Surfactant protein A suppresses reactive nitrogen intermediates by alveolar macrophages in response to Mycobacterium tuberculosis

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Myobacterium tuberculosis attaches to, enters, and replicates within alveolar macrophages (AMs). Our previous studies suggest that surfactant protein A (SP-A) can act as a ligand in the attachment of M. tuberculosis to AMs. Reactive nitrogen intermediates (RNIs) play a significant role in the killing of mycobacteria. We have demonstrated that RNI levels generated by AMs were significantly increased when interferon-γ–primed AMs were incubated with M. tuberculosis. However, the RNI levels were significantly suppressed in the presence of SP-A (10 µg/ml). The specificity of SP-A’s effect was demonstrated by the use of F(ab’)2; fragments of anti–SP-A monoclonal antibodies and by the use of mannosyl-BSA, which blocked the suppression of RNI levels by SP-A. Furthermore, incubation of deglycosylated SP-A with M. tuberculosis failed to suppress RNI by AMs, suggesting that the oligosaccharide component of SP-A, which binds to M. tuberculosis, is necessary for this effect. These results show that SP-A–mediated binding of M. tuberculosis to AMs significantly decreased RNI levels, suggesting that this may be one mechanism by which M. tuberculosis diminishes the cytotoxic response of activated AMs.


Introduction

Myobacterium tuberculosis remains an important cause of pulmonary infection and is the leading cause of morbidity and mortality worldwide (1–3). After inhalation, M. tuberculosis organisms enter alveolar macrophages (AMs), survive and replicate in the host cell, and eventually produce pathogenic effects (4–11). Thus, attachment of M. tuberculosis to AMs is likely a necessary requirement for the establishment of the initial tuberculous infection in the alveolar spaces. However, much remains unknown about the mechanisms by which M. tuberculosis survive and replicate inside AMs.

Several studies have suggested the involvement of multiple receptors (CR1, CR3, mannose receptor, transferrin receptor, CD14, C2a component of complement, and an unknown receptor that is inhibited by β-glucan) on the surface of macrophages that mediate the binding and phagocytosis of M. tuberculosis organisms (12–18). All of these receptors have been implicated as potential mediators of attachment of M. tuberculosis. The diversity of entry mechanisms suggests that the route of entry may influence the fate of ingested tubercle bacilli in AMs (19).

Macrophages have the ability to recognize and kill invading microorganisms (20). After inhalation, M. tuberculosis are rapidly ingested by AMs. Ingestion of M. tuberculosis by AMs leads to the secretion of cytokines that subsequently affect the intracellular survival of mycobacteria (21). Activation of macrophages has been identified as important for controlling growth of the microorganisms. Activated macrophages produce reactive nitrogen intermediates (RNIs) that are highly toxic to various intracellular pathogens (22). These include RNIs produced by the nitric oxide synthase/arginine–dependent pathway in macrophages and are thought to represent a major killing mechanism of mycobacteria in vivo (23–25). Administration of M. tuberculosis in vivo induces production of RNI by rat AMs (26). Furthermore, nitric oxide synthase knockout mice are highly susceptible to M. tuberculosis infection (27).

Surfactant protein A (SP-A) has a multifunctional role in the lung (28). SP-A, the major protein component of surfactant, is a C-type lectin and contains a region on the molecule known as the carbohydrate recognition domain (29, 30). The carbohydrate recognition domain shares several structural features with complement factor C1q and mannose-binding protein. SP-A functions as a nonimmune opsonin for a variety of bacterial pathogens and viruses (31–34). It is also thought that SP-A plays an important role in the modulation of the inflammatory and immunological response (35). Recent studies suggest that SP-A alters oxygen radical production (36, 37) and blocks the costimulatory signals crucial for in vitro T-lymphocyte cell activation (38). AMs incubated with SP-A have a decrease in superoxide production, indicating a dampening of the respiratory burst (36, 38) and suggesting that SP-A has a protective role against the oxidant injury caused by AMs in the lung.
Others, however, have found SP-A to stimulate the respiratory burst of AMs (39, 40). The reasons for these different findings are not completely understood but may be related to different methods used to purify SP-A.

