

NF κ B Is Persistently Activated in Continuously Stimulated Human Neutrophils

Veronika Miskolci,^{1,2} Janet Rollins,¹ Hai Yen Vu,¹ Chandra C Ghosh,¹ Dennis Davidson,² and Ivana Vancurova^{1,2}

¹Department of Biological Sciences, St. John's University, New York; ²Division of Neonatal-Perinatal Medicine, and Center for Immunology and Inflammation, The Feinstein Institute for Medical Research, North Shore – Long Island Jewish Health System, New Hyde Park, NY, USA

Increased activation of the transcription factor NF κ B in the neutrophils has been associated with the pathogenesis of sepsis, acute lung injury (ALI), bronchopulmonary dysplasia (BPD), and other neutrophil-mediated inflammatory disorders. Despite recent progress in analyzing early NF κ B activation in human neutrophils, activation of NF κ B in persistently stimulated neutrophils has not been previously studied. Because it is the persistent NF κ B activation that is thought to be involved in the host response to sepsis and the pathogenesis of ALI and BPD, we hypothesized that continuously stimulated human neutrophils may exhibit a late phase of NF κ B activity. The goal of this study was to analyze the NF κ B activation and expression of I κ B and NF κ B proteins during neutrophil stimulation with inflammatory signals for prolonged times. We demonstrate that neutrophil stimulation with lipopolysaccharide (LPS) and tumor necrosis factor- α (TNF α) induces, in addition to the early activation at 30–60 min, a previously unrecognized late phase of NF κ B activation. In LPS-stimulated neutrophils, this NF κ B activity typically had a biphasic character, whereas TNF α -stimulated neutrophils exhibited a continuous NF κ B activity peaking around 9 h after stimulation. In contrast to the early NF κ B activation that inversely correlates to the nuclear levels of I κ B α , however, in continuously stimulated neutrophils, NF κ B is persistently activated despite considerable levels of I κ B α present in the nucleus. Our data suggest that NF κ B is persistently activated in human neutrophils during neutrophil-mediated inflammatory disorders, and this persistent NF κ B activity may represent one of the underlying mechanisms for the continuous production of proinflammatory mediators.

Online address: <http://www.molmed.org>

doi: 10.2119/2006-00072.Miskolci

INTRODUCTION

Neutrophils (polymorphonuclear leukocytes, or PMN) are a crucial part of the innate immune system and play a vital role in the inflammatory response that characterizes sepsis, acute lung injury (ALI), and bronchopulmonary dysplasia (BPD) (1-4). In addition to their phagocytic and killing properties, neutrophils synthesize numerous proinflammatory cytokines and chemokines that may amplify the inflammatory process (5-9). Furthermore, recent studies have shown that neutrophil apoptosis is delayed in patients with sepsis, ALI, and

BPD (10-12). Expression of many of these proinflammatory and antiapoptotic genes is regulated at the level of transcription by the transcription factor NF κ B (13-22).

In most resting cells other than neutrophils, NF κ B is present in the cytoplasm bound to the inhibitory protein I κ B α (13,16,23). Cell stimulation with inflammatory signals leads to phosphorylation of the cytoplasmic I κ B α , followed by its rapid ubiquitination and degradation by the proteasome. This process releases the NF κ B proteins from the inhibitory complex, and they then

translocate to the nucleus and bind to the NF κ B-responsive promoters (13-17). Thus, in this classic model of NF κ B regulation, NF κ B activity is regulated by the cytoplasmic degradation of I κ B α , and by the nuclear translocation of NF κ B subunits.

Despite remarkable progress in understanding the NF κ B regulation in other human cells as well as in animal models (13-17), much remains unknown about the mechanisms regulating NF κ B activity in human neutrophils. Previous studies from our laboratory have demonstrated that human neutrophils differ from monocytic and other cells, in that they contain predominant amounts of I κ B α in the nucleus of resting cells (24). Our results have shown that neutrophil stimulation with inflammatory signals such as LPS and TNF α result in the degradation of both cytoplasmic and nuclear I κ B α ,

Address correspondence and reprint requests to Ivana Vancurova, Department of Biological Sciences, St. John's University, 8000 Utopia Parkway, New York, NY 11439. Phone: 718-990-6409; Fax: 718-990-5958; E-mail: vancuroi@stjohns.edu

Submitted September 6, 2006; Accepted for publication January 22, 2007.

and that the NF κ B activity induced by LPS and TNF α stimulation for 30–60 min inversely correlates with the nuclear levels of I κ B α (25,26). In addition, we have shown that the increased nuclear level of I κ B α in human neutrophils is associated with the inhibition of NF κ B activity and increased neutrophil apoptosis (25). A recent study has provided evidence for the nuclear I κ B α degradation in neutrophils by the constitutively expressed nuclear I κ B kinase complex (27). Thus, nuclear I κ B α is crucial for the regulation of NF κ B activity and neutrophil apoptosis (24–26). However, the exact mechanisms by which nuclear I κ B α regulates the NF κ B-dependent transcription remain unknown.

Increased activation of NF κ B in neutrophils has been associated with the pathogenesis of ALL, BPD, sepsis, and other inflammatory diseases (28–33). Whereas the acute NF κ B activation induced by neutrophil stimulation for 30–60 min has been well documented (34–41), NF κ B activation during the prolonged neutrophil stimulation that is likely to occur in inflammatory disorders has not been previously studied. In this study, we describe a previously unrecognized persistent NF κ B activation in human neutrophils stimulated with TNF α and LPS for up to 12 h. Whereas the early activation of NF κ B is regulated by the nuclear levels of I κ B α , the newly synthesized nuclear I κ B α induced by continuous neutrophil stimulation appears to be no longer sufficient to inhibit the persistent NF κ B activity. These data suggest that the NF κ B activity persistently increased during neutrophil-mediated inflammatory disorders is regulated by a new, I κ B α -independent mechanism.

