Repeated inoculations of *Mycobacterium bovis* Bacille Calmette-Guérin (BCG) are needed to induce a strong humoral immune response against antigens expressed by the bacteria*

Monique C. da Silva¹, Elena B. Lasunskia¹, Wilmar Dias da Silva²#

¹Laboratory of Biology of Recognition, Universidade Estadual do Norte Fluminense, Campos, Rio de Janeiro, Brazil
²Laboratory of Immunochemistry, Institute Butantan, São Paulo, Brazil; #Corresponding Author: wds@butantan.gov.br

Received 18 April 2013; revised 16 May 2013; accepted 30 May 2013

Copyright © 2013 Monique C. da Silva et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

**ABSTRACT**

The cellular immune response elicited by *Mycobacterium bovis* Bacille Calmette-Guérin (BCG) has been carefully investigated, but the humoral immune response has been partially neglected. BALB/c mice were immunized with BCG strain used to immunize humans. Anti-BCG antibodies, as assayed by ELISA, began to appear in the sera after the third week of immunization and plateaued three weeks after the 8th immunization. The total immunoglobulins (Igs) were purified by caprylic acid method from pooled serum collected after the 8th immunization. Anti-BCG antigen antibodies were detected in the total Igs preparation as well as in IgG, IgM, IgA, IgG1, IgG2a, and IgG2b, but not in the IgG3. Distinct BCG proteins were recognized the IgGs in Western blot analysis. Opsonization of BCG bacilli by the purified Igs potentiated internalization of the bacteria by murine Raw 264.7 macrophages. The intracellular BCG elimination coincided with the induction of NO production, which was more pronounced in cells infected with opsonized BCG compared to those infected with the non-opsonized bacteria. Coincidently, the production of NO was also higher in macrophages infected with opsonized BCG (maximal NO production at 48 h of incubation). The obtained results demonstrate that repeated inoculations of BCG effectively activate the humoral immune response, justifying the use of BCG as a live recombinant vaccine vector to insert genes encoding virulence factors controlled by antibodies.

**Keywords:** *Mycobacterium bovis*; Bacille Calmette-Guérin; BCG; Antibodies; Opsonization; Bacterial Killing

**1. INTRODUCTION**

Bacille Calmette-Guérin (BCG) is an attenuated strain of *Mycobacterium bovis* first used to vaccinate children with a high risk of acquiring tuberculosis (TB) [1]. BCG has become the most widely used vaccine, reaching 3 billion vaccinated individuals [2]. A meta-analysis of individuals vaccinated with BCG indicates that the risk of acquiring active TB is reduced by approximately 50 percent [3]. Widely recovered epidemiological data suggest that the efficacy of the BCG vaccine depends on several associated factors, such as the immune state of the recipient [4], the intensity and frequency of the exposure to the mycobacteria prior to vaccination [5], and the recipient age [6,7]. Therefore, the capacity of BCG to induce an immune response is indirectly indicated. Reinforcing this capacity is the well-recognized ability of BCG to live and multiply in macrophage phagosomes, which facilitates the interaction between the BCG antigens and MHC class II molecules, resulting in the activation of a CD4+ Th1 cell immune response. The observed presence of lipoarabinomannan on the BCG outer cell surface and the ability of this complex molecule to act as an immune adjuvant may increase the potential immunogenicity of BCG [8]. This probable immunogenicity

*The results were used by Monique Camila da Silva to obtain her Masters Degree in Biosciences and Biotechnology from the Laboratory of Biology of Recognition, Universidade Estadual do Norte Fluminense-Darcy Ribeiro, with a fellowship from “Coordenação de Apefeiçoamento de Pessoal de Nivel Superior,Brasilia, Brasil” (CAPES).

The authors have no financial conflict of interest.
underlines the inclusion of mycobacteria into complete Freud’s adjuvant to improve the immune response in experimental protocols [9].

The immune response induced by BCG is predominantly T cell-dependent [10]. The cellular immune response has been followed essentially by evaluating in parallel the activated T cell subtypes and the cytokine profile [11]. Studies addressing the T cell response have considered that antibodies play a minor role in the immune protection against tuberculosis. However, antibodies may facilitate mycobacterial elimination by increasing bacterial opsonization [12].

A direct investigation aimed to detect the eventual antibody production in response to BCG was performed by comparing the humoral immune responses induced by BCG strains grown in different culture media [13].