Subjects with HIV are at increased risk for tuberculosis even before there is significant depletion of CD4+ lymphocytes (41). A recent study conducted by our laboratory indicates that bronchoalveolar lavage (BAL) fluid from HIV-infected individuals increases attachment of M. tuberculosis to AMs (42). The factor in the lavage fluid that increased M. tuberculosis attachment is SP-A. However, it is not clear how SP-A might affect the survival or growth of M. tuberculosis within AMs.

To determine possible underlying mechanisms, we have examined the production of RNI by interferon-γ (IFN-γ) –activated murine AMs in response to M. tuberculosis and have demonstrated that SP-A–mediated attachment of M. tuberculosis to AMs inhibited RNI production by AMs. This RNI inhibitory effect of SP-A was reversed significantly by the addition of anti–SP-A antibody or mannosyl-BSA. Furthermore, deglycosylated SP-A did not have a significant effect on RNI production, suggesting that the oligosaccharide component of SP-A is necessary for this inhibitory effect. Finally, in addition to inhibiting RNI production, SP-A–mediated attachment was associated with enhanced growth of M. tuberculosis in AMs, suggesting one possible mechanism by which the mycobacteria may have enhanced survival.

**Methods**

*M. tuberculosis isolation.* The H37Ra strain of *M. tuberculosis* (American Type Culture Collection, Rockville, Maryland, USA) was cultured at 37°C in a 5% CO2 atmosphere in dispersed form in Middlebrook 7H9 broth (Difco Laboratories, Detroit, Michigan, USA) containing albumin, dextrose, and catalase as enrichments. Bacilli was used in the assay. Each batch of the bacterial suspension was briefly sonicated (20 W for 15 min) and stirred for 30 min. The butanol extract was centrifuged at 15,000 × g and finally resuspended in 5 mM Tris buffer (pH 7.4) containing albumin, dextrose, and catalase as enrichments. Bacterial cultures 10–14 days old were centrifuged, washed once with saline, and the final concentration of the bacterial suspension was adjusted to 10⁶ organisms/ml. Bacterial concentration was determined using a Spectronic 20D spectrophotometer (43) (Milton Roy Co., Rochester, New York). To achieve a single cell suspension, the mycobacterial suspension was briefly sonicated (20 W for 5–10 s). Then the suspension was gently agitated and allowed to stand for 5 min. The top portion of the suspension containing bacilli was used in the assay. Each batch of the bacterial suspension was stained with Kinyoun stain (Midatlantic Biomedical Inc., Paulsboro, New Jersey, USA) and observed under a microscope to check the purity of the suspension. Routinely, samples of bacteria were also grown on 7H11 Middlebrook medium (Difco Laboratories) and maintained as stock.

**SP-A isolation and purification.** Human SP-A was isolated according to Wright et al. (44). Briefly, a surfactant pellet was obtained by centrifugation (10,000 g for 60 min) of BAL fluid from patients with alveolar proteinosis then extracted with butanol. The butanol-insoluble proteins were resuspended in octylglucoside to solubilize serum proteins. SP-A purified by this method was dialyzed extensively against 5 mM Tris buffer (pH 7.4), then quantified by the Lowry protein assay (45). SP-A purity was verified by SDS-PAGE (46). Deglycosylated SP-A was prepared according to the method of Gaynor et al. (47).

Rat SP-A was isolated according to Kuroki et al. (48), with minor modifications. BAL was obtained from rats that had received intratracheal injection of silica (0.25 mg/rat) 4 weeks before sacrifice. The lavage was centrifuged at 22,000 g for 18 h. The pellet (4 ml from 10 rats) was injected into 200 ml of butanol and stirred for 30 min. The butanol extract was centrifuged at 15,000 g for 30 min. The pellet was resuspended in butanol, centrifuged, and finally resuspended in 5 mM Tris buffer (pH 7.4) and dialyzed against the same buffer. Insoluble material was removed by centrifugation at 60,000 g for 1 h. The supernatant was brought up to 10 mM CaCl₂, passed through a fucose-binding agarose column (Sigma Chemical Co., St. Louis, Missouri, USA), and washed with approximately 10 ml of 5 mM Tris containing 10 mM CaCl₂ (pH 7.8). Calcium-dependent fucose-binding proteins were eluted with 5 mM Tris, 2 mM EDTA. SP-A stocks were incubated with polymyxin agarose to reduce endotoxin contamination to undetectable levels (49). Human SP-A was used in most experiments to determine the effects of SP-A on AM response to M. tuberculosis; rat SP-A was used in select studies to verify that the effects were not simply species specific.

