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Vitamin D Inhibits the Occurrence of Experimental Cerebral Malaria in Mice by Suppressing the Host Inflammatory Response

Xiuyue He,* Juan Yan,† Xiaotong Zhu,‡ Qinghui Wang,§ Wei Pang,† Zanmei Qi,* Meilian Wang,* Enjie Luo,* Daniel M. Parker,† Margherita T. Cantorna,* Liwang Cui,‡ and Yaming Cao*

In animal models of experimental cerebral malaria (ECM), neuropathology is associated with an overwhelming inflammatory response and sequestration of leukocytes and parasite-infected RBCs in the brain. In this study, we explored the effect of vitamin D (VD; cholecalciferol) treatment on host immunity and outcome of ECM in C57BL/6 mice during Plasmodium berghei ANKA (PbA) infection. We observed that oral administration of VD both before and after PbA infection completely protected mice from ECM. VD administration significantly dampened the inducible systemic inflammatory responses with reduced circulating cytokines IFN-γ and TNF and decreased expression of these cytokines by the spleen cells. Meanwhile, VD also resulted in decreased expression of the chemokines CXCL9 and CXCL10 and cytoadhesin molecules (ICAM-1, VCAM-1, and CD36) in the brain, leading to reduced accumulation of pathogenic T cells in the brain and ultimately substantial improvement of the blood–brain barriers of PbA-infected mice. In addition, VD inhibited the differentiation, activation, and maturation of splenic dendritic cells. Meanwhile, regulatory T cells and IL-10 expression levels were upregulated upon VD treatment. These data collectively demonstrated the suppressive function of VD on host inflammatory responses, which provides significant survival benefits in the murine ECM model. The Journal of Immunology, 2014, 193: 1314–1323.

Malaria is widespread throughout the tropical and subtropical regions, causing >300 million acute illnesses and resulting in more than 600,000 deaths annually. Cerebral malaria (CM) is the most severe complication of Plasmodium falciparum infection and a major cause of death in children under the age of 5 y. The mechanisms leading to CM in humans is not well understood and appears to be multifactorial. Cytoadhesion of parasitized RBCs (pRBCs) to the brain endothelium is thought to cause mechanical obstruction of the brain microvessels leading to CM pathology (1). In addition, excessive inflammatory responses characterized by high levels of proinflammatory cytokines are also thought to contribute to CM (1). Inflammatory cytokines upregulate expression of the adhesion molecules such as ICAM-1 and VCAM-1 on brain endothelial cells, further enhancing cytoadherence and sequestration of pRBCs in the brain. A better understanding of the mechanisms of CM and identification of effective adjunct therapies of CM are of high priority.

Rodent malaria infections such as the Plasmodium berghei ANKA (PbA) infection in C57BL/6 mice have been used widely as animal CM models because they share several features with human CM (2–4). In the mouse CM models, Th1 responses play a critical role in CM pathogenesis. Th1 responses are characterized by the increased production of IFN-γ and decreased production of the Th2 cytokines such as IL-4. Appropriate induction of Th1 cytokines is needed for successful control of parasitemia and resolution of malaria infection (5, 6), whereas excessive levels of these cytokines are implicated in the pathogenesis of CM (7, 8). Thus, regulation of the magnitude and timing of the Th1 response is essential for producing optimized immune responses that inhibit the malaria parasites without causing immunopathology. Regulatory T cells (Tregs) are important player participating in the control of overwhelming responses to infections (9–12). In the mouse CM model of infection, Treg expansion inhibits the development of pathogenic Th1 cells and CM (13, 14).

Vitamin D (VD) is a fat-soluble vitamin that is either synthesized in the skin after exposure to solar UVB radiation or provided in the diet. In addition to its traditionally known roles in regulation of bone metabolism and calcium–phosphorus homeostasis, VD has been increasingly recognized to have prominent regulatory functions on both innate and adaptive immune systems (15). The active form of VD (1,25(OH)2D3, 1,25D3) primarily affects dendritic cell (DC) maturation and macrophage differentiation (16, 17) and inhibits the production of the cytokines IL-12 and IL-23. In addition, 1,25D3 inhibits the production of Th1 cytokines (IL-2 and IFN-γ) and Th17 cytokines (IL-17 and IL-21) but stimulates Th2 cytokine production.

Abbreviations used in this article: BBB, blood–brain barrier; CM, cerebral malaria; DC, dendritic cell; ECM, experimental cerebral malaria; mDC, myeloid DC; MHC II, MHC class II; PbA, Plasmodium berghei ANKA; pDC, plasmacytoid DC; p.i., postinfection; pRBC, parasitized RBC; Treg, regulatory T cell; VD, vitamin D; VDR, VD receptor.