A recent analysis of the immunological response after oral re-vaccination with M. bovis BCG Moreau RDJ in adults who were intradermally immunized in infancy demonstrated an increase in the humoral immune response without tolerance, and a switch from IgG to IgA was detected [14].

Altogether, these data indicate that BCG immunization induces not only cellular immunity but also the humoral immune response. Immunization protocols that lead to high levels of antibody production in response to BCG should be established.

Recently, BCG vaccine strains began to be intensively investigated as vehicles to deliver heterologous antigens to induce both the humoral (mucosal and systemic) and cell-mediated immune responses [15,16], reinforcing the interest in the induction of antibody production by BCG.

Here, we describe an immunization scheme based on repeated booster inoculations with the vaccine and the antibody profiles of prevalent anti-BCG immunoglobulin classes and subclasses induced in mice. Additionally, we evaluated the capacity of these antibodies to opsonize live BCG bacilli for phagocytosis and subsequent destruction within macrophage phagolysosomes. The results demonstrate that the attenuated M. bovis strain BCG is a potent immunogen, which induces a strong humoral response under hyperimmunization conditions, leading to the production of different classes and subclasses of specific immunoglobulins with opsonizing properties.

2. MATERIALS AND METHODS

2.1. Animals

BALB/c mice (8 weeks old, 18 - 22 g, 5 animals per group) were supplied by “Biotério Central da Universidade Estadual do Norte Fluminense-Darcy Ribeiro (UENF)” and maintained at the animal facility of the Laboratório de Biologia do Reconhecer, Centro de Biociências e Biotecnologia, UENF. All animals were cared for under ethical conditions according to International Welfare Recommendations [17,18]. The protocols used were approved by the Institutional Animal Care and Use Committee CBB/UENF, RJ, Brazil.

2.2. Bacterial Strains, Growth Conditions and Whole Extracts

All experiments were performed using the id-BCG Moreau Copenhagen strain (1978) currently utilized in the Brazilian TB immunization program. The BCG samples were supplied by the Institute Butantan, São Paulo, SP, in lyophilized form at the concentration of 1.0 mg/ml -1.0 × 10^5 bacilli/mL. Liquid cultures of BCG were grown in Middlebrook 7H9 media (MB7H9; Difco, MI, USA) supplemented with albumin-dextrose-catalase (ADC) (MB7H9/ADC) with gentle shaking at 37°C. To determine the number of colony-forming units (CFU) in the bacterial suspensions, the culture samples were serially diluted and plated onto Middlebrook 7H10 agar plates supplemented with OADC (MB7H10/OADC). The plates were incubated for 3 weeks at 37°C, and the CFU were counted.

To obtain whole cell lysates of BCG (BCG-WCL) samples of the BCG suspensions containing 1.0 × 10^8 bacilli/mL were centrifuged at 4000 g at 4°C for 50 min. The sediments were collected and homogenized in lysis buffer (Tris-HCl 10 mM, NaCl 10 mM, 0.5% SDS, 25 mM EDTA) at pH 7.8. The bacilli were submitted to 12 alternating cycles of freezing in liquid N2 and thawing in a water bath at 37°C. The disrupted bacilli were then sonicated (50 Hz for 30 min), the extracts were centrifuged at 4000 g for 15 min, and the protein concentrations were determined.

2.3. Mouse Immunizations

For the immunizations, the BCG strain was grown in MB7H9 for 2 weeks until the late stationary phase (O.D._600 nm = 1.0). The bacteria were collected by centrifugation (4000 rpm at 4°C for 10 min), washed twice and suspended in PBS to 10^6 CFU/mL. Each mouse was intradermally inoculated eight times with 2 × 10^7 bacilli, 0.1 mg in 100 µL PBS, with an interval of 3 weeks between the first and the second inoculations followed by 2 week intervals between the next six immunizations. Control groups of mice received 100 µL PBS. Before the immunizations, pre-immune serum was collected. The presence of anti-BCG antibodies in the sera from immunized and control mice was analyzed by ELISA, and the BCG protein bands recognized by the serum were visualized by Western blotting.