**Production of murine monoclonal antibodies to human SP-A.** Murine anti–human SP-A monoclonal antibodies (MAbs) were generated according to the method of Kohler and Milstein (50), with minor modifications. Briefly, BALB/c mice were immunized with 100 μg of purified SP-A emulsified in Hunter’s Titermax adjuvant (CytRx Corp., Norcross, Georgia, USA) on days 0 and 14. A booster injection without the adjuvant was given on day 28. Three days after the booster injection, spleenocytes harvested from the immune spleen were mixed with mouse myeloma cell lines SP2/O.Ag.14 (American Type Culture Collection) and fused in the presence of 1 ml of 50% polyethylene glycol (Sigma Chemical Co.) for 1 min. After removing the fusing agent by washing with RPMI-1640 (Sigma Chemical Co.), the hybridoma cells were resuspended in hypoxanthine, aminopterin, and thymidine selectors.
tive media containing 10% FCS, glutamine (300 µg/ml), penicillin (100 U/ml), streptomycin (100 µg/ml), gentamycin (40 µg/ml), and nystatin (25 IU/ml) (Sigma Chemical Co.). The hybridoma cells were plated into 24-well tissue culture plates with a concentration of 105 cells/ml and incubated for 2 weeks at 37°C in 5% CO2. After 2 weeks, the culture supernatants were tested for the presence of anti-SP-A antibodies by ELISA. Cultures containing antibodies were cloned by limiting dilution into 96-well tissue culture plates containing feeder cells (51). The positive clones were recloned until the clonal efficiency approached monoclonality. After again screening for the presence of antibodies by ELISA, these MABs were confirmed by Western blotting. The positive clones that secreted antibodies were expanded in the ascites of pristane-(Sigma Chemical Co.) injected BALB/c mice. MABs from the ascites were purified by MAPS II kit (Bio-Rad Laboratories Inc., Hercules, California, USA) and quantified by BCA protein assay (Pierce Chemical Co., Rockford, Illinois, USA). The F(ab')2 fragments were produced according to the manufacturer’s directions (Bio-Rad Laboratories Inc.).

**Interaction of AMs with M. tuberculosis.** Ten-week-old Swiss-Webster pathogen-free mice were purchased from Harlan Sprague Dawley (Indianapolis, Indiana, USA). Murine AMs were isolated by BAL as described previously (52). Briefly, the mice were sacrificed by intraperitoneal injection of Beuthanasia-D solution (Schering-Plough Animal Health Corp., Kenilworth, New Jersey, USA). The trachea was cannulated after a midline neck incision, and the lungs lavaged 10 times with 1.0 ml of 0.9% saline. The organs were cut in small pieces and digested by collagenase (100 U/ml), streptomycin (100 µg/ml), gentamycin 40 µg/ml, and amphotericin B 0.5 µg/ml. Approximately 7–8 ml of lavage fluid was obtained from each mouse. AMs were separated from the lavage fluid by centrifugation at 600 g for 10 min, washed three times with normal saline, and resuspended in DMEM (BioWhittaker Inc., Walkersville, Maryland, USA) with glutamine 300 µg/ml, NaHCO3, and 20 mM HEPES (pH 7.2). Preparations were prepared to contain 98% AMs by routine Diff-Quick (Herleco, Aguada, Puerto Rico) staining and to be >95% viable by trypan blue exclusion. AMs were plated at a density of 2.5 × 106/100 µl/well in 96-well tissue culture plates in an antibiotic-free DMEM. All assays were conducted at 37°C in a 5% CO2 atmosphere.