MATERIALS AND METHODS

Materials

Ficoll-Paque PLUS, dextran T-500, T4 polynucleotide kinase, poly(dI-dC), and Sephadex G25 spin columns were purchased from Pharmacia (Piscataway, NJ, USA). Hanks balanced salt solution, RPMI 1640, and endotoxin-tested, heat-

inactivated fetal calf serum were obtained from Life Technologies (Grand Island, NY, USA). *E. coli*-expressed purified recombinant human TNF α was purchased from R&D Systems (Minneapolis, MN, USA). [³²P] γ -ATP was purchased from Perkin Elmer (Boston, MA, USA). Purified polyclonal antibodies against human I κ B α (sc-371), I κ B β (C-20; sc-945), I κ B β (N-20; sc-969), I κ B ϵ (sc-7156), p50 NF κ B (sc-7178), c-Rel NF κ B (sc-70X), histone H2A (sc-10807), lamin B (sc-6216), p21^{CIP1} (sc-469), and I κ B α -agarose conjugate (sc-203AC) were purchased from Santa Cruz Biotechnology (Santa Cruz, CA, USA). Purified polyclonal antibody against p65 NF κ B (SA-171) was obtained from Biomol (Plymouth Meeting, PA, USA), and lactate dehydrogenase (LDH) antibody (20-LG22) from Fitzgerald Industries International (Concord, MA, USA). Horseradish peroxidase (HRP)-conjugated anti-rabbit and anti-goat secondary antibodies were from Amersham (Arlington Heights, IL, USA). All other reagents were molecular biology grade and were purchased from Sigma (St Louis, MO, USA). All reagents and plasticware used throughout the experiments were pyrogen-free.

Neutrophil Isolation and Culture

Fresh blood was obtained from healthy adult human volunteers and collected in heparinized preservative-free tubes as described (24,36). The study was approved by the Institutional Review Board of the North Shore–Long Island Jewish Health System, and informed written consent was obtained from all subjects. Neutrophils were purified under endotoxin-free conditions using Ficoll-Paque centrifugation, followed by dextran sedimentation and hypotonic lysis of residual erythrocytes as described (24,36). This neutrophil isolation procedure takes approximately 3 h and yields neutrophils that are 95% to 98% pure and 98% to 99% viable (24,36). Purified neutrophils were resuspended in RPMI 1640 supplemented with 10% low-endotoxin fetal calf serum at a final concentration of 5 \times

10⁶ cells/mL and incubated in 48-well cell culture plates at 37°C in a 5% CO₂ humidified atmosphere.

Electrophoretic Mobility Shift Assay (EMSA)

The nuclear extracts were prepared, and EMSA assays were performed as described (24,42).

Western Blot Analysis

Cytoplasmic extracts (CE) and nuclear extracts (NE) were prepared as described (24–26). Denatured proteins were separated on 12% denaturing polyacrylamide gels, and immunoblotting analysis was performed as described (24–26). The images were analyzed by densitometry by using image analysis software (*UN-SCAN-IT* gel v. 5.1 from Silk Scientific, Orem, UT, USA) as described (36).

Immunoprecipitation

Nuclear extracts prepared as described (24,25) were immunoprecipitated (4 h, 4°C) on anti-I κ B α -agarose. The immune complexes were washed 4 times with PBS buffer, and the resulting immunoprecipitated proteins were resolved on 10% SDS gel and detected with I κ B α and p65 antibodies as described (25).

Indirect Immunofluorescence Microscopy

Neutrophils were fixed with 2% paraformaldehyde. After washing, the cells were resuspended in PBS, cytopun onto slides, and permeabilized with 0.5% Triton X-100 for 10 min. After blocking (1 h at room temperature and overnight at 4°C) in PBS containing 10% bovine serum, the cells were incubated (2 h at room temperature) with anti-I κ B α antibody diluted 1:20, anti-p65 antibody diluted 1:100, or anti-p50 antibody diluted 1:20 in PBS containing 2.5% bovine serum and 0.01% Tween-20. The cells were washed and incubated (1 h at room temperature) with FITC-conjugated secondary anti-rabbit IgG antibody diluted (1:100) in PBS containing 2.5% bovine serum and 0.01% Tween-20. The cells were washed, incubated (10 min) with

4'6'-diamidino-2-phenylindole (DAPI) to visualize DNA, and after washing, mounted onto coverslips. The slides were observed using a Nikon Eclipse 800 microscope, and only experimental series that showed no signal in the absence of primary antibodies were analyzed.

Confocal Microscopy

Neutrophils were fixed and permeabilized as described above. After blocking, the cells were incubated (2 h at room temperature) with anti-I κ B α antibody diluted 1:10 in PBS containing 2.5% bovine serum. The cells were washed and incubated (1 h at room temperature) with FITC-conjugated secondary anti-rabbit IgG antibody diluted 1:50 in PBS containing 2.5% bovine serum. The cells were washed, incubated (10 min) with propidium iodide (PI) to visualize DNA, and mounted onto coverslips. The slides were observed using a Leica TCS SL confocal microscope.