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of DC activation, IL-10 production, and expansion of the Tregs. Moreover, 1,25D3 favors development of Tregs via modulation of DCs (19). Because many autoimmune diseases such as inflammatory bowel disease, multiple sclerosis, and arthritis are the result of overwhelming Th1 responses, 1,25D3 treatments suppressed Th1 responses and ameliorated Th1-mediated experimental autoimmunity (20). Paradoxically, even though VD inhibits Th1 and Th17 responses, a number of infectious diseases are not made more severe by treatments with active VD (21).

The immunoregulatory functions of VD especially its inhibitory effect on Th1 responses have prompted us to examine the role of VD in experimental CM (ECM). In P. falciparum malaria, plasma VD level did not vary during the course of infection and VD status was not associated with incident malaria (22, 23). In a rodent malaria model, oral VD treatment of mice for 2 wk prior to P.berghei infection has been reported to decrease parasite growth and expand the life span of infected mice (24). However, a recent study showed that three weekly i.p. injections of 0.5 μg/kg VD had no effect on susceptibility of wild-type mice to PbA infection (25). In this paper, we explored the effect of VD on ECM and showed that oral supplementation with VD protected mice from ECM. Oral VD administration before and after PbA infection completely prevented the occurrence of ECM. We show that the protective effect of VD was through the inhibition of a strong host proinflammatory (IFN-γ and TNF) response mediated directly and indirectly through reduction of DC activation, IL-10 production, and expansion of the Tregs.

Materials and Methods

Animals and experimental infection

Female C57BL/6 mice (6–8 wk old) were obtained from Beijing Animal Institute and maintained at the Animal Care Facilities of the China Medical University. Infections were initiated by i.p. injection of 1 × 10⁸ PbA pRBCs into each mouse. Parasitemia was monitored by counting the number of pRBCs per 1000 RBCs by light microscope examination of Giemsa-stained thin smears from tail blood. Mice were monitored daily for survival and neurologic signs of CM such as ataxia, paralysis, and coma. For the mortality and parasitemia experiment, 10 mice were used in each group, and the experiments were repeated for four times. All experiments were performed in compliance with local animal ethics committee requirements.

VD treatment of mice

VD (cholecalciferol) was obtained from Sigma-Aldrich (St. Louis, MO) and delivered in mice using two routes: injection and feeding. For injection, VD was dissolved in 100% ethanol and diluted to a working solution of 0.1 μg/ml in 0.02% Tween 80. Mice in the treatment group was injected with a 0.5 μg/kg dose of VD every other day, starting 3 d before the PbA infection (25), whereas mice in the control group received equal volumes of 0.02% Tween 80. For oral administration, VD was dissolved in soybean oil before use. For the dose range experiment, mice were orally administered a daily dosage of 0, 10, 50, and 250 μg/kg VD for 4 d after PbA infection. For subsequent experiments, mice were randomly divided into seven groups: naive mice (uninfected), PbA-infected VD supplemented before PbA infection (VD+PbA), two groups with VD supplemented after PbA infection (PbA+VD), and two uninfected but VD treated control groups. Mice in the VD+PbA group were orally administered a daily dosage of 50 μg/kg VD for four consecutive days before PbA infection, whereas the PbA+VD group mice were orally administered 50 μg/kg VD daily for four successive days at 2 d (PbA+VD2d) or 5 d (PbA+VD5d) after PbA infection. The two control groups received the same volume of soybean oil at the identical time points as the VD+PbA and PbA+VD groups. On day 5, blood was collected, and VD content was measured using the 25(OH)D-Vitamin D direct ELISA Kit (Immundiagnostik, Bensheim, Germany), according to the manufacturer’s instructions. Samples were measured in duplicate and averaged.

Histopathology and immunohistochemistry

When PbA-infected mice began showing neurologic symptoms on day 6, five mice from each of the experimental groups were selected for histological examination. Brains were removed, fixed in 4% paraformaldehyde for 24 h, and embedded in paraffin. Serial 4-μm-thick horizontal sections were made, stained with H&E, and examined for microvascular obstruction and leakage. To detect ICAM-1, VCAM-1, and CD36 along the endothelial lining, immunohistochemical staining was performed with specific rabbit polyclonal Abs against ICAM-1 (Abcam, Cambridge, MA), VCAM-1, and CD36 (Santa Cruz Biotechnology, Santa Cruz, CA) as previously described (26), which was followed by biotin-conjugated goat anti-rabbit IgG Ab. Finally, streptavidin-conjugated peroxidase was added, and color development was done using 3- amino-9-ethylcarbazole as the substrate. Finally, the sections were counterstained with hematoxylin, washed, and mounted. ICAM-1–, VCAM-1–, and CD36-positive vessels were visualized by microscopy at ×400 magnification. The number of positive vessels in 20 fields was counted for each mouse.