AMs were primed with or without murine IFN-γ (GIBCO BRL, Grand Island, New York, USA) for 24 h at 37°C in a 5% CO2 atmosphere. Freshly isolated M. tuberculosis organisms were added to adherent AMs in 96-well plates in the ratio of 10:1, and the cultures were incubated for 2 h. After the infection, the AMs were washed to remove any unbound organisms. Fresh medium containing IFN-γ was then added and incubated for various time intervals. To determine the optimal concentration of IFN-γ, AMs were incubated in the presence of IFN-γ (0–1,000 U IFN-γ/ml) for 48 h. After the incubation, the cell-free media was tested to determine the amount of RNI accumulated as described below.

**Measurement of RNIs.** Nitric oxide production was assessed by measuring nitrite. The amount of nitrite formed from NRI was determined from the cell-free supernatants of AMs and M. tuberculosis-infected cultures by the Griess reagent according to Green et al. (53). Briefly, 50 µl of the cell-free supernatant was mixed with 50 µl of freshly prepared Griess reagent (1% sulfanilamide, 0.1% naphthylethylene-diamide-dihydrochloride, and 2.5% H3PO4) and incubated for 10 min at room temperature. After the incubation, the plate was read at 540 nm in an automatic Titertek Multiscan ELISA reader (Flow Laboratories, McLean, Virginia, USA). Nitrite concentrations were determined from a standard curve using sodium nitrite (0–500 µM). The results are expressed as µmol nitrite.ml. Cell viability of AMs after incubation with M. tuberculosis was determined at various time intervals after infecting AMs with M. tuberculosis by the trypan blue exclusion method. The RNI assay was determined using AM cultures with M. tuberculosis alone, in the presence and absence of SP-A (0–40 µg/ml), IFN-γ (100 U/ml), and lipopolysaccharide (LPS) (10 µg/ml).

To determine the possible mechanisms involved in SP-A modulation of RNI production, the assay was performed in the presence or absence of α-methylene-β-mannosepyanidose (mannosyl-BSA) (1,000 µg/ml with a molar ratio of mannose/albumin of 26:1), and F(ab')2 fragments of anti-human SP-A MABs (100 µg/ml), heat-inactivated SP-A (100°C for 10 min), and deglycosylated SP-A alone (10 µg/ml).

Additional experiments were conducted to determine whether SP-A scavenges RNI products. SP-A (10 µg/ml) was incubated with increasing concentrations of sodium nitrite (NaNO3) from 1 µM to 500 µM for 24 h at 37°C. After the incubation, the amount of RNI that accumulated in the supernatant in the presence or absence of SP-A was determined using the Griess reagent.

**Endotoxin assay.** The amount of endotoxin present in all the reagents (media, buffers, bacterial cultures, IFN-γ, SP-A) was assayed by the quantitative chromogenic Limulus amoebocyte lysate assay (BioWhittaker Inc.) in the presence or absence of 100 µg/ml of polymyxin B/ml (Calbiochem-Novabiochem Corp., San Diego, California, USA). Endotoxin concentrations present in the unknown samples were determined from the standard curve using known endotoxin concentrations (0–500 µM). The sensitivity of the assay is <1 ng/ml. Only endotoxin-free SP-A and reagents were used in this study. In addition, all the experiments were conducted in the presence or absence of polymyxin B (100 µg/ml), which has been shown to completely inhibit endotoxin function (54).

**Growth of M. tuberculosis determined by radiometric BACTEC method.** The growth of M. tuberculosis within AMs was determined using a radiometric BACTEC 460 TB system (Becton Dickinson Diagnostic Instrument Systems, Sparks, Maryland, USA) (55, 56). The accuracy of the BACTEC method was confirmed with colony-forming units, a standard method to measure M. tuberculosis growth. The BACTEC method is based on use of 14C substrate present in the medium by mycobacteria and subsequent release of 14CO2 into the atmosphere above the medium. Briefly, AMs were infected with M. tuberculosis at a...
ratio of 1:10 (AMs/organisms). After incubation, AMs infected with M. tuberculosis organisms were washed with DMEM to remove unbound M. tuberculosis organisms. The AMs were lysed by the addition of 50 μl of 0.25% SDS per well for 10 min. After adding 20 μl 20% BSA to neutralize the effect of SDS, the total volume was brought up to 0.5 ml with saline and inoculated into BACTEC 12 Middlebrook vials containing 0.1 ml PANTA, an antibiotic supplement (Becton Dickinson Diagnostic Instrument Systems). The vials were incubated at 37°C, with growth of M. tuberculosis being determined every 24 h by a 460 BACTEC TB instrument (Becton Dickinson Diagnostic Instrument Systems) and expressed as a growth index from 0–999, with an index >10 indicating significant growth.