Subcellular Fractionation

The subcellular fractions were prepared by the procedure of He et al. (43). Briefly, the cells were lysed in cytoskeleton (CSK) buffer (10 mM PIPES, pH 6.8, 0.1 M NaCl, 300 mM sucrose, 3 mM MgCl₂, 1 mM EGTA, 0.5% Triton X-100, 2 mM DTT) containing the protease and phosphatase inhibitors as described (24) and centrifuged (3 min, 5000 rpm =1700 g). The supernatants were collected and labeled as soluble proteins. The nuclear pellets were washed with CSK buffer; the chromatin associated proteins were extracted by incubation with DNase I (15 min, 37°C), followed by incubation with 0.25 M ammonium sulfate (5 min, 4°C). The samples were centrifuged (5 min, 5000 rpm =1700 g), and the supernatants were collected and labeled as a chromatin fraction. The pellets were washed with CSK buffer containing 2M NaCl; the supernatants were collected and labeled as a chromatin wash. Supernatants from each extraction step and the final nuclear matrix pellets were boiled in SDS sample buffer and analyzed by SDS-electrophoresis and immunoblotting.

ELISA

TNF α and IL-8 release was measured in cell culture supernatants using commercially available ELISA kits (R&D) as described (26).

Statistical Analysis

The results represent at least three independent experiments. Numerical results are presented as means \pm SE. Data were analyzed using an InStat software package (GraphPAD, San Diego, CA, USA). Statistical significance was evaluated using Mann-Whitney *U* test with Bonferroni correction for multiple comparisons, and *P* < 0.05 was considered significant.

RESULTS

Prolonged Stimulation of Human Neutrophils with LPS or TNF α Induces Persistent Activation of NF κ B That Is Independent of the Newly Synthesized I κ B α

Because it is the persistent NF κ B activation that is associated with the pathogenesis of sepsis and ALI (28-33,44), we hypothesized that human neutrophils stimulated with proinflammatory signals for prolonged times may exhibit persistent or intermittent activation of NF κ B. To measure the NF κ B activity in persistently stimulated human neutrophils, neutrophils were stimulated with LPS (100 ng/mL) or TNF α (10 ng/mL) for up to 12 h, and NF κ B DNA binding activity was measured in nuclear extracts by EMSA. As illustrated in Figure 1A (left panel), neutrophil stimulation with LPS induced NF κ B activity that typically had a biphasic character: the previously observed NF κ B activity peaking around 30 min (24,25), and a second phase of NF κ B activation peaking around 9 h after stimulation. Neutrophil stimulation with TNF α induced, in addition to the early NF κ B activation at 30-60 min (34-41), a strong and sustained activation of NF κ B, which in terms of magnitude of activation, far exceeded the early phase of NF κ B activation induced by neutrophil stimulation for 30 min. This persistent

NF κ B activity reached the maximum around 9 h after stimulation, and lasted up to 12 h (Figure 1A, right panel).

To determine whether the persistent NF κ B activity was composed of the p50 and p65 NF κ B subunits like the NF κ B activity induced at 30 min (36), we performed a supershift analysis of the NF κ B complexes induced by LPS and TNF α stimulation for 9 h. Figure 1B shows that the NF κ B activity induced by neutrophil stimulation with LPS (left panel) and TNF α (right panel) for 9 h consisted of subunits similar to those of the 30-min induced NF κ B activity (36): p50/50 homodimers and p50/65 heterodimers.

We have previously shown that the immediate NF κ B activation induced by neutrophil stimulation for 30 min is mediated by degradation of nuclear I κ B α , which is later resynthesized and translocates again to the nucleus (25). Figure 1C illustrates the cytoplasmic and nuclear levels of I κ B and NF κ B proteins during neutrophil stimulation with LPS (left panels) and TNF α (right panels) for up to 12 h. As shown in Figure 1C, after the cytoplasmic and nuclear I κ B α was degraded within 30 min after neutrophil stimulation, it was followed by reappearance of I κ B α in both cytoplasm and nucleus. Although I κ B α levels were reduced during the later time points (6-12 h), there were still considerable amounts of I κ B α present in the nucleus at the time of maximal NF κ B activity 9 h after stimulation. The cytoplasmic and nuclear levels of NF κ B p50 and p65 proteins during neutrophil stimulation with LPS and TNF α did not correlate with the NF κ B activity (Figure 1).

To determine whether NF κ B activity in the neutrophils might be regulated by other I κ B proteins, we analyzed the intracellular levels of I κ B β and I κ B ϵ . The I κ B β expression was analyzed by two antibodies raised against the C-terminus (C-20) and N-terminus of I κ B β (N-20). Although these antibodies previously identified I κ B β in other human cells (42), no reacting protein was detected in the neutrophils, indicating that human neutrophils do not express the I κ B β protein

