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Impaired Germinal Center Responses and Suppression of Local IgG Production during Intracellular Bacterial Infection

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Germinal centers (GCs) are specialized microenvironments in secondary lymphoid organs that facilitate the development of high-affinity, isotype-switched Abs, and immunological memory; consequently, many infections require GC-derived IgG for pathogen clearance. Although Ehrlichia muris infection elicits a robust expansion of splenic, IgM-secreting plasmablasts, we detected only very low frequencies of isotype-switched IgG-secreting cells in mouse spleens, until at least 3 wk postinfection. Instead, Ag-specific IgG was produced in lymph nodes, where it required CD4 T cell help. Consistent with these findings, organized GCs and phenotypically defined splenic GC B cells were found in lymph nodes, but not spleens. Ehrlichial infection also inhibited spleen IgG responses against a coadministered T cell-dependent Ag, hapten 4-hydroxy-3-nitrophenyl acetyl (NP)-conjugated chicken γ globulin in alum. NP-specific B cells failed to undergo expansion and differentiation into GC B cells in the spleen, Ab titers were reduced, and splenic IgG production was inhibited nearly 10-fold when the Ag was administered during infection. Our data provide a mechanism whereby an intracellular bacterial infection can compromise local immunity to coinfecting pathogens or antigenic challenge. The Journal of Immunology, 2010, 184: 5085–5093.

The germinal center (GC) reaction is essential for the production for high-affinity, isotype-switched Abs during the adaptive immune response (1, 2). The formation of GCs requires the interaction of B and T cells with resident follicular dendritic cells in secondary lymphoid organs (3, 4). In GCs, Ag-specific B cells undergo rapid expansion, class-switch recombination, and affinity maturation. Although many GC B cells undergo apoptosis, a select few high-affinity B cells survive the T cell-dependent (TD) selection process and differentiate into isotype-switched B cell clones (2, 5, 6).

Class-switch recombination and affinity maturation typically require as long as 2 wk (5). Therefore, to prevent pathogen dissemination during acute infection, activated B cells proliferate and differentiate into Ab-secreting cells or plasmablasts in the red pulp of the spleen, adjacent to the T cell zone (7–9), or in the medullary cords of the lymph nodes (LNs) (10). Plasmablasts secrete primarily IgM against T cell-independent and TD Ags, and in this manner provide an early, typically short-lived, component of host defense.

Our previous study documented such an IgM-producing plasmablast response during Ehrlichia muris infection in the C57BL/6 mouse. E. muris is a tick-transmitted rickettsia that is closely related to Ehrlichia chaffeensis, the etiologic agent of human monocytotropic ehrlichiosis. The plasmablasts elicited during E. muris infection had the unusual characteristic of exhibiting low surface expression of CD11c, a marker more commonly associated with dendritic cells. The CD11c-expressing extralocular plasmablasts accounted for >15% of mononuclear splenocytes at peak infection and were responsible for the CD4 T cell-independent production of Ag-specific IgM (11) that is likely responsible for protective immunity. In the current study, we show that the plasmablast response that we described previously is associated with an impaired GC response in the spleen, but not the LNs. The impairment of the GC responses resulted in the inhibition of the production of isotype-switched Abs, against both ehrlichial Ags and coadministered irrelevant Ags. These studies provide a mechanism whereby ehrlichial infection can compromise local immunity to a coinfecting tick-borne pathogen.

Materials and Methods

Mice

The mice used in these studies were obtained from The Jackson Laboratories (Bar Harbor, ME) or were bred in the Animal Care Facility at the Wadsworth Center under microisolator conditions, in accordance with institutional guidelines for animal welfare. Mice were gender-matched for each experiment, and were 6–12 wk old. The studies were performed in accordance with institutional guidelines for animal welfare.

Infection and immunizations

Details regarding the bacterial strains and infection protocols have been described previously (12). Mice were infected i.p. at 6–12 wk old, with ~50,000 copies of E. muris. For the immunization studies, 4-hydroxy-3-nitrophenyl-acetyl-γ globulin conjugate (NP-CGG; Biosearch Technology, Novato, CA) was precipitated in alum (Imject; Sigma-Aldrich, St. Louis, MO) at a ratio of 1:5 (NP-CGG: alum) for 30 min. E. muris-infected and mock control mice were immunized i.p. or s.c. (in the left and right shoulders) with 100 μg NP-CGG/alum on the indicated day postinfection.