To determine if SP-A would modulate nitrite-mediated killing of M. tuberculosis in the absence of AMs, NaNO₃ (100 μM) was incubated with M. tuberculosis in the presence of 0.25% SDS per well for 10 min. After adding 20 μl 20% BSA to neutralize the effect of SDS, the total volume was brought up to 0.5 ml with saline and inoculated into BACTEC 12 Middlebrook vials containing 0.1 ml PANTA, an antibiotic supplement (Becton Dickinson Diagnostic Instrument Systems) and expressed as a growth index from 0–999, with an index >10 indicating significant growth.

To determine if SP-A would modulate nitrite-mediated killing of M. tuberculosis in the presence of AMs, NaNO₃ (100 μM) was incubated with M. tuberculosis in the presence of 0.25% SDS per well for 10 min. After adding 20 μl 20% BSA to neutralize the effect of SDS, the total volume was brought up to 0.5 ml with saline and inoculated into BACTEC 12 Middlebrook vials containing 0.1 ml PANTA, an antibiotic supplement (Becton Dickinson Diagnostic Instrument Systems) and expressed as a growth index from 0–999, with an index >10 indicating significant growth.

**Results**

**Effect of IFN-γ on RNI production by AMs.** AMs were treated with various concentrations of IFN-γ for 48 hours at 37°C. RNI production increased as the concentrations of IFN-γ increased, with maximum production occurring at 100 U/ml (Fig. 1). No significant RNI levels were detected with media and IFN-γ alone (0.16 ± 0.9 μM). AMs in the presence of M. tuberculosis alone produced 4.5 ± 2.5 μM of RNI. When AMs were primed with various concentrations of IFN-γ for 24 hours and subsequently incubated with M. tuberculosis for an additional 24 hours at 37°C, RNI production was further increased compared with AMs alone (Fig. 1). Therefore, on the basis of these results, all subsequent experiments with AMs were incubated with 100 U/ml of IFN-γ.

**Effect of SP-A concentrations on RNI production by AMs.** To determine the effect of SP-A on RNI production by IFN-γ-primed AMs, AMs were incubated with various concentrations of SP-A (0–40 μg/ml) for 24 hours and subsequently incubated with M. tuberculosis for an additional 24 hours at 37°C, RNI production was further increased compared with AMs alone (Fig. 1). Therefore, on the basis of these results, all subsequent experiments with AMs were incubated with 100 U/ml of IFN-γ.

**Effect of SP-A on RNI generation.** (a) AMs were incubated with M. tuberculosi alone, and in presence or absence of IFN-γ (100 U/ml), SP-A (5 μg/ml), or LPS (10 μg/ml). These results suggest that SP-A significantly decreased RNI production induced by IFN-γ (P < 0.001); however, addition of LPS to LPS-primed AMs did not significantly decrease RNI levels (P > 0.05). (b) SP-A (10 μg/ml) was incubated with increasing concentrations of sodium nitrite (NaNO₃) from 1 μM to 500 μM for 24 h at 37°C. After the incubation, the amount of RNI that accumulated in the supernatant was determined using the Griess reagent. The RNI levels increased in an NaNO₃ concentration-dependent manner, and there was no significant effect of SP-A on RNI determination. Thus, there is no evidence that SP-A directly scavenges NaNO₃. Results are expressed as mean ± SEM of three experiments performed in triplicate. LPS, lipopolysaccharide.

