Maternal Modulation of Neonatal Immune System Development

OMAR R. FAGOAGA and SANDRA L. NEHLEN-CANNARELLA*

Department of Pathology, Immunology Center, Loma Linda University School of Medicine and Medical Center, 11234 Anderson St. Room 2578, Loma Linda, CA 92354-2870 USA

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Changes in programming of neonatal immune development were effected through maternal immune modulation (Leishmania major inoculation). In progeny of these dams, immune profiles in both blood and spleen were changed throughout the neonatal period and were pronounced after weaning. White blood cell (WBC) and lymphocyte counts in blood of 45-day-old progeny were two-fold less than control animals. In blood, proportions of B cells were greater, while T helpers, Tc/s and NK cells were less than in controls. In contrast, proportions of splenic B and NK cells were greater than controls. But, proportions of all T and Tc/s cells on d20 and 45 were lower than controls. In blood, absolute numbers of all T, Th naive and Th memory cells were lower than in controls. In contrast, in the spleen, numbers of NK, T and Th naive and memory cells were up to 200% greater than in control pups. Cytokine responses of splenic lymphocytes stimulated through CD3 ligation revealed no difference in IL-4 production. In contrast, IL-2 and IFNγ were lower on d45 and 5, respectively, in the experimental compared to control mice. These data support the hypothesis that maternal immune events during gestation can modulate the pattern of immune development in offspring.

INTRODUCTION

Challenges of the maternal immune system during, and even before, pregnancy have been shown to be capable of modulating neonatal immune responses. Recent data suggest that this modulation is mediated by T helper cells and, depending on the antigen, may be expressed as a reduction in IgM and IgG responses to specific immunization in the 6-week-old pup or as an increase in IgM levels in fetal circulation (Fujii and Yamaguchi, 1992). The latter suggests an advancement in isotype switching in the neonate. Perhaps, this temporal advancement is the result of maternal factors crossing the placenta. Furthermore, not only humoral but also cell-mediated immunity has been modulated (Field and Caspary, 1971; Cramer et al., 1974; Russell, 1975). These investigators reported neonatal immunity to antigens recognized by the maternal immune system before conception or during pregnancy and proposed such explanations as transplacental passage of antigen, transfer factor, antibodies and sensitized maternal cells leading to active sensitization of the neonate.

The mechanism for fetal sensitization has been proposed to be a result of either maternal cell passage, vertical antigen transmission or maternal transfer of soluble factors. Cell traffic across the placenta has been well documented in humans (Desai and Greger, 1963; Russell, 1975; Pollack et al., 1982) and in the mice (Beer et al., 1977; Piotrowski and Croy, 1996). Maternal lymphoid cells naturally enter the allogeneic fetus and take up residency in liver, spleen and bone marrow of wild type mice (Piotrowski and Croy, 1996). Passage of maternal cells into the fetus could explain the observed B cell unresponsiveness to non-inherited maternal HLA specificities documented in adults that are otherwise highly reactive to alloantigens (Claas et al., 1988). Findings of these studies indicate that maternal cells, transiting the uteroplacental interface, could directly convey information to the fetus about pathogens in the environment.

Vertical transmission of antigens, however, appears to be a viable alternative explanation for fetal sensitization to maternal antigens (Stastny, 1965; Horton and Oppenheim, 1976). Still others (Barnetson et al., 1976) have proposed the presence of a soluble lymphocyte factor by demonstration of sensitization to Mycobacterium leprae in newborn of infected mothers. On the other hand, maternally derived cytokines are potential candidates for modulating fetal and, subsequently, neonatal immunity.

*Corresponding author. Tel.: +1-313-966-0936. Fax: +1-313-966-0934. E-mail: snehlsen@dmc.org
The placental barrier to cytokines appears to be selective. Granulocyte colony-stimulating factor readily passes the uteroplacental barrier (Medlock et al., 1993) while others such as IL-8 (Reisenberger et al., 1996) and erythropoietin (Koury et al., 1988) do not. Although effects of transplacental passage of cytokines on fetal immune function are clearly possible, it is known that cytokines do not convey antigen-specific information that could direct antigen-specific perinatal host immunity. However, their transfer across the placenta raises the possibility that the antigen-specific perinatal host immunity. However, their function are clearly possible, it is known that cytokines do transplacental passage of cytokines on fetal immune development.