Abs and flow cytometry

The following Abs were used for flow cytometry: CD95-PE (clone FAS/APO-1), CD19-PerCP (clone ID3; both from BD Biosciences, Franklin Lakes, NJ), GL7-FITC (clone Ly77), and CD38-allophycocyanin (clone 90; eBioscience, San Diego, CA); 4-hydroxy-5-iodo-3-nitrophenylacetic acid

Abbreviations used in this paper: GC, germinal center; LN, lymph node; MZ, marginal zone; NF, 4-hydroxy-3-nitrophenyl acetyl; PNA, peanut agglutinin; OMP-19, outermembrane protein 19; RT, room temperature; TD, T cell-dependent.

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hydroxycinnamimide ester-haptened PE (NIP-PE) was used to identify NP-specific B cells. For flow cytometric analyses, spleen and LN cells were obtained by mechanical disruption in HBSS containing 2% FBS, and the tissues were disaggregated using a 70-μm pore size nylon strainer (Falcon; BD Biosciences, San Jose, CA). Erythrocytes were removed from the single-cell suspension of splenocytes by hypotonic lysis. For cell surface staining, single-cell suspensions (2 × 10^6 cells) were incubated with Fc blocking solution (1 μg/ml anti-CD16/CD32; FcγRIII/II; clone 2.4G2) in 10% normal goat serum/HBSS/0.1% sodium azide, prior to staining with the mAbs. The cells were incubated on ice for 20 min, washed twice, and analyzed without fixation. Negative controls were used to establish the flow cytometer voltage settings, and single-color positive controls were used to adjust compensation. Data from stained samples were acquired on a FACS Calibur with Cell Quest software (Becton Dickinson, Mountain View, CA), and were analyzed with FloJo Software (Tree Star, Ashland, OR).

ELISPOT and ELISA analyses

The detection of Ab-producing spleen and LN B cells was conducted using an ELISPOT assay. Nitrocellulose plates (Multiscreen-HA; Millipore, Bedford, MA) were coated overnight at 4°C with hapten 4-hydroxy-3-nitrophenyl acetyl (NP)-conjugated BSA (NP33-BSA; used at 5 μg/ml; Biosearch Technology, Novato, CA), or purified recombinant *E. muris* outer membrane protein-19 (OMP-19; at 10 μg/ml) (13). The assay plates were then incubated in blocking solution (IMDM supplemented with 10% FBS), for 2–3 h at 37°C. Cells were cultured in IMDM supplemented with 2 mM L-glutamine, 100 U/ml penicillin, 100 U/ml streptomycin, 50 μg/ml M-2-ME, and 10% FBS, and seeded in 96-well plates at a concentration of 1 × 10^5 cells/ml, in triplicate, in a volume of 100 μl; the cells were further diluted in the microwell plate using 2-fold doubling dilutions. After 18 h incubation at 37°C in 5% CO₂, bound IgM or IgG was detected using goat anti-mouse IgM or IgG conjugated to alkaline phosphatase (Southern Biotechnology Associates, Birmingham, AL); 5-bromo-4-chloro-3-indolylphosphate/nitro blue tetrazolium (Sigma-Aldrich) was used as the substrate. Spots were enumerated with a CTL immunospot S5 Core Analyzer, and the data were analyzed by CTL ImmunoSpot software (Cellular Technology, Shaker Heights, OH). IgM and IgG serum titers were determined by ELISA using purified recombinant *E. muris* OMP-19, as described previously (12), or NP33-BSA (BSA; 5 μg/ml).