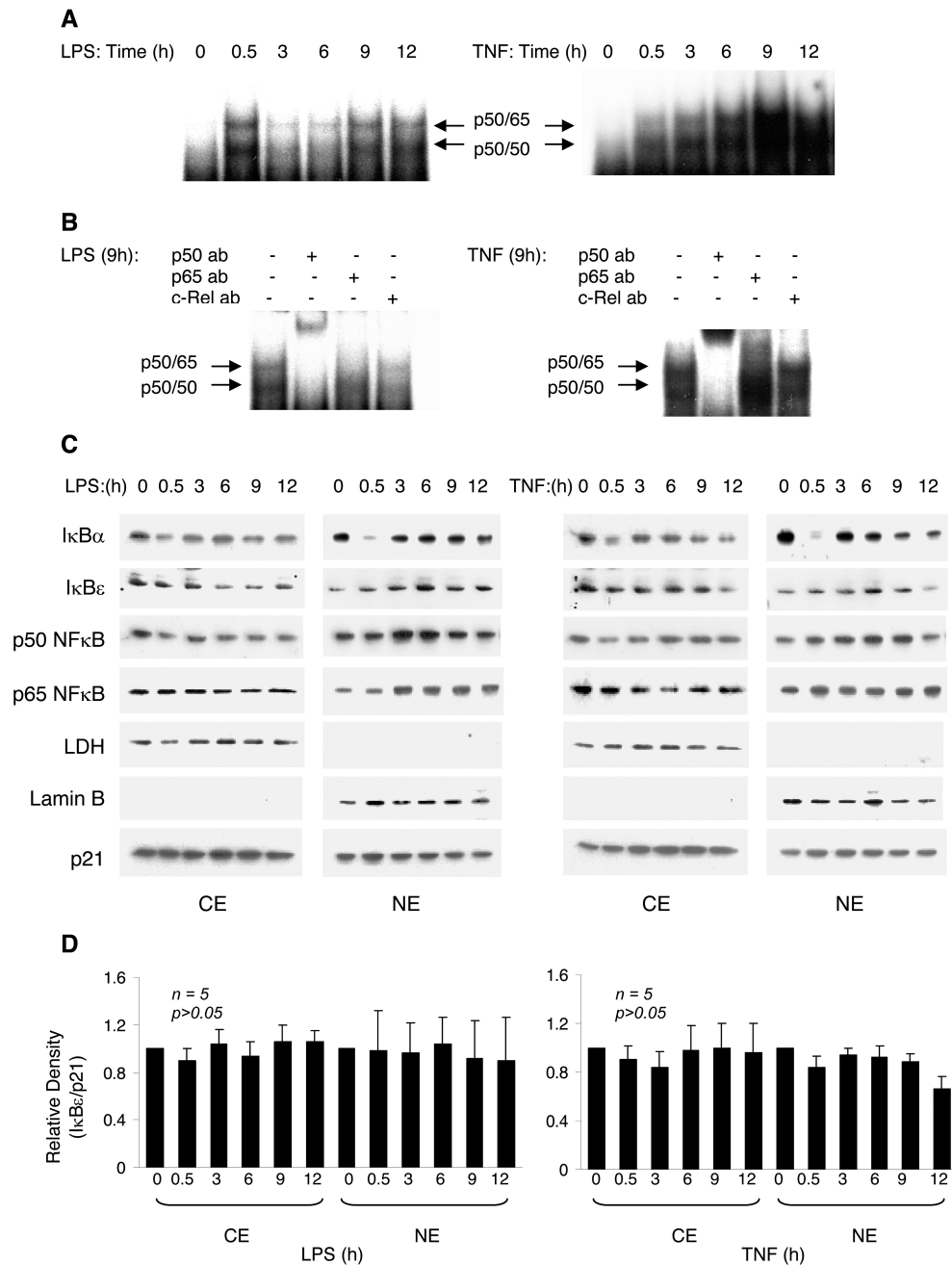


Figure 1. NFκB activity and cytoplasmic and nuclear levels of IκB and NFκB proteins during persistent neutrophil stimulation with LPS and TNFα. (A) Neutrophils were stimulated with LPS (100 ng/mL) or TNFα (10 ng/mL) for 0, 0.5, 3, 6, 9, and 12 h, and NFκB activity was measured in nuclear extracts by EMSA. (B) Supershift analysis of NFκB complexes induced in human neutrophils after 9-h stimulation with LPS and TNFα. Nuclear extracts prepared from neutrophils stimulated with LPS (100 ng/mL) and TNFα (10 ng/mL) for 9 h were incubated with NFκB antibodies specific against p50, p65, and c-Rel subunits. (C) Neutrophils were stimulated with LPS (100 ng/mL) or TNFα (10 ng/mL) for 0, 0.5, 3, 6, 9, and 12 h, and the cytoplasmic (CE) and nuclear (NE) levels of IκBα, IκBε, and p50 and p65 NFκB proteins were analyzed by immunoblotting. The presence of cytoplasmic proteins in the nuclear fraction was monitored using lactate dehydrogenase (LDH) antibody. Nuclear contamination in the cytoplasmic fraction was assessed using lamin B-specific antibody. Each lane contains approximately 5×10^5 cells. (D) Densitometric evaluation of IκBε levels in stimulated human neutrophils. The IκBε bands were scanned and the densities were normalized to densities of p21 used as a loading control. The values at T = 0 for CE and NE were arbitrarily set to 1, and the other values are presented relative to those values. The data represent the means of five independent experiments \pm SE.

(data not shown). Our data showed that IκBε is expressed in human neutrophils and is localized in both cytoplasm and nucleus (Figure 1C). Because the IκBε protein has not been previously described in human neutrophils, we analyzed the intracellular levels of IκBε in LPS and TNFα-stimulated neutrophils by densitometry (Figure 1D). The changes in the IκBε cytoplasmic or nuclear levels were not statistically significant ($P > 0.05$), and the IκBε levels did not correlate with the NFκB activity, suggesting that IκBε degradation does not play an important role in the regulation of transcriptional NFκB activity in human neutrophils.