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**RESULTS**

Pregnant (gestational d13) CD-1 mice were inoculated with *L. major* extract (intraperitoneal). This immunization, designed to induce maternal inflammatory (Th1) immune responses, was tested by challenging maternal spleen lymphocytes (collected on the day of delivery) and assessing IFNγ and IL-2 production. Based on historical reports, it was hypothesized that progeny of these immunized dams would be sensitized to *L. major* antigen and have an advanced repertoire of immune cell phenotypes and Th1 functional capacity.

After 8 days of *L. major* inoculation (175 µg/animal), both IL-2 and IL-4 concentrations in serum from pregnant and non-pregnant controls were undetectable, Table I; while IFNγ, although measurable, was not different between groups. IL-2 concentrations in cell culture supernatants from pregnant animals were significantly (t test, p < 0.05) increased in experimental compared to control animals. Similarly, IL-2 concentrations in non-pregnant animals were significantly higher (p < 0.05) than controls. In contrast to IL-2, IFNγ concentrations were not different between the cultures of controls and experimental animals. In culture supernatants, no IL-4 was detected in any group. This study confirmed that in vivo treatment with *L. major* extract could elicit a Th1 immune response (IL-2 and IFNγ) in both pregnant and non-pregnant CD-1 mice. It also confirmed that 16.6 µg *L. major* extract/well was sufficient antigen concentration for in vitro testing of sensitization.

**Ontogenic Development of Immunophenotypes in Progeny Born to Dams Inoculated with Leishmania major**

White blood cell (WBC) and total lymphocyte counts in blood of 45-day-old experimental progeny were lower than those in control animals. In blood, the WBC on d20 was 1.5-fold greater than control cohorts (two-way ANOVA, interaction between factors, p < 0.05). However, by d45, WBC of experimental progeny were significantly lower (2-fold) from those of control progeny (Student’s t-test, p < 0.05). Absolute numbers of lymphocytes in blood of experimental progeny were significantly lower only on postnatal d45 compared to controls (two-way ANOVA, interaction between factors, p < 0.05). In contrast to blood,
Proportions of immune cell phenotypes in the blood of pups at various ages delivered from dams treated during gestation with saline or *L. major* (175 μg/mouse i.p. in saline). Data points are mean ± SE of 5 samples/age, except 6 for d20 saline group; and 5 samples/age for *L. major*. Samples represent pools of randomized pups (d3-10, 5–14/Pool; d20, 2/Pool; d45, 1 pup). Statistical significance (p < 0.05; df = 1; F = 4.18 for CD19; and F = 4.13 for CD8) for treatment effects among groups was derived from two-way ANOVA. Post hoc analyses were performed with Student’s t-test to reveal differences among groups per age (*, p < 0.05). See “Statistical Analysis” section for further details. At some ages, symbols encompass area of SE bars.

In spleen, treatment effects were observed only in absolute numbers of lymphocytes; these were 1.3-fold higher on d45 than in control mice (Student’s t-test, p < 0.05). Thus, treatment effects were observed but they were not apparent until after weaning. Proportions of lymphocyte subsets in offspring born to experimental or control dams were then compared for treatment effects.

### Proportions of Immune Cell Phenotypes in Blood

In blood of experimental progeny, proportions of B cells were greater (Fig. 1, panel A) while T cell subsets and NK cells were less than those in control progeny (Fig. 1, panel B). T cell effects were reflected in decreased Tc/s cells (two-way ANOVA, main effects, treatment vs. control, p < 0.05), that were lowest on d45 (Student’s t-test, p < 0.05). Proportions of NK and naïve Th (CD62L<sup>low</sup>CD44<sup>low</sup>) in blood were significantly affected, changes were dependent on age (two-way ANOVA, interaction between factors (age vs. treatment), p < 0.05). NK proportions in experimental animals were lower on d3 and higher on d10 (Student’s t-test, p < 0.05), and tended (t-test, p = 0.07) to be higher on d45 when compared to control progeny of the same ages. In contrast, proportions of naïve Th were significantly lower on d4 (Student’s t-test, p < 0.05) and tended (p = 0.12) to be higher on d45. Thus, *L. major* inoculation during pregnancy had a modulating effect on the proportions of specific immune cell phenotypes in circulation.