2.4. Purification of Anti-BCG Antibodies

Blood was collected by venomous puncture from
anesthetized mice. The blood samples were allowed clotting at room temperature for 1.0 h and kept for 16 h at 4°C. The clotted blood was centrifuged at 2000 rpm for 20 min, the serum was withdrawn, diluted 1/3 in pH 4.0, 60 mM acetate buffer, and the IgG immunoglobulins were purified by negative precipitation with caprilic acid [19]. The pH of the resulting purified IgG was adjusted to pH 7.2, by adding 3M Tris solution. The IgG immunoglobulins, were further purified by adding ammonium sulfate, at 4°C - 8°C, until 45% saturation. The precipitated IgG was separated by centrifugation at 5000 g, for 15 min, dissolved in 0.15 M NaCl and dialyzed against carbonate buffer (0.015 M Na2CO3, 0.035 M NaHCO3), pH 9.7. After washing, the whole BCG extract diluted in carbonate buffer (0.015 M Na2CO3, 0.035 M NaHCO3), pH 9.7. After washing, the plates were blocked with 100 µL of 10% FBS in PBST and the wells were washed five times with PBST, and 50 µL of peroxidase-labeled isotype-specific secondary detection antibodies (rabbit anti-mouse IgG (total) or anti-mouse IgG1, IgG2a, IgG2b, IgG3, IgA and IgM; Santa Cruz Biotechnology Inc., USA) diluted 1:2000 were added. The plates were incubated for an additional 1 h at 37°C and washed. The peroxidase activity was measured using o-phenylenediamine (OPD) and hydrogen peroxide as a substrate by spectrophotometry at the wavelength of 492 nm. The resulting OD was plotted against the test sera or IgG dilutions. The antibody titers were referred to either as the “end point” or the U-ELISA amount. The “end point” was generated by the first OD attaining maximal values. The U-ELISA was calculated by using the closest OD to 0.2. One U-ELISA, defined arbitrarily as the smallest amount of an antibody to yield an OD of 0.2 under the ELISA conditions, was calculated according the following equation [20]:

\[
U\text{-ELISA} = \left(\frac{(DO - 0.200)(Dil.)}{DO}\right) + Dil. \times 10,
\]

where DO was obtained from the equine antibody dilution closest to 0.200; Dil., the IgG (T) dilution corresponding to DO, and 10 refers to per mL of undiluted antibody. In this work, the antibody quantity in micrograms was used as a substitute for the antibody dilution to calculate the U-ELISA. All the values were recorded after the appropriate blank correction. Similar assays were used to quantify the anti-BCG antibodies in the purified total immunoglobulin IgG and the IgG1, IgG2a, IgG2b, IgG3, IgA and IgM antibody subclasses.

2.6. Detection of BCG Protein Bands by Western Blotting

Whole-cell protein extracts from the BCG strain were resolved by SDS-PAGE (15%) and subsequently transferred onto a nitrocellulose membrane. After transfer, the nitrocellulose sheets were probed with the mouse IgG immunoglobulins isolated from the sera of the BCG-immunized mice, followed by treatment with anti-mouse IgG conjugated to hors eradish peroxidase as the secondary antibody. The purified IgG prepared before immunization or from non-immunized mice were used as negative controls. The membranes were developed with a chemiluminescent kit (Millipore) and exposed on an ImageQuant LAS 4000 (GE, USA).

2.7. BCG Opsonization

One-hundred microliters of the bacterial suspension were incubated for 30 min at 37°C in 1.0 mL of phosphate-buffered saline (PBS) with or without the mouse anti-BCG total serum or IgG. One hundred microliters of PBS or serum obtained from non-immunized mice or collected just before immunization were used for opsonization as negative controls. The suspension of opsonized bacteria was centrifuged at 3000 rpm for 10 min at 4°C, and the supernatant was discarded. Immediately before the assay, the pellet was suspended in 900 µL of warm (37°C), colorless Hanks’ solution (Gibco) buffered to pH 7.4 with 10 mM HEPES, containing 10 - 5 M luminol (Sigma-Aldrich, St. Louis, MO). All opsonization tests were performed in triplicate.

2.8. Macrophage Culture, Infection and BCG Phagocytosis

Murine macrophage-like RAW 264.7 cells (ATCC) were expanded in 25 cm² or 75 cm² flasks (Corning) containing DMEM-F12 supplemented with 10% of BS, at 37°C for 5 days. For experiments, the cells (1 × 10⁶ cells/mL) were seeded onto 24-well plates. The macrophage monolayers were infected with BCG pre-treated with the murine anti-BCG IgG at a ratio 10/1 bacilli/macrophage. Samples of BCG incubated with non-immune IgG or PBS were used for infection of the control cultures (always running in parallel). The macrophage/bacteria interactions were stopped at 1 h, 3 h, 6 h, 9 h, 12 h, 24 h, 48 h and 72 h after infection. The cells were washed with PBS three times to remove any extracellular.
bacteria and examined for phagocytosis of mycobacteria by immunofluorescent microscopy.