Blood–brain barrier integrity

To evaluate integrity of the blood–brain barrier (BBB), 200 μl 2% (w/v) Evans blue dye (Sigma–Aldrich) in PBS (pH 7) was injected i.v. into each mouse. One hour later, mice were euthanized and perfused with 20 ml PBS. Brains were isolated and incubated in 2 ml formamide for 48 h at 37°C. The amount of Evans blue in 100 μl of the brain tissue extracts was determined by measuring absorbance at 630 nm (27).

RNA extraction and real-time PCR

Total RNA was extracted from isolated brains by TRIzol (Invitrogen, Carlsbad, CA), according to the manufacturer’s instructions. Contaminating DNA was removed by subjecting 2 μg total RNA to DNase I digestion. The resulting total RNA was used for reverse transcription with the oligo(dT) primer. One-fifth volume of the reverse transcription reaction mixture was used for real-time PCR with the primer pairs for CXCL9, CXCL10, IFN-γ, TNF, and IL-10 (Supplemental Table I). PCR was performed with the SYBR Green PCR Master Mix for 40 cycles in an ABI PRISM 7700 apparatus (Applied Biosystems, Foster City, CA). Threshold values were obtained using PE Biosystem software, and mRNA was quantified. β-actin was used as an internal control, and the ratio of each target gene to β-actin was determined. An untreated control sample value was taken as 100%, and treated values were calculated based on the control. The specificity of the PCR was confirmed by melting-curve analysis.

Quantification of cytokine production by splenocytes

Splenocyte culture was performed as described previously (28). Briefly, spleens from control and infected mice were removed aseptically and pressed through a sterile fine-wire mesh with 10 ml RPMI 1640 medium supplemented with 10% heat-inactivated FCS, 25 mM HEPES, 0.12% gentamicin, and 2 mM glutamine. Cell suspensions were collected by centrifuging at 350 × g for 10 min. Erythrocytes were lysed with cold 0.17 M NH₄Cl, and cells were washed twice with fresh medium. Spleen cell viability was determined by trypan blue exclusion and was >90%; Spleen cells were adjusted to a final concentration of 10⁷ cells/ml in RPMI 1640 medium supplemented with 10% heat-inactivated FCS. Aliquots of 5 × 10⁷ cells/well were incubated in 24-well flat-bottom tissue culture plates (Falcon) in triplicate for 48 h at 37°C in a humidified 5% CO₂ incubator. Cytokines (IFN-γ, TNF, and IL-10) in the culture supernatant and plasma samples were measured by ELISAs (R&D Systems, Minneapolis, MN).

Flow cytometry

For each experimental group, five mice were sacrificed at the indicated time points for flow cytometric measurement of Th1 cells, Tregs, DCS, and CD11c⁺ DCS expressing the costimulatory marker MHC class II (MHC II), CD86, TRL-4, and TLR9 in splenocytes. Unless otherwise indicated, Abs were purchased from BD Biosciences (San Jose, CA). To measure Th1-type cells (CD4⁺ T-bet⁺ IFN-γ⁺), 10³ fresh splenocytes were stimulated in 12-well plates with 50 ng/ml PMA and 1 nM ionomycin (Sigma–Aldrich) for 5 h at 37°C and 5% CO₂ in the presence of 1 μM monensin to inhibit cytokine secretion. Cells were harvested and surface stained with FITC-conjugated anti-CD4 mAb (clone H1.2F3). After fixation and permeabilization with the intracellular fixation kit (eBioscience, San Diego, CA), cells were stained with PE-Tet-PE (clone 4B10) and anti-IFN-γ (clone XMG12.2). Tregs were surface stained with FITC-anti-CD4 and PE-anti-CD25 Abs in 100 μl PBS supplemented with 3% FCS and were then stained for intracellular Foxp3 with allophycocyanin-anti-Foxp3 (clone FJK16s) as described earlier (29). For DCS, cells were double-stained with FITC-conjugated CD11c mAb (clone HL3) and PE-conjugated anti-CD11b (clone M1/70). CD86 (clone GL1), MHC II mAb (clone M5/114.15.2), CD11c (clone I C35/R/220) (clone IC35/R/220) (clone 233/220), and anti-TLR4 (clone MTS510). To assess the expression of TLR9 in CD11c⁺ DCS, spleen cells were stained first with FITC-conjugated CD11c mAb. After fixation and permeabilization, cells were incubated with biotinylated anti-
To determine the migration of CD4+ and CD8+ T cells to the brain, brain mononuclear cells were isolated from the brains on day 5 postinfection (p.i.) following a published procedure (30). A single-cell suspension was obtained by grinding the tissues and resuspending the resulting cells in 5 ml RPMI 1640 medium, which contained 100 U/ml IV collagenase (Invitrogen), and incubated at 42°C for 45 min. Cells were pelleted at 300 × g for 10 min, resuspended in 30% Percoll in PBS (Sigma-Aldrich), layered on 70% Percoll, and centrifuged at 515 × g for 30 min at room temperature. Cells at the interface were isolated, washed twice, resuspended in PBS, and labeled with the following Abs: FITC-anti-CD4, PerCP-anti-CD8 (clone 53-6.7), and allophycocyanin-anti-CD3 (clone 145-2C11). Cells were incubated at 4°C for 30 min and washed twice with PBS. This brain cell isolation technique collects both intravascular and extravascular cells, thus allowing the evaluation of both already trafficked and recruited cells. Flow cytometric analysis was performed using a FACSCalibur, and data were analyzed with the FlowJo software.