### Proportions of Immune Cell Phenotypes in Spleen

Proportions of splenic phenotypes in experimental progeny were increased compared to control progeny (Fig. 2). Main treatment effects were observed across all ages (two-way ANOVA, main effects, treatment vs. control group, p < 0.05) for CD19, CD3, CD4 and CD8. Overall, B cells were higher and T cells lower in experimental pups. Proportions of T cells on d20 and Tc/s cells on d20 and 45 were significantly below normal (Student’s t-test, p < 0.05). NK cells were strongly affected by treatment, and were dependent on age (two-way ANOVA, interaction between factors, p < 0.05). On d10, the NK population was 1.7-fold higher than controls (Student’s t-test, p < 0.05). In summary, proportions of B cells in blood and spleen were higher, while T cells were lower in experimental progeny compared to control progeny.

### Numbers of Immune Cell Phenotypes in Blood

Absolute numbers of blood cells in experimental progeny were lower than control progeny, including T, naïve Th (CD62L<sup>low</sup>CD44<sup>low</sup>) and memory Th (CD62L<sup>low</sup> CD44<sup>high</sup>), (Fig. 3), and was the main treatment effect (two-way ANOVA, main effects, control vs. treatment group, p < 0.05). Furthermore, decreases in numbers of all subsets of T cells were dependent on age (two-way ANOVA, interaction between factors, p < 0.05). T cells and T cell subsets were 1.9- to 3.4-fold lower than controls on d45, but Tc/s rose 1.5-fold higher on d20 (Student’s t-test, p < 0.05). In general, *L. major* inoculation during gestation decreased the number of blood immune cell phenotypes in progeny, but most of the effect was not apparent until adulthood. The specific time when the effects were more noticeable post-weaning could not be determined because data between d20 and 45 was not collected.

### Numbers of Immune Cell Phenotypes in Spleen

The principal treatment effects observed in spleen cells were higher NK and naïve Th cells (Fig. 4; two-way ANOVA, main effects, control vs. experimental progeny, p < 0.05). Additional effects of treatment, significantly
dependent on age (two-way ANOVA, interaction between factors, \( p < 0.05 \)), were greater numbers of NK, T and Th naıve and memory cells than found in control pups. Absolute numbers of naıve and NK cells on d45 were significantly higher (Student’s \( t \)-test, \( p < 0.05 \)), 140 and 200%, respectively, than those in control mice. Thus \( L. major \) treatment during pregnancy had an up-regulating effect on populations of T, naıve and memory Th and NK cells in spleens of progeny and this effect was pronounced at post-weaning age. Missing data (no assessment between d20 and 45) prevents accurate determination of when these changes actually manifest.

**Ontogenic Development of Cytokine Response Capabilities of the Progeny Born to \( L. major \) Treated Dams**

Functional capabilities of splenic lymphocytes were tested as previously described (CD3-ligated mixed cell cultures and cytokine levels in supernatants). IL-4 production did not differ between \( L. major \)- and saline-treated groups. In contrast, Th1 cytokines decreased in response to CD3 ligation. IL-2 was significantly lower (\( p < 0.05 \)) on d45. The modulating effects of \( L. major \) inoculation on IFN\( \gamma \) production was associated with age (two-way ANOVA, interaction between factors, \( p < 0.05 \)). Post hoc comparisons with Student’s \( t \)-test revealed significantly (Fig. 5, Student’s \( t \)-test, \( p < 0.05 \)) lower IFN\( \gamma \) concentrations on d5. Thus, data of this study have documented that induction of immune reactivity in dams during gestation induces modulation in cytokine response capabilities of offspring.

**Evaluation of Sensitization to \( L. major \) in Progeny**

To determine whether progeny of inoculated dams had developed antigen-specific sensitization, spleen cells were challenged (72-h culture) with \( L. major \) extract (17 mg). Supernatants were then analyzed for cytokines (IL-4, IL-2 and IFN\( \gamma \)). No cytokines were detected in progeny from saline or \( L. major \) treated dams. These data indicate that the pups born to experimental dams may not have been sensitized to \( L. major \) in utero.