2.9. Immunofluorescence Examination

The washed cells were fixed with 4% paraformaldehyde for 15 min and permeabilized with 0.5% Triton × 100 for 30 min at room temperature. After washing with PBS, the cells were incubated in DMEM-F12 supplemented with 10% FCS and treated with rabbit anti-mouse IgG-conjugated with phycoerythrin (1:1000 dilution) for 1.0 h at 4°C, in the dark. The cells were covered with ProLong Gold plus DAPI (Invitrogen), and the slide borders were sealed before examination by fluorescence microscopy.

2.10. Quantification of Intracellular BCG after Phagocytosis by Macrophages

Macrophages were plated onto 96-well plates (4 × 10^5 cells/mL). After adhesion, the cells were infected with BCG bacilli opsonized with immune mouse IgG (100 µg/ml) or non-opsonized bacilli at a bacillus:macrophage ratio of 10:1 for 15 min at 37°C under constant stirring. The plates were transferred to a CO2 incubator and incubated for different periods of times (1 h, 3 h, 6 h, 9 h, 12 h, 24 h, 48 h, 72 h). At the indicated intervals, the non-phagocytosed BCG bacilli were removed by three washes with sterile PBS. The bacterial uptake and survival in macrophages was quantified by the CFU test. The infected macrophages were disrupted with 0.1% saponin, homogenized and serially diluted. After stirring, 60 µL samples were plated onto Petri dishes containing 7H10 Middlebrook solid medium. The Petri dishes were incubated for 30 days at 37°C, and the numbers of CFU were counted.

2.11. Quantification of NO Production by Infected Macrophages

The production of NO was indirectly quantified by measuring the amount of NO_2− in the macrophage supernatants. Fifty microliter samples of the supernatants from each macrophage culture infected with BCG for 24 h were plated onto 96 wells plates containing 50 µl Griess reagent (1.0% sulfanilamide, 0.1% N-naftilenediamine and 2.5% phosphoric acid). After 10 min of incubation at room temperature, the O.D. was read at 540 nm. The amounts of released NO were calculated by projecting the O.D. obtained onto a previously constructed standard curve based on eight dilutions starting with 200 µM NO_2−.

2.12. Statistical analysis

The results were submitted to two-way ANOVA tests using GraphPad Prism 5.0 software (GraphPad, San Diego, CA). A p-value of <0.05 was considered indicative of a significant difference.

3. RESULTS

3.1. Production and Purification of the Anti-BCG Antibodies

Specific anti-BCG antibodies were quantified in the serum of immunized mice by ELISA assay using either rabbit anti-mouse total Immunoglobulin antibody or antibodies specific to different Ig classes (IgG, IgM, IgA, or the subclasses, IgG_1, IgG_2a, IgG_2b and IgG_3). Anti-BCG antibodies began to appear after the second immunization (n = 5; median = 8.8152 ± SD = 0.110642) compared with the sera collected before immunization (n = 5; median = 0.20712 ± SD = 0.079421). The titers increased thereafter, reaching 2.54-fold higher at the 8th booster inoculation with the antigen (n = 5; median = 2.0712 ± 0.116721) (Figure 1). The anti-BCG antibodies were differently distributed among the IgG subclasses and classes as follows: IgG_1 (n = 3; median = 1.9983 ± SD = 0.168197); IgG_2a (n = 3; median = 1.467667 ± 0, 061744); IgG_2b (n = 3; median = 1.470333 ± 0.050003);

Figure 1. Anti-BCG antibodies were detected in serum of mice immunized with whole BCG. ELISA plate wells were precoated with BCG whole extracts and incubated for 1 h at 37°C. After washes and incubation with peroxidase labeled rabbit anti total mouse IgG, the peroxidase activity was measured using o-phenylenediamine (OPD) and hydrogen peroxide as a substrate by spectrophotometry at the wavelength of 492 nm. Anti-BCG antibodies started to appear after the second immunization (n = 5; median = 8.8152 ± SD = 0.110642) as compared with sera collected before immunization, and difference between the titers observed after the 2nd and 8th immunizations are statistically significant (p < 0.001). The assays were performed in triplicates and the results were analyzed by the one way ANOVA test, p < 0.001.
IgG3 (n = 3; median = 0.095 ± 0.001732); IgM (n = 3; median = 1.30333 ± 0.060069); and IgA (n = 3; median = 1.141 ± 0.179084. Anti-BCG antibodies were not detected in the IgG3 fraction or in the serum collected before immunization (Figure 2).