Immunohistochemistry

Spleen and inguinal LN sections were prepared as previously described (11). The sections were stained in succession with rat anti-mouse Thy1.2 (BD Biosciences; overnight), biotinylated rabbit anti-rat IgG (Vector Laboratories; 2 h at RT), and streptavidin-Alexa Fluor-350 (Invitrogen, Carlsbad, CA; 30 min); they were washed extensively in PBS and then blocked with 10% FBS for 30 min at RT. For the detection of GCs, the sections were next incubated with biotin-conjugated rat anti-mouse B220 (BD Biosciences; 2 h at RT), and streptavidin-Alexa Fluor-647 (Invitrogen; 60 min). For the detection of GCs, the same sections were blocked with 2% BSA for 60 min at RT. Next, the sections were incubated with biotinylated peanut agglutinin (PNA) (Vector Laboratories; 1 h at RT) followed by streptavidin-Alexa Fluor-488 (Invitrogen; 30 min). When biotinylated Abs were used, a streptavidin-biotin blocking kit (Vector Laboratories) was used between each of the Ab incubations. The stained sections were mounted in anti-fading reagent (Slow Fade Gold; Invitrogen). Images were acquired using an epi-fluorescence microscope (Axioskop2; Zeiss, Peabody, MA) equipped with a Hamamatsu camera (Hamamatsu Photonic Sciences, Burlingame, CA; 60 min at room temperature [RT]), and images were mounted in anti-fading reagent (Slow Fade Gold; Invitrogen). Erythrocytes were removed from the single-cell suspension by hypotonic lysis. For cell surface staining, single-cell suspensions (10^6 cells) were incubated with Fc blocking solution (1 μg/ml anti-CD16/CD32; FcγRIII/II; clone 2.4G2) in 10% normal goat serum/HBSS/0.1% sodium azide, prior to staining with the mAbs. The cells were incubated with biotinylated peanut agglutinin (PNA) (Vector Laboratories; 1 h) and were analyzed with the mAbs. The sections were next stained with rat anti-mouse B220 (BD Biosciences; 2 h at RT), and streptavidin-Alexa Fluor-350 (Invitrogen, Carlsbad, CA; 30 min); they were washed extensively in PBS and then blocked with 10% FBS for 30 min at RT. For the detection of GCs, the sections were next incubated with biotin-conjugated rat anti-mouse B220 (BD Biosciences; 2 h at RT), and streptavidin-Alexa Fluor-647 (Invitrogen; 60 min). For the detection of GCs, the same sections were blocked with 2% BSA for 60 min at RT. Next, the sections were incubated with biotinylated peanut agglutinin (PNA) (Vector Laboratories; 1 h at RT) followed by streptavidin-Alexa Fluor-488 (Invitrogen; 30 min). When biotinylated Abs were used, a streptavidin-biotin blocking kit (Vector Laboratories) was used between each of the Ab incubations. The stained sections were mounted in anti-fading reagent (Slow Fade Gold; Invitrogen). Images were acquired using an epi-fluorescence microscope (Axioskop2; Zeiss, Peabody, MA) equipped with a Hamamatsu camera (Hamamatsu Photonic Systems, Bridgewater, NJ), and were processed with OPENLAB software (Zeiss, Peabody, MA).

Statistical analysis

The Mann-Whitney *U* test was used to assess statistical significance with values of *p* < 0.05 considered significant.

Results

Impaired IgG production in the spleens of *E. muris*-infected mice

Our previous study demonstrated that *E. muris* infection generates a large population of CD11c-expressing plasmablasts that produced the majority of total IgM, and almost all of the Ag-specific IgM within the spleen (11). That study focused primarily on IgM-producing cells, because we were unable to detect Ag-specific IgG-producing cells in the spleen on day 10 postinfection. To address the IgG response in greater detail, in this study, we infected C57BL/6 mice i.p. with *E. muris* (5 × 10^4 cells) and then measured Ab titers against the immunodominant Ag, OMP-19. OMP-19–specific IgG was detected in the sera of *E. muris*-infected mice beginning on day 9 postinfection, although Ab titers at this time were relatively low (<100; Fig. 1A). By day 28 postinfection, there was a 10-fold increase in OMP-19 IgG titer; OMP-19–specific IgG of the subclasses IgG2b, IgG2c, and IgG3 were detected at titers greater than 200, but IgG1 was not detected (Fig. 1B). The absence of a detectable IgG1 response was unexpected, although other infections associated with strong Th1 responses also favored switching to the IgG2a, IgG2b, and IgG3 subclasses (14). These data revealed that IgG was detectable at a low titer in the serum until at least 4 wk postinfection.