**Results**

**Effect of IFN-γ on RNI production by AMs.** AMs were treated with various concentrations of IFN-γ for 48 hours at 37°C. RNI production increased as the concentrations of IFN-γ increased, with maximum production occurring at 100 U/ml (Fig. 1). No significant RNI levels were detected with media and IFN-γ alone (0.16 ± 0.9 μM). AMs in the presence of M. tuberculosis alone produced 4.5 ± 2.5 μM of RNI. When AMs were primed with various concentrations of IFN-γ for 24 hours and subsequently incubated with M. tuberculosis for an additional 24 hours at 37°C, RNI production was further increased compared with AMs alone (Fig. 1). Therefore, on the basis of these results, all subsequent experiments with AMs were incubated with 100 U/ml of IFN-γ.

**Effect of SP-A concentrations on RNI production by AMs.** To determine the effect of SP-A on IFN-γ-primed AMs, AMs were incubated with IFN-γ (100 U/ml) and various SP-A concentrations (0–40 μg/ml). RNI production was decreased by SP-A in a concentration-dependent manner (Fig. 2), with maximal inhibition occurring at 10 μg/ml SP-A (P < 0.001). Similarly, RNI
production by IFN-γ-primed AMs incubated with M. tuberculosis was decreased by SP-A in a concentration-dependent manner (Fig. 2), with the maximal decrease occurring at 10 μg/ml SP-A (P < 0.001). Longer incubations of AMs with M. tuberculosis and SP-A had no significant effect on RNI production.

In the presence of LPS, AMs infected with M. tuberculosis generated RNI levels of 7.0 ± 2.4 μM, and the addition of SP-A to the same reaction mixture generated RNI levels of 4.0 ± 2.4 μM (P = 0.294) (Fig. 3a). Furthermore, there was no evidence that SP-A scavenges RNI products (Fig. 3b). Thus, the ability of SP-A to reduce RNI generation by AMs does not appear to be secondary to interference with the nitrite assay.

To determine the specificity of the SP-A effect on RNI production, the assay was performed in the presence of 100 μg/ml purified F(ab')2 fragment of anti–human SP-A. Addition of anti–SP-A F(ab')2 fragments to the mixture of SP-A, M. tuberculosis, and AMs significantly increased RNI production from 10.6 ± 0.7 μM to 18.3 ± 1.2 μM (Fig. 4). A control antibody (normal IgG) did not block the inhibitory effect of SP-A (data not shown). Both human SP-A and rat SP-A significantly inhibited RNI production from 17.5 ± 0.8 μM to 8.1 ± 0.7 μM and 8.8 ± 0.5 μM, respectively (P < 0.05 for both comparisons). To determine possible mechanism(s) involved in the SP-A–decreased RNI production, AMs and M. tuberculosis were preincubated with SP-A in the presence or absence of mannosyl-BSA. Mannosyl-BSA significantly blocked the effects of SP-A on RNI production by IFN-γ–primed AMs in response to M. tuberculosis (Fig. 5a). Furthermore, neither deglycosylated SP-A nor heat-inactivated SP-A (Fig. 5b) had a significant effect on RNI production (17.7 ± 1.7 μM and 16.1 ± 1.6 μM, respectively) compared with control (22.1 ± 2.2 μM). These results indicate that blocking SP-A with mannosyl-BSA or anti–SP-A F(ab')2 fragments prevents its inhibitory effect on RNI production of AMs stimulated by IFN-γ and M. tuberculosis. These results suggest that the oligosaccharide component of SP-A is involved in the SP-A–mediated decrease in RNI production by AMs in response to M. tuberculosis infection.

Effect of SP-A on growth of M. tuberculosis in AMs. Incubation of AMs and M. tuberculosis with SP-A significantly enhanced the growth of M. tuberculosis organisms (expressed as growth index) from 145.6 ± 14.0 to 842.4 ± 97.8. Addition of anti–SP-A F(ab')2 fragments to the mix-

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**Figure 5**

Effect of mannosyl-BSA, denatured SP-A, and deglycosylated SP-A on IFN-γ–induced RNI production by M. tuberculosis–infected AMs. (a) RNI production was examined after preincubation of the IFN-γ–primed AMs and M. tuberculosis with mannosyl-BSA (1,000 μg with a 26:1 molar ratio of mannose/albumin). Mannosyl-BSA significantly blocked the effect of SP-A on the RNI production. (*P < 0.05, comparison with either AMs + M. tuberculosis or AMs + M. tuberculosis + SP-A + mannosyl-BSA.) (b) Incubating either denatured SP-A or deglycosylated SP-A with IFN-γ–primed AMs infected with M. tuberculosis did not significantly suppress RNI production (P > 0.05, both comparisons). Results are expressed as mean ± SEM for six experiments performed in triplicate.