The immunoglobulins were purified by negative caprylic acid precipitation of pooled sera collected from mice immunized with BCG one week after the last booster (8th inoculation). The precipitated non-immunoglobulin proteins were removed by centrifugation, and the immunoglobulin-rich supernatant was dialyzed against PBS. The resulting purified immunoglobulin preparation containing 1.0 mg of total protein/mL upon analysis by PAGE under reducing conditions manifested the typical immunoglobulin H and L chains as the major protein bands (Figure 3).

The anti-BCG antibodies present in these immunoglobulin preparations were quantified by ELISA. Various concentrations of the purified immunoglobulin preparations (0.78 µg to 100 µg) were added to wells pre-coated with 2 µg of whole BCG extract as the primary antibody, and rabbit anti-mouse IgG was added as secondary antibody. Anti-BCG antibodies were significantly detected in a dose-dependent manner even at the lowest doses (Figure 4). As indicated above, the O.D. at 492 nm corresponding to 0.2 was used to calculate the U-ELISA. This immunoglobulin preparation contained 1794.8 U-ELISA/mg. This value corresponds to the specific activity of the purified mouse anti-BCG immunoglobulin preparation. The purified immunoglobulin preparation was capable of recognizing protein bands with different molecular masses ranging from 7 kDa to 122 kDa in whole BCG extracts, as revealed by Western blot analysis. Compatible with the different anti-BCG antibody titers present within the IgG subclasses, 10 protein bands were clearly detected by IgG1 and 4 by IgG2a and IgG2b, but none were detected with IgG3 (Figure 5).
3.2. BCG Opsonization by Anti-BCG Antibodies

Suspensions of BCG (4 × 10⁶ bacilli per sample) were prepared as indicated in the Materials and Methods, washed with ice-cold PBS and pretreated with the indicated amounts of purified immunoglobulins for 15 min at 37°C. After incubation, the bacilli were washed and used for infection of RAW 264.7 cells at a bacteria/macrophage ratio of 10/1. The uptake of the opsonized BCG was examined by immunofluorescence and phase contrast microscopic analyses of the infected macrophages. The results presented in Figures 6(A)-(C) demonstrate the presence of opsonized BCG bacilli inside macrophages. The left panels illustrate stained opsonized bacilli (red) within macrophages (blue nuclei), while the right panels show that the BCG bacilli were also visible inside macrophages by light microscopic examination. The non-opsonized bacilli were detected in macrophages under light microscopy but not by the immunofluorescence test. Figure 6(D) shows macrophage not incubated with BCG.

3.3. Anti-BCG IgG Modulates Bacilli Phagocytosis by Macrophages

To determine the effect of the BCG pretreatment with anti-BCG antibodies on phagocytosis, the Raw 264.7 cell monolayers (1 × 10⁵ cells) were infected with opsonized bacilli by anti-BCG antibodies. Raw 264.7 cell monolayers were infected for 3.0 h at 37°C with viable BCG bacilli, pretreated (A), (B) or not treated (C) with mice anti-BCG antibodies. The bacteria/macrophage ratio was 10/1. After incubation, the cells were washed to remove non-phagocytosed bacilli. The cells were fixed, permeabilized, and treated with rabbit anti-mouse IgG-conjugated with phycoerythrin and stained with DNA fluorochrome, DAPI. The resulted cell monolayers were examined using immunofluorescent (A)-(D) or light (E)-(H) microscopy. In Left panel, only the opsonized BCG bacilli (A), (B), but not non-opsonized bacilli (C) were detected inside macrophages with nuclei stained in blue. The opsonized bacilli stained in red are pointed by arrows. In Right panel, the opsonized and non-opsonized intracellular bacilli are pointed by arrows (A)-(D), negative controls; macrophages incubated without BCG bacilli or non-opsonized BCG (BCG/macrophage ratio of 10/1) at 37°C. After different incubation periods, the cell monolayers were washed three times with PBS to re-
move outside bacilli. The macrophages were disrupted with 0.1% saponin, and 10 µL of the resultant cellular lysate was plated onto Petri dishes containing solid 7H10 culture medium. Next, the plates were incubated for 30 days at 37°C, and the numbers of CFU were counted. Figure 7(A) shows that the numbers of BCG bacilli recovered from macrophages infected with the opsonized BCG were significantly higher within the first 12 h of incubation compared with the cells infected with non-opsonized bacilli. The opsonized BCG/non-opsonized BCG phagocytosis ratio, which are greater than 1 until 12 h of incubation, was inverted after 48 h of incubation (Figure 7(B)). These results demonstrate that the numbers of intracellular bacteria began to decrease after 24 h of incubation, suggesting the activation of putative bacterial killing mechanisms in the host macrophages. The observed reduction was more pronounced in the cells infected with opsonized bacilli, suggesting that the bactericidal mechanisms were more potent in these cells.