To analyze the cellular levels of IκBα and NFκB proteins in resting and stimulated neutrophils at the level of individual cells, we performed indirect immunofluorescence microscopy. As shown in Figure 2, in the resting neutrophils, IκBα, p50, and p65 NFκB proteins displayed both cytoplasmic and nuclear staining. After neutrophil stimulation with LPS for 30 min, IκBα was degraded and exhibited reduced cellular staining. No substantial differences were observed in the immunofluorescence staining of the NFκB p50 and p65 subunits. These results, and the immunoblotting data shown in Figure 1, indicate that in human neutrophils stimulated with inflammatory signals for 30 min, the cellular levels of IκBα, and not the NFκB subunits, serve as a better marker for NFκB activity. When neutrophils were stimulated with LPS for 9 h, the newly synthesized IκBα was no longer degraded, and its cellular levels were comparable to those in the resting neutrophils. Similarly, the nuclear and cytoplasmic levels of NFκB p50 and p65 subunits at 9 h after stimulation were essentially unchanged from the resting neutrophils (Figure 2). Similar results were observed when neutrophils were stimulated for 30 min and 9 h with TNFα instead of LPS (data not shown). Figure 3 illustrates the intracellular localization of IκBα analyzed by confocal immunofluorescence microscopy in resting neutrophils and neutrophils stimulated with LPS for 3 h. Consistent with the indirect immunofluorescence microscopy (Figure 2), in both resting and stimulated neutrophils, IκBα was local-

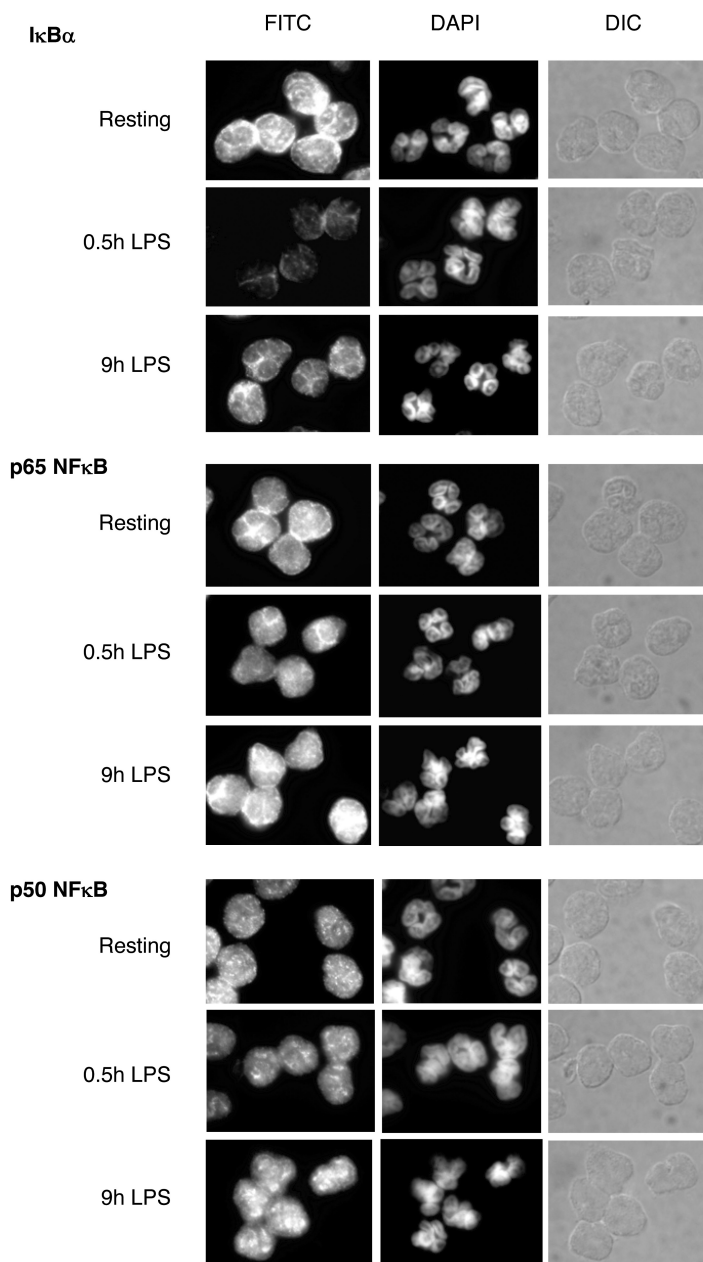


Figure 2. Indirect immunofluorescence microscopy of IκBα and NFκB p50 and p65 proteins during neutrophil stimulation with LPS (100 ng/mL). The left panels show FITC staining of IκBα and NFκB p50 and p65 proteins. The middle panels illustrate DNA nuclear staining with 4′,6′-diamidino-2-phenylindole (DAPI). The right panels show differential interference contrast (DIC) microscopic images of the cells.

ized both in the cytoplasm and in the nucleus; however, the punctuated staining of IκBα in the nucleus indicated that it might be binding to intranuclear components or structures.