**DISCUSSION**

In this study, changes in programming of neonatal immune development were effected through maternal immune modulation (response to \( L. major \) inoculation). In progeny of these dams, maturational profiles of immune cells in both blood and spleen were changed throughout the neonatal period, and these changes were more pronounced...
after weaning. Blood leukocytes, lymphocytes and specifically T cells of experimental progeny were lower than in control progeny and this difference was noted to be marked on d45. In contrast, spleen lymphocytes, T cells, naïve and memory Th cells and NK cells of experimental progeny were higher on d45 compared to control progeny, suggesting an increased homing of circulating lymphocytes to splenic compartments. That effects of maternal immune experience during gestation are most apparent in progeny reaching adulthood, may be a result of yet further maternal-mediated immune regulation, perhaps through passive transfer of immunoglobulins during late gestation and via milk during the suckling period (Newman, 1995).

Some of these regulatory products in milk include antibodies, cells (lymphocytes, macrophages and neutrophils), lactoferrin, B12 binding protein, bifidus factor, fibronectin, gamma-interferon, hormones, growth factors and lysozyme.

Expectations were that this experiment would result in up-regulation of Th1 function; instead, it appears that Th2 functions may have been induced. Spleen cells of progeny had decreased production of IL-2 and IFN\(\gamma\) compared to normal controls. It is possible that a state of tolerance rather than inflammatory reactivity was induced in dams and pups, evidenced by the decrease in Th1 cytokine production in response to polyclonal stimulation. The decrease in Th1 cytokine production can be explained by taking into account three facts: soluble antigens primarily induce Th2 reactions (Zinkernagel and Kelly, 1997); inoculation in mid-gestation can lead to tolerance (Beer et al., 1977); and down-regulation of Th1-type cytokines is a consequence of tolerance induction (Chen and Field, 1995). If dams had been tolerized and this immune status passed to their progeny, a shift in cytokine profile away from Th1 would be expected. Cytokines production might have been modified by the abundant immune products that

**FIGURE 3** Absolute numbers of Th (CD4, Panel A), naïve Th (CD62\(^{lo}\)/CD44\(^{lo}\), Panel B) and memory Th (CD62L\(^{hi}\)/CD44\(^{hi}\), Panel C) cells in blood of pups of various ages delivered from dams treated during gestation with *L. major* (175 \(\mu\)g/mouse i.p.) or saline. Data points are mean ± SE of 5 samples/age, except 6 for d20 in saline group; and 5 sample/age in *L. major* group. Samples represent pools of randomized pups (d3-10, 5–14/pool; d20, 2/pool; d45, 1 pup). Statistical significance (\(p < 0.05\); df = 1; \(F = 4.9\) for Th; 2.4 for naïve and 5.3 for memory) for treatment effects among groups was derived from two-way ANOVA. Interaction between treatment and age suggest that effects are seen only on d45 (\(F\) statistic: naïve = 2.4 and memory = 5.3; df = 5, \(p < 0.05\)). Post hoc analyses with Student’s t-test (two-tail) revealed differences among groups per age (*, \(p < 0.05\); df = 8; t statistic of −3.35 (CD4), −3.64 (naïve), and −3.64 (memory)). See “Statistical analysis” section for further details. At some ages, symbols obscure SE bars.
are delivered through suckling (Newman, 1995), thus, explaining the observation of more significant differences being noted post-weaning. However, this conclusion cannot be validated with the current data and further experimentation is required to explore the influence of milk immune products acquired through suckling in this model.

The present study raises the possibility that maternal immune experiences may equip offspring in utero to defend against specific pathogens encountered outside the womb while preventing vertical transmission of infection. If this had been the case, sensitization of experimental offspring to \textit{L. major} antigen would have been induced. The memory recall experiments, performed by challenging spleen cells of experimental offspring with \textit{L. major} extract, produced no detectable cytokines after three days of culture. These results indicate lack of sensitization to \textit{L. major} in the offspring. Alternatively, absence of response may be due to challenging neonatal cells with too high antigen concentration; it has been reported by one investigative team that neonatal antigen presentation is 100-fold more efficient than adults (Van Tol \textit{et al.}, 1983; 1984). The dose of \textit{L. major} antigen used in the memory