The putative bacterial killing mechanisms employed by macrophages that phagocytosed the opsonized BCG bacilli were further investigated. To this end, we analyzed the induction of nitric oxide production in macrophages infected with opsonized and non-opsonized BCG. Supernatants from the macrophage cultures were collected, and the NO production was indirectly quantified by determining the amounts of NO2−-using Griess reagent (Figure 8(A)) demonstrates that after 24 h of incubation, NO production increased to reach a maximum at 72 h of incubation. The production of NO by macrophages phagocytosing opsonized BCG was at least 2-fold higher compared with the non-opsonized bacilli. The ratios of NO production induced in macrophages by the opsonized BCG versus non-opsonized BCG (ratio opsonized/non-opsonized BCG) began to increase after 9 h of incubation (ratio of 16.6), progressively increasing to reach the highest value at 24 h (ratio of 6.7), then decreasing until 48 h (ratio of 1.68) (Figure 8(B)).

4. DISCUSSION

The attenuated Mycobacterium bovis (BCG) strain, while retaining the typical characteristics of the species, does not express the relevant components responsible for the original virulence [21]. BCG is an intracellular bacterium that resides within mononuclear phagocytes and preferentially induces a cellular immune response [22]. Accordingly, previous research has largely focused on the activation of T cell subsets and the resulting cytokine profiles [23-25]. Except for some ancillary approaches, the humoral immune response putatively activated by BCG has been neglected. The exceptions include an analysis of the effects of the BCG growth conditions [15] and different BCG preparations, such as oral BCG, on the induction of the humoral immune response [16]. The lack of virulence factors and the intrinsic antigenic and adjuvant properties of BCG have long been exploited to protect humans against some forms of tuberculosis (M. tuberculosis) and hanseniasis, either through subcutaneous or oral administration. The adjuvant properties of the vaccine qualify BCG to be a vector for virulence factors of other pathogens. Accordingly, reports about the generation of recombinant BCG expressing virulence factors from Bordetella pertussis or enteropathogenic Escherichia coli have been recently published [16,17]. Detailed investigations of the capacity of BCG to induce humoral immunity are lacking.

The present study was designed to generate data about the pattern of the humoral immune response elicited by BCG in mice, specifically regarding the major qualities of the resulting anti-BCG antibodies. The strategy involved a longer immunization schedule of 7 repetitive boosters following the primary immunization. The implicit objective was to provide ample access of the antigens prevalent in BCG to the different antigen-presenting...
Cells (APC), dendritic cells, macrophages and B cells, leading to the subsequent activation of naïve T cells, effector T cells and B cells [26] of anti-BCG antibodies were obtained, as assessed by ELISA (Figure 1). These data indicate that the employed immunization strategy permitted successful MHC class II-dependent presentation of BCG antigens processed by APCs. The resulting consequences of the binding and cross-linking of immunoglobulin receptors, leading to B cell activation and the upregulation of costimulatory molecules including B7-1, B7-2, and CD40, were apparently achieved. The observed variety of Ig classes and subclasses produced in response to repeated BCG inoculations demonstrated that the expansion and differentiation of B cell precursors into Ab-secreting plasma cells, immunoglobulin H-chain class and subclasses switching [27] and antibody production occurred (although not investigated in this work). This idea is based on extensively published related data. These events led to anti-BCG antibody production, resulting in high antibody titers and unequal class and subclass distribution (Figure 2). The lowest levels of anti-BCG antibodies in the IgG3 subclass may be explained, at least in part, by the known naturally low prevalence of this IgG subclass in the mouse molecular immunoglobulin gene organization [28]. The anti-BCG antibodies of various immunized mouse immunoglobulin classes and subclasses recognized several distinct protein bands from the BCG extracts (Figure 5). Therefore, the protective possibility of the BCG antibodies can at least be double expanded.