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**Figure 6**

Effect of SP-A on M. tuberculosis growth. Effect of SP-A on M. tuberculosis growth was determined by the BACTEC radiometric method. AMs were infected with M. tuberculosis at a ratio of 10:1 (M. tuberculosis:AM) in the presence of IFN-γ (100 U/ml), SP-A (5 μg/ml), or anti–SP-A antibodies (100 μg/ml). AMs were washed with DMEM to remove unbound M. tuberculosis organisms, and IFN-γ, SP-A, and SP-A antibodies were added back with the fresh medium for 24 h. The AMs were then lysed and inoculated into the BACTEC vials. The vials were then incubated at 37°C, with growth of M. tuberculosis being determined every 24 h by the BACTEC TB instrument. Preincubation of AMs and M. tuberculosis with SP-A significantly increased M. tuberculosis growth. These results suggest that blocking the effect of SP-A again restores the ability of AMs to inhibit the growth of M. tuberculosis.
ture of SP-A, *M. tuberculosis*, and AMs significantly decreased the growth of *M. tuberculosis* from 842.4 ± 97.8 to 124.8 ± 18.7 (Fig. 6). It did not appear that SP-A had any direct effect independent of macrophages, i.e., on diminishing the toxicity of RNIs to the *M. tuberculosis* organisms. When *M. tuberculosis* organisms were grown in the absence of macrophages but in the presence of NaNO₃ (100 μM), growth was decreased from 85.5 ± 143.6 to 476.0 ± 185.4. However, this inhibitory effect of *M. tuberculosis* growth was not affected by the presence of SP-A (465.0 ± 138.1). Thus, SP-A did not alter the growth-inhibiting effects of RNIs on *M. tuberculosis*.

**Discussion**

The results of this study suggest that SP-A suppresses RNI production in IFN-γ-treated AMs in response to *M. tuberculosis* and that the oligosaccharide component of SP-A is critical in this interaction. SP-A suppressed RNI production by IFN-γ-stimulated AMs in a concentration-dependent manner in response to *M. tuberculosis*, and the inhibitory effect of SP-A could be blocked with both anti–SP-A F(ab)², fragments and mannosyl-BSA. Deglycosylated or heat-inactivated SP-A failed to suppress the RNI production. This suggests SP-A–mediated attachment/phagocytosis of *M. tuberculosis* represents one possible mechanism by which *M. tuberculosis* organisms can safely enter its host cell without triggering a cytotoxic response. In this way, AMs may provide a potentially safe habitat whereby the organisms are able to survive and multiply.

Recent investigations have focused on the role of RNIs in host defense against *M. tuberculosis* (22). For instance, IFN-γ-priming of L-arginine–dependent RNI production plays a major role in growth inhibition or killing of *M. tuberculosis* (23–25). Likewise, *in vivo* RNI production was shown to be greatly reduced in IFN-γ knockout mice, and this resulted in unrestricted growth of *M. tuberculosis* (58). CD4 T lymphocytes appear to be the major source of IFN-γ early during the course of infection with *M. tuberculosis* (59). The cytokine IL-12 appears to upregulate IFN-γ, whereas IL-10 downregulates IFN-γ in response to *M. tuberculosis* infection (60, 61). In our study, AMs primed with IFN-γ and incubated with *M. tuberculosis* markedly increased RNI production in a concentration-dependent manner, with maximum production occurring at 100 U IFN-γ/ml.

