Densitometric analysis was performed using Fujifilm MultiGauge v. 3.0.

**Flow cytometry**

Cells were starved overnight as described above, and then stimulated with different concentrations of LPS. After two hours, 3 μg/mL of brefeldin A (eBioscience 00-4506) was added, followed by 1 μg/mL ionomycin (Sigma 10634-1MG), 20 ng/mL phorbol 12-myristate 13-acetate (PMA) (Sigma P8139-1MG), and additional brefeldin eighteen hours later. At twenty-four hours, cells were harvested and processed using a fixation and permeabilization kit (BD Biosciences 554715). Flow cytometry was performed using antibodies against IL-12 (BD Biosciences 554480) and analyzed on BD FacsDiva software.

**Statistics**

Statistical analysis by Holm-Sidak pairwise comparison or Student’s t test was performed using Sigma Plot 11 software (SigmaPlot) as detailed in the figure legends. Results were considered to be statistically significant at p < 0.05.

**Results**

**Dynamic modulation of IL-12 expression by increasing doses of LPS**

IL-12 is a classic pro-inflammatory cytokine induced by stimulation of macrophages through TLR4 [8]. Thus, we chose to investigate the effects of different LPS doses on the expression of IL-12 by murine BMDM. As shown in Figure 1, rising doses of LPS (from 100 pg/mL to 10 ng/mL) induced elevated expression of IL-12 in wild type bone marrow-derived macrophages (BMDM) as shown by flow cytometry. Intriguingly, as the LPS concentration rose above 10 ng/mL, the expression of IL-12 failed to increase further and was reduced instead.

**The biphasic expression pattern of IL-12 depends on the presence of negative regulators Lyn or IRAK-M**

The biphasic responses of IL-12 expression to low and high doses LPS may be due to competing cellular circuits in macrophages. Previous studies suggest that both pro- and anti-inflammatory processes may be induced by the Toll-like-receptor pathways [20]. We hypothesized that higher doses of LPS may be more efficient in engaging negative regulators of inflammation, and thus causing

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**Figure 1:** Effects of different LPS doses on IL-12 production by WT BMDM. Cells were stimulated for 24h with the indicated concentration of LPS, with addition of brefeldin at 2h and PMA and ionomycin at 20h, at the conclusion of which cytokine production was analyzed by flowcytometry. A) Representative MFI plots. B) IL-12 production by WT BMDM peaks at 10 ng/mL LPS, while higher doses induce less IL-12. Letters denote statistically significant differences between groups (* as compared to untreated control, p< 0.05, Holm-Sidak).
compensatory reduction of IL-12 expression. To test this approach, we examined the involvement of two key negative regulators Lyn and IRAK-M. IRAK-M is a known negative regulator of TLR signaling [18], and Lyn has been implied to be a negative regulator proximal to TLR complex [13,14].

As shown in Figure 2A, the biphasic response pattern of IL-12 induction by LPS was not present in Lyn deficient macrophages. Instead, IL-12 expression levels steadily rose in Lyn deficient cells with increasing amount of LPS. Likewise, the expression levels of IL-12 were not ablated with rising LPS concentrations (Figure 2B).

CREB activity corresponds with reduced IL-12 production in BMDM

The potential molecular mechanisms for the compensatory suppression of inflammatory mediators may include the transcriptional modulator CREB, which promotes the production of IL-10 [21] and has also been linked to the suppression of IL-12 [22]. CREB is activated by phosphorylation at Serine 133, which results in increased transcription of anti-inflammatory genes and skew macrophages towards an alternatively-activated phenotype [22-24]. In particular, Lyn has been shown to play a role in CREB activation through ERK in the context of brain ischemia [25]. In light of this, we hypothesized that higher doses LPS may induce CREB phosphorylation through Lyn or IRAK-M, and investigated the phosphorylation status of CREB in cells treated with varying doses LPS. As shown in Figure 3A, only higher doses of LPS induced robust phosphorylation of CREB. In contrast, the levels of CREB phosphorylation were reduced in Lyn deficient and IRAK-M deficient macrophages, particularly with respect to nuclear translocation of activated CREB (Figure 3B).

Suppression of pro-inflammatory phenotype by adenosine

To further test that CREB phosphorylation may be associated with the compensatory inhibition of IL-12 expression, we treated cells with adenosine, an established CREB agonist with anti-inflammatory effects in neutrophils [26,27]. We observed that co-stimulation of BMDM with 500 nM adenosine and varying dosages of LPS suppressed the expression of IL-12 in wild type BMDM (Figure 4A). As expected, adenosine failed to exert its inhibitory effect in IL-12 expression in either Lyn or IRAK-M deficient cells (Figure 4B, Figure 4C).

Discussion

The patterns of IL-12 production in response to LPS stimulation of macrophages show complex dynamics. The phenomena of LPS priming and tolerance are defined as different responses to secondary LPS challenge after a primary stimulation [11], but it is clear that even the primary response can vary widely depending on the LPS dosage. In a typical dose-dependent response, production of a cytokine or activity of a gene increases or decreases in a manner correlating with the concentration of the stimulant. However, the IL-12 response to LPS is more complex. Rather than a simple increase or plateauing of IL-12 production, the patterns of the LPS response in WT cells peak at an intermediate dose, while further increase in the stimulatory

Figure 2: Effects of different LPS doses on IL-12 production by Lyn^-/^- and IRAK-M^-/- BMDM. Cells were stimulated for 24h with the indicated concentration of LPS, with addition of brefeldin at 2h and PMA and ionomycin at 20h, at the conclusion of which cytokine production was analyzed by flow cytometry. Upper panel comprises representative dot plots, lower panel analyzes production of IL-12 by A) Lyn^-/- and B) IRAK-M^-/- BMDM in response to varying dosages of LPS. Letters denote statistically significant differences between groups (" as compared to untreated control, p< 0.05, Holm-Sidak).
dose actually results in decreased IL-12 production. Removal of Lyn or IRAK-M changes the dynamics of this response, resulting in undiminished or continually increasing production of IL-12 as the stimulatory dose of LPS increases.

The dynamic modulation of IL-12 is critically important for host physiology during microbial infection. Robust induction of IL-12 is essential to mount an effective Th1 type immune defense [28,29]. On the other hand, down-regulation of IL-12 may serve as a compensatory mechanism to avoid excessive inflammation. Indeed, as shown in this report and other previous studies, immune macrophages dampen their IL-12 expression when challenged with higher dosages of endotoxin, as dysregulation of IL-12 suppression increases mortality of mice in the context of endotoxic shock [30].

Mechanistically, our study reveals that high-dose LPS may engage negative regulators such as IRAK-M and Lyn in the compensatory suppression of IL-12. This is consistent with previous findings that IRAK-M and Lyn may serve as negative regulators for the TLR signaling circuit [13,19,31,32]. Our work further extends this concept and reveals a differential engagement of these negative regulators by varying dosages of LPS.

The importance of CREB in the modulation of IL-12 is further buttressed by the observation that treatment of BMDM with adenosine consistently suppresses IL-12 expression in response to LPS, especially at higher LPS doses. In WT cells, co-stimulation with LPS and adenosine resulted in clear suppression of IL-12, and promoted the increased activation of CREB. Lyn- and IRAK-M-
References


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