Accumulating evidence has implicated both IgG and IgA antibodies in the host defense against bacterial infections [29]. IgG first interacts with the pathogenic cells or with their soluble components through the Fab-hypervariable regions. The resulting immune complexes formed after the reaction of specific IgGs with the epitopes expressed by the pathogens can activate cell or humoral effector mechanisms.

The effector cell functions of macrophages, monocytes and polymorphonuclear cells are initiated by the interaction of the IgG Fc fragments with the Fc receptors, FγRI (CD64), FγRII (CD32), and FγRIII (CD16), expressed by these cells [30]. The activation of humoral effector mechanisms involves the complement system (CS) as one important source of both anaphylatoxins and C3b fragments. The CS can be activated by three pathways: the classical pathway (CP), the MBL pathway (MBLP) and the alternative pathway (AP). The CP is activated after the interaction of the Fc of immunocomplexes with the C1 component; the MBLP is initiated by the interaction between mannan-rich bacterial cell components and MAPs; and the AP is initiated after the spontaneous hydrolysis of C3-thioester bonds and the immediate ligation of exposed OH− or NH+ pathogen cell surface components [31]. The activation of these pathways may result in the production of various CS fragments that can combine with specific effector cell-bound receptors, resulting in pathogen destruction. The pathogen killing can be executed by phagocytosis, antibody-dependent cell-mediated cytotoxicity, and respiratory...
burst [32].

In this work, we have demonstrated that the anti-BCG antibodies are potentially able to prepare the bacilli for phagocytosis and lysis upon combination with the FcRs of the phagocytes (Figures 6(A)-(D)). The internalization of the opsonized BCG by macrophages during the first hours after infection was significantly higher compared with the non-opsonized BCG (Figure 7(A)). Once inside the phagosomes, the presence of Ig on the BCG cell surface accelerated the bacterial destruction (Figure 7(B)). The observed bactericidal effect was associated with a significant increase in nitric oxide (NO) production (Figure 8(A)). This finding suggests an increase in the production of inducible nitric oxide synthase indicated by a clear inversion of the amounts of BCG recovery from the infected macrophages which was increasing higher until 24h of the macrophage with opsonized BCG (opsonized/non-opsonized BCG ratio of 36.7), then decreasing until 48 h (ratio of 1.68) (Figure 8(B)). The production of NO, results from the metabolism of arginine to citrulline, and a number of number of enzymes, including nucleoside reductase, resulting in impaired bacterial growth [33-36].

Pathogen-associated patterns (PAMPs) can be recognized extracellularly by germline-encoded pattern-recognition receptors (PRRs) as Toll-like receptors (TLRs) [37], and intracellularly by the cytoplasmic nucleic acid receptors RIG-1 and Mda5 [38,39]. Once inside, cells antibody-coated particles are bound to cytosolic IgG receptor, tripartite motif-containing 21 (TRIM21) [40] via its carboxy (C) terminal PYRSPRY domain with highest binding affinity [41]. Antibody-coated virus once into the cytoplasm interacts with TRIM21 are degraded by a clear inversion of the amounts of BCG recovery from the infected macrophages which was increasing higher until 24h of the macrophage with opsonized BCG (opsonized/non-opsonized BCG ratio of 36.7), then decreasing until 48 h (ratio of 1.68) (Figure 8(B)). The production of NO, results from the metabolism of arginine to citrulline, and a number of enzymes, including nucleoside reductase, resulting in impaired bacterial growth [33-36].

The work was supported with research funds from FAPERJ—Programa-Cientistas de Nosso Estado (Proc. No.: E-26/100.628/200); FAPESP, (Proc. No.: 09/52804-0); CNPq, Bolsa de Produtividade (WDS), Nível 1A, (Proc. No.: 301836/2005-1). Angela Alice Amadeu and Estevam José Baldon for the kind assistance in the manuscript organization.

REFERENCES


[33] Shiloh, M.U., MacMicking, J.D., Nicholson, S., Brause,