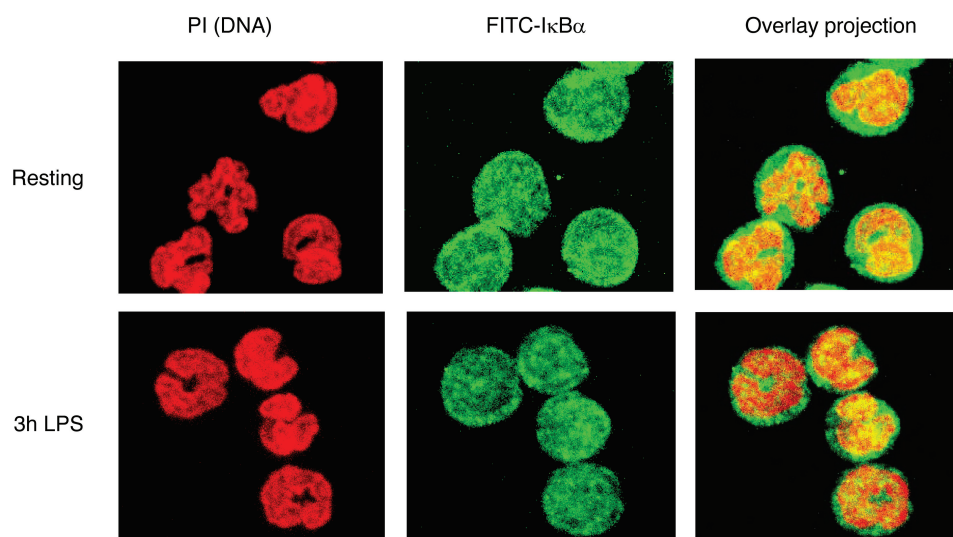


Figure 3. Confocal immunofluorescence microscopy of $\text{I}\kappa\text{B}\alpha$ in LPS-stimulated neutrophils. Resting neutrophils and neutrophils stimulated for 3 h with LPS (100 ng/mL) were fixed and analyzed by confocal laser scanning microscopy using anti- $\text{I}\kappa\text{B}\alpha$ antibody. Left panels show DNA staining with propidium iodide (PI). The middle panels illustrate $\text{I}\kappa\text{B}\alpha$ localization by using FITC staining. The right panels show overlap of the DNA and FITC staining.

$\text{I}\kappa\text{B}\alpha$ and NF κB p50 and p65 Proteins Associate with the Nuclear Matrix in Human Neutrophils

To gain insight into the subnuclear localization of $\text{I}\kappa\text{B}\alpha$ and NF κB proteins during neutrophil stimulation, we isolated soluble chromatin and nuclear matrix fractions by the sequential extraction procedure of He et al. (43). In the first extraction step, soluble (cytoplasmic and nuclear) proteins are removed by extraction with Triton X-100. Chromatin-bound proteins are then released by DNase digestion and extraction with ammonium sulfate. After washing with 2 M NaCl, the last fraction is composed of structural nuclear proteins and the nuclear matrix-associated proteins. Supernatants from each extraction step and the final nuclear matrix pellets were analyzed by SDS-PAGE and immunoblotting. The purity of the cellular fractions was monitored by using lamin B (nuclear matrix marker), histone H2A (chromatin marker), and lactate dehydrogenase (LDH; soluble cytoplasmic protein). Surprisingly, as shown in Figure 4A, in the resting neutrophils, the vast majority of $\text{I}\kappa\text{B}\alpha$ was associated with the nuclear matrix, also containing the NF κB p65

and p50 subunits. Neutrophil stimulation with LPS for 30 min induced degradation of $\text{I}\kappa\text{B}\alpha$; however, the intracellular levels and localization of NF κB p50 and p65 proteins remained essentially unchanged. Neutrophil stimulation with LPS for 9 h induced resynthesis of $\text{I}\kappa\text{B}\alpha$, which was again localized in the nuclear matrix. As in the neutrophils stimulated with LPS for 30 min, 9-h stimulation did not have any significant effect on the nuclear levels of NF κB p50 and p65 subunits (Figure 4A). Very similar results were observed when neutrophils were stimulated with TNF α (data not shown).

The above data suggested that the nuclear $\text{I}\kappa\text{B}\alpha$ may associate with NF κB proteins in the nucleus of human neutrophils. To test this hypothesis, we immunoprecipitated $\text{I}\kappa\text{B}\alpha$ from the nuclear extracts of resting and stimulated neutrophils and analyzed the associated p65 NF κB by immunoblotting. As shown in Figure 4B, in the resting neutrophils, p65 NF κB coimmunoprecipitated with the nuclear $\text{I}\kappa\text{B}\alpha$. In neutrophils stimulated with LPS for 30 min, the nuclear $\text{I}\kappa\text{B}\alpha$ was degraded, and thus neither $\text{I}\kappa\text{B}\alpha$ nor p65 NF κB was recovered on anti- $\text{I}\kappa\text{B}\alpha$ -agarose. In neutrophils stimu-

lated with LPS for 9 h, the newly synthesized $\text{I}\kappa\text{B}\alpha$ was again immunoprecipitated on anti- $\text{I}\kappa\text{B}\alpha$ -agarose. Interestingly, p65 NF κB coimmunoprecipitated with the nuclear $\text{I}\kappa\text{B}\alpha$ from the nuclear extracts of neutrophils stimulated for 9 h, indicating that the p65 NF κB binding by the newly synthesized nuclear $\text{I}\kappa\text{B}\alpha$ is not sufficient to prevent the persistent NF κB activation in continuously stimulated neutrophils.

Release of NF κB -Dependent Mediators During Continuous Neutrophil Stimulation

To investigate whether the prolonged NF κB activation correlates with the synthesis of NF κB -dependent mediators, we measured release of IL-8 and TNF α from LPS-stimulated neutrophils. As illustrated in Figure 5, whereas the TNF α release was induced early (around 3 h after stimulation), after the early activation of NF κB in LPS-stimulated neutrophils, IL-8 was released after 9 h of stimulation, after the second wave of NF κB activation. Neutrophils cultured for 18 h in the absence of LPS released only 0.4% of IL-8 compared with LPS-stimulated PMN (200 ± 74 pg/mL compared with $55,533 \pm$

6,043 pg/mL, *n* = 3), and 6% of TNFα compared with LPS-stimulated PMN (11 ± 2 pg/mL compared with 194 ± 21 pg/mL, *n* = 3).