Given that IgG production was significantly delayed relative to the IgM response, we next used ELISPOT analysis to address whether the frequencies and numbers of Ig-producing B cells in LNs and the spleen were affected. *E. muris* does not exhibit any apparent LN tropism, so the studies were performed using pooled cells from the inguinal, brachial, and axillary LNs. The frequency of spleen OMP-19–specific IgM-producing cells was as high as 4% of total B cells, representing ~3 × 10^6 cells on day 14 postinfection, similar to our previously reported data (Fig. 1C) (11). In contrast, OMP-19–specific IgG-producing cells were found at low frequencies in the spleen and were not observed in significant numbers until day 28 postinfection, when ~1.5 × 10^6 cells were
detected. In contrast to the spleen, the LNs had readily detectable IgG-producing B cells between days 10 and 15 postinfection; the cells were present in LNs at a 4-fold higher frequency (0.4% and 0.1% in the LN and spleen, respectively; Fig. 1C).

**CD4 T cells are required for Ag-specific IgG production in LNs**

The spleen is the largest secondary lymphoid organ (15), and some IgG-secreting cells were detected in this tissue early during infection (Fig. 1C). *E. muris* infection causes splenomegaly, which is most pronounced at day 17 postinfection, and is associated with a 4-fold increase in mononuclear cells (11, 16). Thus, although the spleen contained IgG-producing B cells, we hypothesized that the LNs were major secondary lymphoid organs responsible for the production of high-affinity, isotype-switched IgG prior to 3–4 wk postinfection. Because IgG responses typically require CD4 T cell help, we next addressed whether T cells were required. This was determined using MHC class II-deficient mice, which lack CD4 T cells, and TCRβ-deficient mice, which lack αβ T cells. In these studies, mice were infected with *E. muris*, and OMP-19–specific IgM and IgG serum titers were assessed on day 16 postinfection. In the MHC class II and TCRβ-deficient mice, the OMP-19 IgM titers were reduced by 2- and 3-fold, respectively, compared with C57BL/6 mock-infected mice, suggesting that the Ag-specific IgM response was partially dependent on classical T cell help (Fig. 2A). In contrast, Ag-specific IgG was detected in C57BL/6 mice, but was not detected the sera of either MHC class II-deficient or TCRβ-deficient mice (Fig. 2B). Data from ELISPOT analyses supported this conclusion: Ag-specific IgG-secreting cells were largely undetectable (<0.02% of LN B cells) in the spleens and LNs of either the MHC class II-deficient or TCRβ-deficient mice (Fig. 2C). Although MHC class II-deficient mice exhibited a 2-fold decrease in the frequency and number of LN OMP-19 IgM-secreting cells, these cells were undetectable in the LNs of TCRβ-deficient mice. The spleen was the primary source of Ag-specific IgM in both the MHC class II- and TCRβ-deficient mice, but total numbers were reduced by 2- and 3-fold, respectively. Thus, T cells likely provide important helper functions that promote the development of both IgG- and IgM-producing cells in the LNs.

**Impaired GC responses in the spleen, but not the lymph nodes, during *E. muris* infection**

To further address whether the LNs provide a better environment for the production of IgG than does the spleen, we next evaluated whether the development of GCs and GC-derived B cells differed between the two lymphoid tissues. To identify GC B cells, we first analyzed spleen and LN B cells for cell surface expression of the late activation marker GL7 (17, 18). GL7-positive B cells were detected on day 9 postinfection in the LNs, reached a maximum on day 14 postinfection, and remained at frequencies above those in mock-infected mice on day 9 postinfection, when a 2-fold increase in frequency was observed. In contrast, infected and mock-infected mice were detected in the LN and spleen, respectively; Fig. 1C).