![Figure 4](image-url) Natural killer (NK, Panel A) and naive Th cells (CD62LposCD44low, Panel B) in spleens of pups of various ages delivered from dams treated during gestation with saline or \textit{L. major} (175 mg/mouse i.p. in saline). Data points are mean ± SE of 5 samples/age, except 6 on d20 in saline group; and 5 sample/age for \textit{L. major}. Samples represent pools of randomized pups (d3-10, 5-14/pool; d20, 2/pool; d45, 1 pup). Statistical significance (p < 0.05; df = 5; F = 8.27 for NK and 16.2 for naive) for treatment effects among groups was derived from two-way ANOVA. Post hoc analysis with Student’s t-test (two-tail) to reveal differences among groups per age (*, p < 0.05; df = 8; t statistic of 2.99 (NK) and 5.4 (naive)). See “Statistical analysis” section for further details.

![Figure 5](image-url) IFN\(\gamma\) (top panel) and IL-2 (bottom panel) production (stimulated) by spleen cells of CD-1 outbred mice (male and female) delivered from dams treated during gestation with saline or \textit{L. major} (175 mg/mouse i.p in saline). Each culture contained \(1 \times 10^6\) CD3 + CD4 + splenocytes and hamster anti-mouse CD3 monoclonal antibody in complete media. Control cells were cultured in media only. Data are expressed as mean ± SE of samples (control: d5 = 4, d10 = 3, d20 = 10, d45 = 11; experimental: d5 = 7, d10 = 7, d20 = 3, d45 = 5). Samples in stimulated cultures were pools of randomized pups (d5 and 10, 10/pool; d20, 3/pool; d45, 1 pup). Statistical significance (p < 0.05; df = 5; F = 3.6) between groups was derived from two-way ANOVA. Post hoc analyses with Student’s t-test (two-tail) revealed differences among groups per age (*, p < 0.05; df = 9 (IFN\(\gamma\)) and 14 (IL-2); t statistic of -2.63 for IFN\(\gamma\) and -2.45 for IL-2). See “Statistical analysis” section for further details.
recall assays had been optimized with adult splenocytes, a deficiency in this study that needs further investigation. However, assuming that a prozone effect had not occurred, these results were unexpected since Herman et al. (1982) reported sensitization of offspring from Leishmania donovani-infected hamsters. Lack of sensitization in our experimental progeny may also be the result of using a non-pathogenic, purified extract administered i.p.; Herman et al. (1982) administered live amastigotes (active intracardiac infection) to their experimental animals.

Active infection is processed differently than an injection of soluble antigen from a killed parasite. The life cycle of this parasite is complex and during the infection process, promastigotes express abundant gp63 glycoprotein. On the other hand, amastigotes (from which the extract was made) express very low levels of this protein (Kurtzhals et al., 1994; Reiner and Locksley, 1995). The cytokine environment during induction of immune responses to gp63 determines the course of disease resulting from the type of immune responses induced (Kurtzhals et al., 1994; Reiner and Locksley, 1995). Humoral responses are primarily induced by this protein (Kurtzhals et al., 1994) in the early phase of infection and Kemp et al. (1994) showed that cytokine response in visceral Leishmaniasis is primarily a Th2 response, whereas cutaneous Leishmaniasis elicits predominantly Th1 response. This suggests that there are different cytokine environments in visceral vs. cutaneous sites, and that response is also dependent on route of administration (i.p. vs. subcutaneous).

The unexpected result of these experiments may be rooted not only in the type of antigen used to induce Th responses during pregnancy, but also the immunization program (dose, route and timing; Zinkernagel and Kelly, 1997). In this work, L. major extract was injected i.p. and this may be a factor for lack of sensitization in progeny, even though maternal spleens tested in vitro tended to a polarized Th1 response. Time of inoculation during pregnancy may be another factor; in a different model, tolerization resulted when dams were inoculated 5–7 days antepartum (Beer et al., 1977). In the current work, L. major extract was given 7 days antepartum, a time chosen as most likely time to avoid fetal loss due to release of Th1 cytokines (Krishnan et al., 1996). This single immunizing event may not have been sufficient to induce sensitization in the offspring, and instead may have induced tolerance. Sustained chronic immune responses to live L. donovani over the whole course of pregnancy could explain sensitizations realized by Herman et al. (1982).