DISCUSSION

In this study, we demonstrate that TNFα- and LPS-stimulated neutrophils exhibit a previously unrecognized NFκB activation that lasts up to 12 h. In contrast to the early activation of NFκB that is regulated by the nuclear levels of IκBα (24-26), this delayed NFκB activity in 9 h stimulated neutrophils is increased despite the presence of IκBα in the nucleus. Based on this and previous studies from our laboratory (25,26), we propose a new model of NFκB regulation during persistent neutrophil stimulation with inflammatory signals. In the resting neutrophils, IκBα is localized in the nuclear matrix that also contains NFκB p50 and p65 subunits, and by binding to p65 NFκB it inhibits NFκB activity. After neutrophil stimulation with proinflammatory signals, both the nuclear and cytoplasmic IκBα are degraded within 30 min, thus releasing p65 NFκB from the inhibitory complex to bind to NFκB-dependent promoters (25). Our results indicate that during persistent neutrophil stimulation, IκBα is again synthesized and translocates to the nucleus (Figure 1C). This is supported by our previous study demonstrating that cycloheximide treatment prevents the reappearance of both nuclear and cytoplasmic IκBα after 2-h neutrophil stimulation with LPS or TNFα (25). Thus, it seems that the newly synthesized nuclear IκBα that associates with p65 NFκB in the nuclear matrix (Figure 4), is no longer sufficient to inhibit the persistent NFκB activity. The reduced ability of the nuclear IκBα to inhibit NFκB DNA binding at 9 h after stimulation suggests that the newly synthesized nuclear IκBα may be post-translationally modified, and its binding to p65 NFκB does not inhibit, or may even stimulate, the NFκB DNA binding. Alternatively, the concentration of the newly synthesized IκBα in the nucleus might be lower than the concentrations

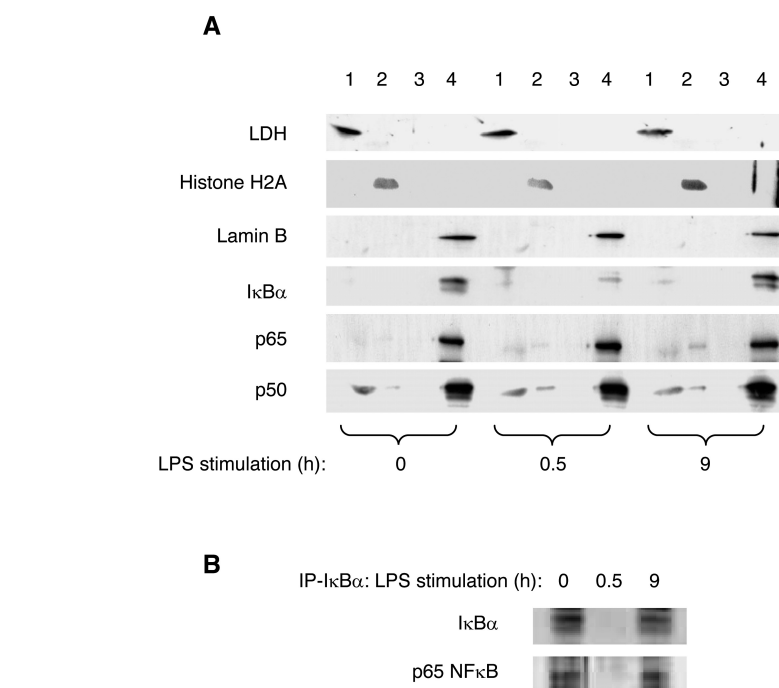


Figure 4. IκBα associates with NFκB in the nuclear matrix of human neutrophils. (A) Neutrophils were stimulated with LPS (100 ng/mL) for 0, 0.5, and 9 h and fractionated to prepare soluble (1), chromatin (2), chromatin wash (3), and nuclear matrix (4) fractions. The fractions were analyzed by Western blotting using LDH, histone H2A, lamin B, IκBα, and p50 and p65 NFκB antibodies. Each lane contains approximately 10⁶ cells. (B) Neutrophils were stimulated with LPS (100 ng/mL) for 0, 0.5, and 9 h, and the nuclear extracts were immunoprecipitated on anti-IκBα-agarose. The immunoprecipitated proteins were detected by immunoblotting using IκBα and p65 NFκB antibodies.

of NFκB proteins. In addition, it cannot be completely ruled out at present that even though IκBε is not degraded after neutrophil stimulation, it can still regulate the transcriptional activity of NFκB. Studies are in progress to discriminate between these models, and to identify the exact mechanisms by which the nuclear IκB proteins regulate the NFκB-dependent transcription in stimulated human neutrophils.

The nuclear matrix is a specialized proteinaceous nuclear structure that serves as a scaffold for chromatin loops. Studies have suggested that the nuclear matrix is tightly associated with transcriptionally active, but not inactive, DNA (45,46). To our knowledge, this study provides the first demonstration that IκBα associates with the nuclear matrix. Our results are consistent with the

morphological study of Trubiani et al. (47) showing that in stimulated epithelial cells, the subnuclear localization of NFκB p50 and p65 proteins is the nuclear matrix. We suggest that the nuclear localization of IκBα and attachment to nuclear matrix represent one of the underlying mechanisms for the decreased activation of NFκB in human neutrophils (34,48) and for the increased apoptosis of these cells.