Statistical analysis

Data are presented as the means ± SEM. VD levels in human participant were compared among three groups: patients infected with P. falciparum, patients infected with P. vivax, and healthy participants with no malaria infection. Kruskal–Wallis ANOVA test was used to look for statistically significant median values across the three groups, whereas an ANOVA was conducted on the log-transformed VD values. Difference in survival of the mice among experimental groups was assessed using the Kaplan–Meier test. For comparisons between two groups, statistical significance was analyzed by either a t test or Mann–Whitney U test, depending on normality of the data. For comparisons between three or more groups, statistical significance was determined using a one-way ANOVA. The data were analyzed using GraphPad Prism software (version 6.01), and a p value < 0.05 was considered significant.

Results

VD administration protects mice against ECM

The effect of VD on PbA infection depended on the delivery route and dose of the VD. Earlier work showed that there was no effect of injection of VD on P. berghei infection in mice (24, 25). Injection using the same protocol reported (25) offered no protection against ECM (Supplemental Fig. 1A, 1B). Preliminary experiments using a dose response of oral VD showed that there was no effect of 10 and 250 μg/kg VD as compared with the controls. In contrast, the mice treated orally with 50 μg/kg PbA infection did not develop ECM (Supplemental Fig. 1C). Therefore, we selected the 50 μg/kg dosage for subsequent experiments.

We applied three oral VD treatment schemes: daily oral VD supplement of 50 μg/kg for 4 d prior to PbA infection (VD+PbA), daily oral VD supplement of 50 μg/kg for 4 d beginning 2 d (PbA+VD2d), and 5 d (PbA+VD5d) after PbA infection. The four doses of VD supplementation prior to and 2 d p.i. was effective at raising the serum 25(OH)D3 levels of mice (Supplemental Fig. 1D). On day 5 p.i., the 25(OH)D3 levels in the VD-treated groups were significantly higher than that in the untreated PbA group. There was no effect of the infection on serum 25(OH)D3 levels because VD status was the same in the healthy control mice and the PbA-infected mice (data not shown).

The three VD treatment schemes offered significant protection of mice against ECM. In the control group, PbA infection resulted in the development of neurologic symptoms on days 5–6, and mice began to die on days 6–7, with all of the mice dead by day 11 p.i. (Fig. 1A). Death following PbA infection occurred in mice that had relatively low parasitemia (<12%) (Fig. 1B). In PbA-infected mice, the vehicle control groups (soybean oil) had symptoms (pathology, survivorship, or parasitemia) identical to the untreated group (data not shown). VD treatments either before or 2 d after PbA infection protected the mice from early mortality and all of the VD-treated mice survived beyond day 16 p.i. (Fig. 1A). VD administration beginning on day 5 p.i. also offered protection of the mice from early mortality with no mortality in this group before day 11. Regardless of the VD treatment all of the PbA-infected mice progressed to death from anemia with high parasitemia after ∼3 wk (Fig. 1A, 1B). There was an advantage to VD supplementation on day 2 p.i. because the PbA+VD2d group survived significantly longer (Kaplan–Meier test, p < 0.05) than the other two VD treatment groups (Fig. 1A). In addition, the parasitemias in both the PbA+VD2d and PbA+VD5d groups were lower than that of the VD+PbA group (Fig. 2B). For subsequent experiments, we focused on the PbA+VD2d group (referred hitherto as PbA+VD) for more detailed pathological and immunological analyses.