In summary, the data indicate that at maturity, T cells in blood augmented their homing into lymphoid compartments as evidenced by significant increases in splenic NK and T cell numbers at postnatal d45 (Fig. 4). Furthermore, these changes in splenic cellularity produced significant decreases in IL-2 (d45) and IFNγ (d5) production by activated spleen cells compared to controls. Taken together, these data support the hypothesis that immune events experienced by the mother during gestation can modulate the pattern of immune development (cell phenotypes and cytokine response capabilities) in the offspring. Although significant effects in development were induced during gestation, a complete understanding of the mechanism must be ascertained before elective manipulation can be entertained. Timing and mode of delivering sensitizing or tolerizing immunogens must be defined, as well as dose and schedule of immunization. Of considerable importance is the need to determine what effect concurrent immune events would have on the intended process since regulatory function promulgated by APC is subject to prevailing signals dictated by local neuroendocrine immune products.

**MATERIALS AND METHODS**

CD-1 mice, a genetically wild-type strain were purchased from Charles River Laboratories (Wilmington, MA). The first pregnant CD-1 mice arrived in the vivarium on gestational d13. All mice were housed individually in small cages in the same room and maintained in a 12- and 12-hour light–dark cycle. Postpartum females, adult males and progeny provided the breeding nucleus needed to generate additional pups for later studies. A “trios” method of breeding was employed. Briefly, one male was mated to two females and housed together for 48 h. Females were then individually housed; a vaginal plug confirmed that copulation had occurred. The limited duration of conspecific exposure coordinated the timing of birth. Body weights of female breeders were measured daily to confirm the successful progress of pregnancy.

**Collection of Blood and Spleen**

Mice, 10 days of age or less were killed by decapitation and blood was collected into heparinized capillary tubes from cut surface of the body. For 20- and 45-days old groups, blood was collected from individual mice by cardiac puncture following pentobarbital anesthesia (40 mg/kg body weight); then mice were killed by decapitation. The spleen was extracted from the peritoneal cavity of each mouse under sterile procedures, weighed without mesenteric fat, and placed in sterile RPMI-1640 media (Fisher Scientific, Pittsburgh, PA) at room temperature. Lymphocytes were harvested within 30 min. Briefly, a spleen cell suspension was prepared by gently pressing tissue through a fine nylon mesh sieve and resuspending in complete media (RPMI-1640 containing prescreened 10% fetal calf serum and 1% penicillin/streptomycin). These reagents were purchased from Fisher Scientific, Pittsburgh, PA. Leukocytes in all blood and spleen cell suspensions were counted with a Unopette blood counting system (Becton Dickinson, Franklin Lakes, NJ) and a hemocytometer.
Pools were created by mixing blood or spleens of randomly-sorted pups from different litters of the same age. Each pool contained eight pups each for ages 0, 1 and 2 days. Although adequate sample volumes were obtained for immunophenotyping, it was impractical to perform experiments for study of cytokine production in mice before 2 days of age.

Immunophenotyping
Flow cytometric analysis was performed as previously described (Fagoaga et al., 2000). Briefly, single cell preparations from whole blood and spleen (1 × 10⁷ cells/tube) were stained with 10 μl fluorochrome-conjugated monoclonal antibodies (mAb). A mAb panel was constructed for two- (fluorosothiocyanate [FITC] and phycoerythrin [PE]) and three-color immunophenotyping (FITC, PE and Cy-Chrome). Two-color analysis was used to enumerate B vs. T cells (CD19*FITC/CD3*PE), T helper/inducer cells (CD4*FITC/CD3*PE), T cytotoxic/suppressor cells (CD8*FITC/CD3*PE) and natural killer cells (CD3*FITC/NK1.1*PE). Three-color analysis was used to assess naïve and memory T helper cells (CD62L*FITC/CD44*PE/CD4*Cy-Chrome). All rat anti-mouse mAb were obtained from PharMingen (San Diego, CA), except CD3*FITC (hamster anti-mouse antibody) obtained from Caltag Laboratories (San Francisco, CA). After incubation (15 min, 4°C), erythrocytes were lysed with 0.8% ammonium chloride solution and samples centrifuged at 300g for 5 min. Each supernatant was aspirated and cell pellets resuspended in 300 μl phosphate buffered saline (PBS).