**Infection inhibits the splenic IgG response to an irrelevant Ag**

Our findings thus far demonstrate that ehrlichial infection inhibits the IgG response in the spleen. To determine whether this phenomenon is limited to ehrlichial Ags, or whether the IgG response to other Ags is also impaired during infection, we immunized uninfected and *E. muris*-infected C57BL/6 mice with the TD Ag NP-CGG, in alum, at various time points after *E. muris* infection. Sera and spleens were harvested for analysis 12 d after NP-CGG immunization (8, 25, 26), and the production of NP-specific Abs was analyzed by ELISA and ELISPOT. Mice immunized with NP-CGG typically generated reciprocal Ab titers ranging from 2000–10,000 within 12 d of immunization. However, when immunization was performed after *E. muris* infection, NP-specific serum Ab titers were much reduced (Fig. 5A). The greatest suppression of NP-specific IgG was observed when the Ag was administered between 2 and 16 d postinfection, although a modest but nonsignificant inhibition was noted at day 23 postinfection. The reduction in NP-IgG titers was likely caused by the impaired production of IgG-secreting cells in the spleen, given that the frequency of splenic NP-specific IgG-secreting cells was reduced by ~10-fold when NP-CGG was administered on day 9 postinfection (Fig. 5B). When NP-CGG was administered at other time points relative to infection, the frequency of IgG-secreting cells was reduced by factors ranging 2–4-fold. Despite the decrease in frequency, the number of NP-specific IgG-secreting cells was significantly reduced only at days 9 and 16 postinfection, because of infection-induced splenomegaly (Fig. 5C). Thus, *E. muris* infection inhibited the production of splenic IgG against both ehrlichial and coadministered Ags.

In the studies described thus far, both the pathogen and the Ag were administered i.p. We next addressed whether the NP-IgG response was affected when Ag was injected s.c., distant from the peritoneal drainage path. Mice were infected via the peritoneum with *E. muris*, and on day 9 postinfection, infected and mock-infected mice were
B cell response (3–5%) in the spleen (28, 27). Because of the lack of the transgenic mice leads to the generation of a modest NP-specific response (27). Pairing of the transgenic H chain with the endogenous H chain genes, the (m+s)lg Tg mice are unable to secrete IgG; however, splenic architecture and the B cell compartment remain intact (28). Because NP-specific IgG serum titers were suppressed as early as day 2 postinfection, we chose this time point for immunization of *E. muris*-infected mice with NP-CGG. Spleens were harvested 12 d postimmunization (day 14 after *E. muris* infection), and NP-specific CD19-positive B cells were detected by their ability to bind the related hapten, NIP (29, 30). Consistent with the results of published studies, we observed an increase in the frequency and number of NP-specific B cells in (m+s)lg Tg mice immunized with NP-CGG in alum, relative to unimmunized mice of the same strain (Fig. 6A). In contrast, expansion of the NP-specific B cell population was not detected in the *E. muris*-infected NP-CGG-immunized mice. In the infected mice, the frequency of NP-specific B cells was reduced by ∼10-fold, relative to the frequencies detected in mock-infected, unimmunized mice (0.21 and 2.67%, respectively), and the number of NP-specific cells was roughly similar in both groups, indicating the lack of specific B cell expansion during infection (Fig. 6B, 6C). To determine whether the reduction of NP-specific response was due to the failure of the Ag-specific B cells to develop into GC B cells during infection, we analyzed NIP+ CD19+ cells in the spleen at least in part to the lack of proper GC formation. The inhibition was not limited to ehrlichial Ags; infection also suppressed local IgG production against an irrelevant, coadministered TD Ag. Although we cannot rule out the possibility of a defective extrafollicular response, our studies suggest that the impaired production of splenic IgG is due to an impeded production during infection, but *E. muris* infection reduced the frequency of GC B cells by ∼10-fold (Fig. 6D, 6E). Furthermore, the total numbers of NIP+ GC B cells in the immunized–infected mice were comparable to those in the mock-infected control mice (Fig. 6F).