VD treatment reduces brain pathology

The pathological changes of ECM that mark the early lethality of mice following PbA infection includes adhesion of leukocytes in the brain microvasculature, microhemorrhages, disruption of BBB integrity, and extravasation of pRBCs into the brain parenchyma (31, 32). Activation and upregulation of endothelial adhesion molecules are responsible for these pathologies (33, 34). Thus, we assessed the brain pathology and expression levels of adhesion molecules in cerebral vessels. On day 5 p.i., the number of vessels containing leukocytes per microscopic field in the PbA+VD group was significantly lower than that in the untreated group (Fig. 2). Immunostaining for VCAM-1, ICAM-1, and CD36 on the endothelium of the microvessels was less intense in PbA+VD than the PbA group (Fig. 2). Meanwhile, a substantial loss of BBB integrity was evident in the PbA group, which showed high levels of dye extravasation. In comparison, VD treatment offered significant protection of the integrity of BBB in PbA-infected mice (p < 0.05) (Fig. 2). VD treatment beginning 2 d after PbA infection protected the brain from pathologic signs of ECM.

FIGURE 1. Treatment with VD improves survival in mice with PbA-induced ECM. The survival curves of infected mice (A) and parasitemia (B) following VD treatment in three groups of mice were compared: PbA control (PbA) (n = 40); daily VD administration for four successive days prior to PbA infection (VD+PbA) (n = 28); daily VD administration for 4 d 2 d after PbA infection (PbA+VD2d) (n = 32), and daily VD administration for 4 d 5 d after PbA infection (PbA+VD5d) (n = 40). Shown in this figure are the cumulative results of four separate experiments. Parasitemia is shown as mean ± SEM.
VD treatment reduces T cell trafficking to the brain

Increased trafficking and accumulation of pathogenic CD8+ T cells in the brain mediated through the IFN-γ–inducible chemokines CXCL9 and CXCL10 are associated with ECM (35–37). To determine whether VD treatment affected the migration and accumulation of T cells in the brains of PbA-infected mice, brain mononuclear cells were isolated and quantified for CD8+ and CD4+ cells. Compared with uninfected naive mice, PbA infection promoted the accumulation of both CD8+ and CD4+ T cells (Fig. 3A). In contrast, the numbers of both T cell populations in the brains of VD-treated PbA-infected mice did not differ significantly from those in uninfected naive mice (Fig. 3A). To determine whether reduced T cell trafficking to the brain was associated with decreased expression of chemokines, we compared the expression of CXCL9 and CXCL10 mRNA in the brains of VD-treated and untreated mice. On day 5 p.i., VD treatment significantly reduced the expression of both chemokines in the brains (Fig. 3B, 3C). These results indicate that VD treatment is sufficient to modulate the cerebral environment to downregulate expression of key chemokines, leading to decreased accumulation of pathogenic T cells in the brain.

VD treatment dampens the Th1 immune response

An overwhelming proinflammatory (Th1) response is essential for ECM (7, 38, 39). To test whether VD treatment inhibits Th1 cells during ECM, we quantified the IFN-γ–producing CD4+ T-bet+ cells...
in the spleens of infected mice and two Th1 cytokines IFN-γ and TNF on days 3 and 5 p.i. In association with the appearance of ECM pathology, IFN-γ and TNF levels were significantly increased in the serum of PbA-infected mice on day 5 p.i. \( (p < 0.01) \) (Fig. 4A, 4B). Significantly increased production of these cytokines also was observed by splenocytes isolated from PbA-infected mice

**FIGURE 3.** VD treatment reduced pathogenic T cell sequestration in the brain. (A) VD treatment of PbA-infected mice reduced the number of CD4\(^+\) and CD8\(^+\) T cells in the brains \((n = 5)\). Values are means plus SEMs (error bars). (B and C) VD treatment significantly reduced the expression of the chemokines CXCL9 and CXCL10 in the brains of PbA-infected mice. The mRNA levels for the two chemokines were normalized to the \(\beta\)-actin mRNA level, and fold changes were calculated against control uninfected mice. \( *p < 0.05, \) significant difference between mRNA levels in infected mice on day 5 and their corresponding baseline levels on day 0; \( \#p < 0.05, \) significant difference between the PbA and PbA+VD groups on day 5 p.i.

**FIGURE 4.** VD treatment reduced expression of proinflammatory cytokines (IFN-γ and TNF). (A and B) Levels of circulating cytokines in serum. (C and D) Production of IFN-γ and TNF by splenocytes. (E and F) mRNA levels of IFN-γ and TNF in the brain of mice. Each experiment was repeated three times \((n = 5/group)\). Values represent mean ± SEM. \( *p < 0.05 \) and \( **p < 0.01, \) significant differences between the values in PbA-infected mice and the baseline levels on day 0, respectively. \( \#p < 0.05 \) and \( \###p < 0.01, \) significant difference between the PbA and PbA+VD groups.
during their in vitro culture ($p < 0.01$) (Fig. 4C, 4D). VD treatment led to a significant reduction of the levels of these two cytokines in both serum and cultured splenocytes of PbA-infected mice (Fig. 4A–D). In addition, the mRNA levels of IFN-γ and TNF in the brains of infected mice on day 5 p.i. also experienced significant declines in VD-treated mice as compared with untreated mice (Fig. 4E, 4F).