The subunit composition of NFκB complexes activated in LPS- and TNFα-stimulated neutrophils at 9 h (Figure 1B) is the same as the NFκB subunit composition at 30 min (36): p50/50 homodimers and p50/65 heterodimers. In addition, Western blotting (Figures 1 and 4) as well as immunofluorescence microscopy (Figure 2) revealed that the p50 and p65 NFκB nuclear protein levels are not

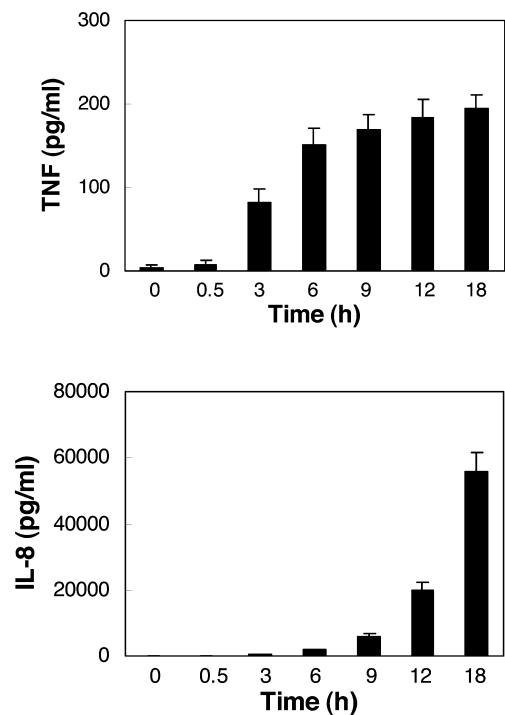


Figure 5. TNF α and IL-8 release from LPS-stimulated neutrophils. TNF α and IL-8 release from LPS (100 ng/mL)-stimulated neutrophils (5×10^6 /mL) was assayed from cell culture media. Data for both panels are expressed as mean values of three independent experiments \pm SE.

substantially changed during neutrophil stimulation with TNF α or LPS, indicating that the persistent NF κ B activity in stimulated human neutrophils is not regulated primarily by the nuclear translocation of NF κ B subunits. Thus, when assessing NF κ B activity during neutrophil-mediated inflammatory disorders, it is important to analyze the extent of NF κ B activity by EMSA, and not only the nuclear levels or translocation of NF κ B subunits.

There is an urgent need for early detection and safer and more specific therapies that minimize tissue injury for ALL, sepsis, and BPD. Neutrophil apoptosis, regulated by NF κ B, plays a critical role in the resolution of inflammation associated with these disorders (28-32,49). Although studies have shown that NF κ B is activated in patients with ALL, sepsis, and BPD (4,30-33,44), the stage of NF κ B activation in these patients is not known. It seems plausible that it is the persistent

NF κ B activity, and not the early NF κ B activation induced by neutrophil stimulation for 30 min, that is associated with the tissue injury in these inflammatory disorders. Whereas TNF α , one of the first mediators of sepsis (50,51), is released early during neutrophil stimulation, IL-8 is released during later time points (Figure 5). We hypothesize that different genes may be activated by NF κ B during different times of neutrophil stimulation. In this scenario, the TNF α transcription would be expected to be induced by the early NF κ B activation in LPS-stimulated neutrophils, and the IL-8 transcription during the second wave. It might be possible that the second wave of NF κ B activation in LPS-stimulated neutrophils is caused by the release of TNF α or other NF κ B-regulated mediator. However, this does not seem very likely, because the second wave of NF κ B activation in LPS-stimulated neutrophils is not associated with the degradation of cytoplasmic or

nuclear I κ B α (Figure 1). A detailed analysis by chromatin immunoprecipitation of the nuclear I κ B and/or NF κ B complexes that are recruited to NF κ B promoters during neutrophil stimulation will be essential to identify the specific NF κ B-regulated proinflammatory and anti-apoptotic genes synthesized during different stages of neutrophil activation.

Previous studies have shown that the early activation of NF κ B is regulated by PKC δ , I κ B kinase, and p38 MAP kinase and by PP1/PP2A phosphatases (24,27,41,26); however, the mechanisms that regulate the persistent NF κ B activity during continuous neutrophil stimulation are unknown. In addition, inhibitors of NF κ B, such as dexamethasone or curcumin, inhibit production of NF κ B-dependent proinflammatory cytokines and induce neutrophil apoptosis (52,53). It will therefore be important to identify the upstream signaling mechanisms that regulate the persistent NF κ B activity and to determine whether this delayed NF κ B activity in continuously stimulated human neutrophils can be inhibited by antiinflammatory or proapoptotic drugs like dexamethasone and curcumin.

Previous work from our laboratory has demonstrated that there are significant differences in the NF κ B regulation between human neutrophils and other cells, such as monocytic cells (24-26). The early stage of NF κ B activation in human neutrophils is regulated predominantly by the nuclear levels of I κ B α (24-26). This study shows that NF κ B is persistently activated in human neutrophils stimulated with LPS or TNF α , but this persistent NF κ B activation is independent of the nuclear levels of I κ B α . A better understanding of the mechanisms regulating the initiation and persistence of NF κ B activation in human neutrophils may provide new information for the development of early detection and safer therapies for BPD, ALL, sepsis, and other neutrophil-mediated inflammatory disorders.

ACKNOWLEDGMENTS

This work was supported in part by National Institutes of Health Grant

HD39643 and by the St. John's University Faculty Research Award to IV.

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