**Discussion**

Our studies demonstrate that *E. muris* infection can suppress IgG production and inhibit the development of GC B cells within the spleen. Ag-specific IgG-secreting cells were largely absent in the spleen until 3–4 wk postinfection (i.e., 2 wk after IgG-secreting cells were first detected in the LNs). We attribute the suppression of IgG production in the spleen at least in part to the lack of proper GC formation. The inhibition was not limited to ehrlichial Ags; infection also suppressed local IgG production against an irrelevant, coadministered TD Ag. Although we cannot rule out the possibility of a defective extrafollicular response, our studies suggest that the impaired production of splenic IgG is due to an inhibition in the GC response, because we demonstrated a clear defect in GC formation. These data provide a possible mechanism whereby a bacterial infection can suppress an Ab response against a coinfecting agent. It is well established that tick-borne pathogens can be cotransmitted by the same tick or by different ticks feeding on the same host (31). *Babesia microti* and *Borrelia burgdorferi*, the respective agents of babesiosis and Lyme disease, exemplify coinfecting pathogens that are commonly transmitted with the ehrlichiae and/or other related rickettsiae (32, 33). Coinfection with *Anaplasma phagocytophilum*, the etiologic agent of human anaplasmosis, and *B. burgdorferi* was associated with increased spirochete burden and exacerbated the early onset of Lyme arthritis (34, 33). Coinfection was proposed to impair the activation of macrophages, resulting in an increase in pathogen burden (34). Our studies suggest the impaired production of splenic IgG as another mechanism that contributes to coinfection-associated immunosuppression.

The inhibition of IgG production during *E. muris* infection was associated with distinct humoral responses in various secondary lymphoid organs. Although splenic B cell function was impaired, B cell responses in LNs were largely intact. Well-developed GCs were observed in LNs by day 14 postinfection, and IgG production required T cell help, indicating that classical B cell GC responses were largely unaffected in LNs during *E. muris* infection. Thus,
the suppression that we describe is likely relevant for understanding impaired immunity to other blood-borne pathogens that gain access to the spleen. In cases in which tick-borne pathogens first access the tissue-draining LNs, early IgG production may be unaffected (35); however, once a pathogen disseminates via the blood stream, the suppression of IgG production in the spleen could affect host defense. For example, the humoral response is required for immunity against infection with *B. burgdorferi* (36–38), given that the depletion of marginal zone (MZ) B cells (which are found only in the spleen) was responsible for a reduction in *Borrelia*-specific IgM and IgG, increased spirochete burden in the spleen, and early onset of Lyme arthritis (39). Another study demonstrated that *B. burgdorferi* infection caused the expansion of the MZ B cell subset, suggesting the importance of these B cells in controlling bacteremia (37). Thus, the local IgG suppression that we have observed during infection by *E. muris*, if it indeed occurs during other ehrlichial and/or rickettsial infections, may contribute to the suppression of IgG responses against coinfecting tick-borne pathogens, such as *B. burgdorferi*.

Impaired GC formation and IgG responses also have been documented in other bacterial (40), viral (41, 42), and parasitic infections (43, 44). *Plasmodium berghei* infection caused disruption of splenic and LN GC architecture during the first 10 d of infection (43). Impairment of the GC response was also reported during MMTV infection, although virus-specific IgG2c titers remained largely unaffected (41). A 3–4-wk delay in splenic GC formation was observed during *Salmonella enterica* serovar *Typhimurium* infection (40). Similar to our findings, systemic IgG was present during acute *Salmonella* infection, but...
the production of high-affinity isotype-switched Abs did not occur until the splenomegaly had been at least partially re-solved. Several studies have also demonstrated that infections can impair the IgG response against irrelevant Ags (45–49). Although a number of mechanisms have been proposed, possible effects of infection on Ag-specific B cells and GC responses have, prior to this study, not been addressed. The findings that infections can impair IgG responses are broadly relevant to the issue of vaccination efficacy, especially in regions where pathogens are endemic (50).

Although GC B cells were detected at a low frequency in the spleen during early infection, appreciable numbers of GC B cells were detected in this tissue because of massive splenomegaly. Although it is possible that these spleen IgG-producing cells provide a significant portion of IgG produced during infection, our data suggest that the quality of the splenic IgG response is poor, given that distinct GC structures were detected only in the LNs and that PNA-binding B cells were found in low numbers and were randomly dispersed throughout the spleen. One prediction from our findings is that splenic B cells generated early during infection will fail to undergo normal TD processes, possibly including affinity maturation, although this has not yet been addressed.