PbA infection was associated with increased expansion of CD4+T-bet+IFN-γ+ T cells. On both days 3 and 5 p.i., the total population as well as the proportion of this Th1 cell subtype in the spleens of infected mice were substantially increased (Fig. 5). VD treatment, however, significantly reduced this Th1 cell subtype in the spleens of infected mice on day 5 p.i. ($p < 0.01$) (Fig. 5).

VD promotes expansion of Tregs and the production of IL-10 in spleen

Both Tregs and IL-10 contribute to the improvement of PbA-specific Th1 response involved in the induction of ECM. Overall, compared with the untreated group, VD treatment promoted the expansion of CD4+CD25+Foxp3+ Tregs in PbA-infected mice, albeit significance ($p < 0.01$) was only observed in total Treg populations on day 3 p.i. (Fig. 6A, 6B). However, the secretion of the anti-inflammatory cytokine IL-10 by cultured splenocytes was significantly higher in VD-treated groups than untreated PbA groups ($p < 0.05$) (Fig. 6C). Meanwhile, the plasma level of IL-10 was also significantly increased in VD-treated mice on day 5 p.i. ($p < 0.01$) (Fig. 6D).

VD inhibits DC differentiation, maturation, and function

The nature of the immune response is critically dependent on the interplay between the innate and adaptive immune systems. Central to this interaction are the DCs (40). We found that the proportions of both myeloid DCs (mDCs) (Fig. 7A, 7C) and plasmacytoid DCs (pDCs) (Fig. 7B, 7D) were significantly higher than those in the VD-treated groups on day 5 p.i. ($p < 0.05$, t test), indicating that VD inhibited differentiation of DCs. In addition, VD treatment also led to a decrease of expression of the costimulatory molecules MHC II on DCs on day 5 p.i. (Fig. 8A, 8D). Furthermore, the expression of TLR4 on DCs was drastically reduced in VD-treated groups on day 5 p.i. (Fig. 8B, 8E), whereas TLR9 expression in the DCs was also reduced to some extent (Fig. 8C, 8F). Meanwhile, the total numbers of DC populations within the spleen of mDCs, pDCs, and DCs expressing MHC II, CD86, TLR4, and TLR9 were mostly downregulated by VD treatment (Supplemental Fig. 2). To further illustrate that VD treatment affected the function of DCs, we measured cytokine production by DCs and found that VD treatment resulted in decreased production of IL12p40 on both days 3 and 5 p.i. and increased production of IL-10 on day 5 (data not shown). Taken together, these results

![FIGURE 5. VD reduces the expansion of CD4+T-bet+IFN-γ+ T cells. Both proportion (A) and the absolute number (B) of the CD4+T-bet+IFN-γ+ T cells in the spleens were quantified. Each experiment was repeated three times. For each experiment, five mice were used per group. Values represent the mean ± SEM (n = 5 mice/group). *$p < 0.05$ and **$p < 0.01$, significant differences between the values in PbA-infected mice and the baseline levels on day 0, respectively. ##$p < 0.01$, significant difference between the PbA and PbA+VD groups.](http://www.jimmunol.org/)

![FIGURE 6. VD stimulated the expansion of Tregs and increased IL-10 production. The absolute number (A) and the proportion (B) of CD4+CD25+Foxp3+ Tregs were quantified by flow cytometry. The concentrations of IL-10 secreted by cultured splenocytes (C) and in the mouse serum (D) were determined by ELISA. Results are representatives of three independent experiments. Values are presented as the mean ± SEM (n = 5 mice/group). *$p < 0.05$ and **$p < 0.01$, significant differences between the values in PbA-infected mice and the baseline levels on day 0, respectively. ##$p < 0.05$ and ###$p < 0.01$, significant difference between the PbA and PbA+VD groups, respectively.](http://www.jimmunol.org/)

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indicate that VD treatment inhibited the differentiation, maturation, and activation of DCs.