It is known that *M. tuberculosis* can interact with various AM receptors (12–18), and yet it frequently avoids being killed by the AMs. Specifically, it has been suggested that SP-A– or complement–mediated attachment/phagocytosis by AMs may avoid triggering a cytotoxic response (36, 37, 62). Our study indicates an SP-A concentration-dependent suppression of RNI production by IFN-γ-primed AMs in response to *M. tuberculosis*. This occurs despite the fact that SP-A increases the attachment/phagocytosis of *M. tuberculosis* by AMs (52), resulting in AMs having higher numbers of *M. tuberculosis* organisms intracellularly.

SP-A is well recognized to be an effective innate opsonin in the alveolar spaces as an important part of innate immunity within the lungs (63). SP-A can stimulate directed chemotaxis by alveolar macrophages to microorganisms (64). Furthermore, the role of SP-A in host defense is supported by the recent study of LeVine et al. (65), which demonstrates that SP-A–deficient mice have increased susceptibility to group B streptococcal infection. In contrast to extracellular pathogens such as streptococci, mycobacterial species must enter and subsequently grow within the alveolar macrophage (4–11). Our study suggests that *M. tuberculosis* may use this SP-A–mediated entry mechanism to its own advantage.

Recent studies from our lab indicated that only the glycosylated, not the deglycosylated, form of SP-A binds to *M. tuberculosis* (52). Gaynor et al. (47) also showed that deglycosylated SP-A does not enhance the adherence of *M. tuberculosis* to monocytes. Therefore, the previous studies and the current data suggest that sugars in the carbohydrate recognition domain of SP-A may be important in the ability of SP-A to recognize *M. tuberculosis* and to mediate an inhibitory effect on the alveolar macrophage response.

Contrary to our current study, others have demonstrated a stimulatory influence of SP-A on AMs (39, 40, 66). The reasons why SP-A in some studies appears to stimulate AMs (39, 40, 66) and in others appears to dampen the AM response (36–38) is not clear. There is some evidence that the method of isolation and purification of SP-A may alter SP-A function (39). Also, SP-A appears to avidly bind LPS (67), and preliminary studies suggest SP-A may enhance presentation of LPS to AMs (Stamme, C.S., and Wright, J.R., unpublished data). Similar LPS-binding properties have been described for SP-D (68); thus, even a trace contamination of endotoxin with SP-A might alter the AM response. Further study is clearly needed in defining the interaction of SP-A, endotoxin, and AMs.

It is also recognized that SP-A likely functions differently *in vitro* than in the complex lipid-rich milieu of the alveolar spaces (69). As a result, our *in vitro* studies may not accurately reflect the interactions of SP-A, *M. tuberculosis*, and AMs as they occur *in vivo*. However, our initial study of SP-A’s effect on *M. tuberculosis* and AMs used human BAL fluid; in contrast to culture media and artificial surfactants, the intact BAL contained at least some of the native alveolar constituents that might influence SP-A function (42). Thus, we believe the current study is relevant to the alveolar conditions that might influence the interactions of *M. tuberculosis* and AMs.

As might be predicted, a decrease in RNI production by AMs incubated with SP-A–coated *M. tuberculosis* organisms was associated with a commensurate increase in growth of *M. tuberculosis* within the AMs. Increased *M. tuberculosis* growth may occur for multiple reasons. It is likely that both increased numbers of *M. tuberculosis* organisms in AMs (52) and a decrease in RNI production permit increased growth (22, 23). Our observations suggest that elevation of SP-A in the lung may represent one reason why HIV-infected individuals may be more susceptible to *M. tuberculosis* infections.

Another pulmonary disorder associated with elevated levels of SP-A is silicosis (70), a condition associated with a lifelong predisposition to tuberculosis (71). The mechanisms for the elevation of SP-A in silicosis are unclear,
but this may represent a compensatory mechanism by the lung to facilitate clearance of silica particles from the lower respiratory tract. HIV infection and silicosis are very different conditions; however, it is interesting that both are associated with elevated SP-A levels and both have an increased risk for pulmonary tuberculosis.

Further studies of the mechanisms of *M. tuberculosis* survival in the intracellular environment of the AM may provide important information regarding the earliest stages in the pathogenesis of the disease, and this may permit the development of novel therapeutic strategies to modulate the disease process.

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