We propose that the impairment of the GC responses is caused by a disruption of spleen architecture, and this loss of tissue integrity is what prevents the development of a GC response. Other studies from our laboratory have shown that the disruption of spleen architecture is temporally associated with extramedullary erythropoiesis and lymphopoiesis during acute E. muris infection (51). The latter events include the massive expansion of Ter119-positive prereticulocytes, as well as spleen extramedullary hematopoiesis. Such changes in structure and function could account for the impaired splenic IgG responses. Alternatively, suppressed T cell function, perhaps owing to impaired dendritic cell maturation, could be responsible; such suppression has been observed in other infection models (46, 48). Our data provide an argument against the possibility that the failed IgG responses are due to a paucity of Ag-specific CD4 T cells, given that we have observed an expansion of Ag-specific, IFNγ-secreting T cells in the spleen during E. muris infection (16). We propose instead that Ag-specific CD4 T cells are unable to form proper associations with

FIGURE 5. E. muris infection suppressed the IgG response to an irrelevant Ag. E. muris-infected mice and uninfected control C57BL/6 mice were immunized i.p. with NP-CGG in alum on the indicated days postinfection. To assess the NP-specific IgG response, serum and spleen tissue were harvested on day 12 postimmunization and were analyzed by ELISA (A) and ELISPOT analysis (B, C) in mice immunized with NP-CGG only (○) and in E. muris-infected, NP-CGG–immunized mice (●). The frequency (B) and number (C) of splenic NP-specific IgG-secreting cells was determined by ELISPOT analysis after subtracting background spots obtained using cells from mock-infected control mice. D, Day 9 E. muris-infected and mock-infected mice were immunized s.c. with NP-CGG in alum, and the NP-specific IgG response was analyzed at day 21 postinfection by ELISPOT analysis in the spleen (top row) and pooled brachial and axillary draining LNs (bottom row). The data are representative of three independent experiments in which three mice were analyzed at each time point. *p ≤ 0.05.
differentiating GC B cells, because of the loss of tissue integrity, as well as the extramedullary hematopoiesis that we have observed in the spleen.

Alteration of the inflammatory cytokine milieu may also contribute to IgG suppression. Increased production of the proinflammatory cytokine IL-12, after TD immunization of Fc receptor-associated γ-chain–deficient mice, was associated with an expansion of short-lived plasmablasts, but resulted in an impairment of GC B cell differentiation (52). Similarly, GC formation was impaired during Salmonella infection, until after the resolution of an extrafollicular plasmablast response (40). Although the role of IL-12 was not investigated in the latter study, Salmonella infection elicits a Th1 response and is associated with the production of proinflammatory cytokines (53). In our studies, IgG responses to NP-CGG immunization were most strongly affected when the Ag was administered at the peak of the plasmablast response. Both the proinflammatory response elicited during ehrlichial infection (54) and the disruption of spleen architecture probably favor the expansion of the CD11c-expressing plasmablasts and the subsequent suppression of GC production.

The origin of the CD11c<sup>+</sup> plasmablasts has not yet been resolved (11), but ongoing studies suggest that MZ B cells likely contribute to the expansion of this population (R. Racine and G.M. Winslow, unpublished data). Although MZ B cells provide an early source of IgM against blood-borne pathogens, by rapidly differentiating into plasmablasts (55), the B cell population can also participate in GC responses to produce isotype-switched Abs (55), the B cell population can contribute to the expansion of this population (R. Racine and G.M. Winslow, unpublished data). Although E. muris infection was associated with an expansion of IgM-positive B cells, the loss of CD21 expression might prevent the efficient capture and shuttling of Ag from the MZ to follicular dendritic cells, thus contributing to the localized suppression of splenic IgG. In contrast, the LNs, which lack MZ B cells, provide a more suitable environment for the formation of GCs and production of isotype-switched Abs.

Thus, E. muris infection causes, via several possible mechanisms, a localized suppression of the GC responses, and this suppression is likely responsible for the inhibition of class-switch recombination and the production of isotype-switched Abs. These finding are relevant to our understanding of how coinfections can exacerbate disease symptoms during a diverse range of microbial infections.

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**Disclosures**

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