**Discussion**

Susceptible mice infected with PbA developed a severe CM syndrome characterized by hyperinflammation, intravascular accumulation of immune T cells, and BBB leakage (38). Although it has been extensively shown that in vivo treatments with 1,25(OH)2D3 inhibit Th1-mediated immune responses (20, 21) and suppress experimental autoimmunity, to our knowledge this is the first study to demonstrate the effectiveness of short-term VD treatments to suppress in vivo Th1-mediated inflammation. It is well established that VD and 1,25(OH)2D3 directly and indirectly via actions on DCs and macrophages inhibit Th1 cell production of IFN-γ. We demonstrated that oral administration of VD improved serum 25(OH)D3 levels and downregulated the levels of circulating inflammatory cytokines (IFN-γ and TNF). The inhibition of Th1 responses in addition with the reduced expression of chemokines CXCL9 and CXCL10 and cell adhesion molecules ICAM-1, VCAM-1, and CD36 resulted in the decreased accumulation of CD8+ T cells in the brain and improved the integrity of the BBB. Interestingly, the timing of the VD supplementation looks to be critical and dosing animals just p.i. seems to be more effective than before infection. There was no effect of infection on serum 25(OH)D3 levels in mice. Analysis of plasma VD levels in humans infected with *P. falciparum* or *P. vivax* malaria from a subtropical area (Kachin State, Myanmar) found highly variable 25(OH)D3 levels including frank VD deficiency (<12 ng/ml) in 10% and VD deficiency <20 ng/ml in 28% of the population (data not shown). It would be interesting to determine whether lower VD status would predict development of CM. Although individuals in tropical malaria regions may not suffer from low sunlight exposure, VD deficiency is still prevalent in tropical climates despite high amounts of sunlight (41–43). VD production in the skin following UV exposure is significantly less in dark skin, and dietary sources for VD is uncommon in most parts of the world. This study suggests that VD might be an effective treatment for reducing lethality from CM in humans, albeit the potential difference in VD physiology between humans and mice demands further investigation.

An overwhelming proinflammatory Th1 response characterized by excessive production of the cytokines IFN-γ and TNF is involved in CM pathogenesis (4). The cerebral pathology during ECM is an IFN-γ–dependent process as evidenced by the complete resistance of the IFN-γ–deficient mice to ECM (39). IFN-γ is responsible for the induction of chemokines CXCL9 and CXCL10 during ECM (44), which subsequently recruit CD4+ and CD8+ T cells to the brain via CXCR3 on the cell surface (35, 36, 45). In addition, the proinflammatory cytokine TNF is proposed to upregulate the production of adhesive molecules in the brain vasculature. In this study, we showed that VD treatment significantly reduced the levels of circulating IFN-γ and TNF, secretion of these cytokines by the immune cells in the spleen, as well as expression of these cytokines in the brains of infected mice. Furthermore, VD inhibited the differentiation of CD4+ IFN-γ−T-bet+ Th1 cells. VD deficiency is associated with an increased risk of disease in Th1-mediated diseases such as inflammatory bowel disease (20). VD via the VD receptor (VDR) and 1,25(OH)2D3 directly inhibits IFN-γ transcription in human and mouse T cells (46, 47). In addition, TNF production is inhibited by 1,25(OH)2D3 in macrophage and T cells (20, 48). Direct inhibition of TNF and IFN-γ following VD supplementation protects against ECM.

**FIGURE 7.** VD treatment inhibited the expansion of DC subsets. The frequency of CD11c+CD11b+ DCs (mDCs), and CD11c+B220+ DCs (pDCs) were measured by flow cytometry. (A and B) Representative plots showing the proportions of the two DC populations. (C and D) Bar graphs show the frequencies of the two DC populations within the spleens. Data are presented as the mean ± SEM (n = 5 mice/group). Results are representatives of three independent experiments. *p < 0.05, significant difference between the values in PbA-infected mice and the baseline levels on day 0. #p < 0.05, significant difference between the PbA and PbA+VD groups.
VD exerts its immunomodulatory effects through interaction with VDR, a member of the superfamily of nuclear hormone receptors, which is expressed in numerous human immune cells such as T cells, B cells, invariant NKT cells, macrophages, and DCs (49). Thus, VD directly acts on T cells to inhibit T cell proliferation and IFN-γ production (50). DCs, the highly specialized APCs, are critical for stimulating the differentiation of effector T cells from naive T cell precursors. There is ample evidence showing that VD also hampers DC maturation from monocytes (51–54). Our data show that VD treatment decreased the number of mDCs and pDCs during P. falciparum infection. This decrease may be due to changes in DC migration and homing. In addition, VD suppressed the expression of Ag-presenting molecules, such as MHC II and CD86 on DCs, and changes in surface marker with corresponding change in DC function such as Ag presentation (55). These results are consistent with the role of VD for inhibiting differentiation, maturation, and activation of DCs. Furthermore, we found decreased expression of TLR4 and TLR9 in DCs upon VD treatment during PbA infection. Because DCs are essential to induce Th1 cell development as well as Th1 cytokine production upon malaria infection (8), our data demonstrate that VD also results in decreased expression of IL-12 and increased production of IL-10 in these APCs, consistent with a decrease of the Th1 response (56). Collectively, these data are in line with earlier descriptions of the suppressive activities of VD with respect to the stimulation of Th1-mediated immunity.