Analytical controls included (a) unstained cells and (b) cells mixed with isotype antibody to evaluate degree of non-specific staining. Furthermore, an unlabeled, purified rat anti-mouse CD16/CD32 (FcγIII/II receptor; PharMingen, San Diego, CA) monoclonal antibody was used as an agent for blocking non-specific binding through the FcγIII/II receptor.

Proportions of lymphocyte populations (CD3, CD3CD4, CD3CD8, NK, CD19) were expressed as percentages normalized to lymphocyte purity (sum of lymphocyte subsets; CD19 + CD4 + CD8 + NK). This normalization excluded debris, other non-lymphoid cells (mainly monocytes) and Tγδ cells (CD3+CD4−CD8−). Normalization was not required for analyzing T helper cells for CD62L and CD44 expression because normalization excluded debris, other non-lymphoid cells (mainly monocytes) and Tγδ cells (CD3+CD4−CD8−). Absolute numbers of T helper cells analyzed for expression of CD62L and CD44 were similarly generated.

T Cell Stimulation Assay
A monoclonal antibody to CD3 was produced from cell culture of clone 145-2C11 (PharMingen, San Diego, CA) as previously described (Fagoaga and Nehlsen-Cannarella, 2000). Splenocytes were collected from spleens as described above. Subsequently, cell cultures were set up in 24-well plates (Fisher Scientific, Pittsburgh, PA). Cultures consisted of either unsorted single cell spleen suspensions (SSCS). To obtain a constant number of CD4 cells in each culture, absolute numbers of CD4 cells in unsorted cell preparations were determined by flow cytometry. An aliquot containing 1 × 10⁶ CD3 + CD4 + splenocytes was added to each well with 1 ml of optimally diluted CD3 monoclonal antibody. Each culture volume was brought up to 2 ml with complete media.

Cytokine Assays
Commercial cytokine ELISA kits from ENDOGEN (Woburn, MA) were used to assess IL-2, IL-4 and IFNγ production by murine splenocytes after CD3 ligation and after in vivo and in vitro L. major challenge. Assay procedures followed manufacturer’s instructions and as previously described (Fagoaga and Nehlsen-Cannarella, 2000).

Determination of In Vivo and In Vitro Leishmania major Dose
L. major extract of killed protozoa (sonicated) (Antibody Systems, Hurst, TX) was suspended in saline (1 mg/ml). A pilot study was conducted with pregnant and non-pregnant (n = 6) mice to determine optimal in vivo dose to induce a Th1 response (IL-2 and IFNγ production) Table I. Each animal received 175 μl (intraperitoneal injection) of either L. major extract (pregnant, n = 3; control, n = 3) or saline (pregnant, n = 3; control, n = 3). This dose was recommended by Antibody Systems for Th1 induction. Pregnant animals were inoculated once (gestational d8, 11 or 13); no further injections were given as a precaution to avoid induction of premature labor and delivery. Spleens and sera were collected from dams on day of parturition (gestational d20) and from non-pregnant controls 9 days post-inoculation.

Successful immunization of these animals was confirmed by assessing cytokine concentrations (IL-2, IFNγ, IL-4) in serum and in supernatants from spleen cells in culture after in vitro challenge with L. major (recall). Sera for these determinations were obtained by cardiac exsanguination under sodium pentobarbital anesthesia as previously described. For the immune memory recall study, spleens were harvested from exsanguinated mice and single cell suspensions prepared as described. Cells were incubated (1 × 10⁶ cells/well) in 96-well microculture plates (Fisher Scientific, Pittsburgh, PA) with 16.6 μg/well of L. major extract (“starting” concentration suggested by supplier) in complete media for 72 h in 5% CO₂ in humidified air at 37°C. At the end of incubation,
cell culture supernatants were collected and assayed for IL-2, IFNγ and IL-4.

Statistical Analysis

One-way ANOVA and Duncan’s post hoc test for multiple comparisons among different ages were calculated. If test for homoscedasticity was significant (greater than the critical value for F-max test for homogeneity of variances), data were log transformed. After transformation, if significant homogeneity of variances persisted, a Kruskal Wallis ANOVA with multiple comparisons was performed to assess differences among age groups. Two-way ANOVA was used to determine treatment effects of L. major extract inoculations. Furthermore, Duncan’s test for multiple comparisons and t-tests were used as post hoc tests to determine group differences. A value of p < 0.05 was considered significant.

References

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