Tregs expand during Plasmodium infection (57, 58) and have been shown to inhibit the development of Th1 immune responses (59). The critical role of Tregs and the anti-inflammatory cytokine IL-10 in inhibiting CM, expansion of Tregs and elevation of IL-10 levels would improve the outcome of the ECM. VD has been shown to be required for the optimal development and function of Tregs. VD increased the Treg population (Foxp3) (60), resulting in the blockade of the Th1 response (61). Importantly, treatment of naive CD4+ T cells with VD potently induced the development of Tregs (62, 63). Meanwhile, VD alone or in combination with dexamethasone induced IL-10–producing Tregs in an APC-free in vitro system (64, 65). Also, a VD analog triggered the emergence of a CD4+CD25highCD127low Treg phenotype and selectively induced IL-10 expression within the CD4+ T cell subset (66). These results highlighted the impacts of VD on the differentiation of Tregs and expression of IL-10. Our results provided further evidence to show that VD treatment expanded the Treg population, which was accompanied by increased levels of IL-10, conditions that favor host protection during CM.

The role of VD regulation of infection has been more difficult to determine. For many infections including malaria strong Th1
responses are needed to clear the parasite. Our work suggests that VD is protective against ECM but not PbA infection because the VD treated mice survive the ECM but then several weeks later show increased parasitemia and eventual death caused by anemia. Interestingly, murine models of infectious diseases, where IFN-γ are required for host protection, were not affected by 1,25(OH)2D3 treatments (20, 21). Our data support a model where VD and production of increased expression of IL-10. The direct effects of VD on IFN-γ maturation, and functioning of DCs, which together resulted in that VD treatments expanded Tregs and inhibited differentiation, APCs and innate immunity. In addition, our results demonstrate the Th1 cells as well as indirectly through the inhibition of the host Th1 response by VD is mediated directly by its action on the Th1 cells as well as indirectly through the inhibition of the APCS and innate immunity. In addition, our results demonstrate that VD treatments expanded Tregs and inhibited differentiation, maturation, and functioning of DCs, which together resulted in increased expression of IL-10. The direct effects of VD on IFN-γ and TNF and the indirect effects of VD on DCs and production of IL-10 serve to suppress the Th1-driven ECM following VD treatment. Importantly, administration of VD either before or after Plasmodium infection was effective in preventing ECM, suggesting that both prophylactic treatment and p.i. therapy could benefit human CM patients.

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Disclosures
The authors have no financial interests of conflict.

References


Table S1. Primers used for real-time RT-PCR analysis

<table>
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**Figure S1.** (A) Injection of VD did not offer protection against cerebral malaria in PbA infected mice. The survival of infected mice was compared between VD injected (VD+PbA) and control (PbA) groups. Mice in the VD+PbA group (n=10) were injected every other day with 0.5 µg/kg/day of VD three days before PbA infection, while the control group (n=10) received equal volumes of 0.02% Tween 80. (B) Parasitemias in the control and VD-injected groups shown in (A). (C) Survival curves of PbA-infected mice in the control group (n=10) received vegetable oil only and in three groups of mice received daily oral supplement of VD at 10, 50 and 250 µg/kg for four days before PbA infection. (D) 25(OH) vitamin D levels in PbA-infected mice. PbA – mice received only vegetable oil (n=6); VD+PbA – mice received daily vitamin D oral supplement for four days prior to PbA infection (n=9); PbA+VD – mice received daily vitamin D oral supplement for four days 1 day after PbA infection (n=10). VD levels were measured on day 5 post infection. # and ## indicate significant differences at P=0.05 and P=0.01, respectively (t-test).
**Figure S2.** VD treatment inhibits DC differentiation, maturation and activation. The absolute numbers of DC populations within the spleen were determined by flow cytometry. (A) mDCs; (B) pDCs; (C) DCs expressing MHC II; (D) DCs expressing TLR4; (E) DCs expressing TLR9, and (F) DCs expressing CD86. Data are presented as the mean ± SEM (n=5 per group). Results are representatives of three independent experiments. * and ** indicate significant differences between the values in PbA-infected mice and the baseline levels on day 0 at $P<0.05$ and $P<0.01$ (t-test), respectively. # and ## indicate significant difference between the PbA and PbA+VD groups at $P<0.05$ and $P<0.01$ (t-test), respectively.