B Cell: T Cell Interactions Occur within Hepatic Granulomas during Experimental Visceral Leishmaniasis

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Abstract

Hepatic resistance to Leishmania donovani infection in mice is associated with the development of granulomas, in which a variety of lymphoid and non-lymphoid populations accumulate. Although previous studies have identified B cells in hepatic granulomas and functional studies in B cell-deficient mice have suggested a role for B cells in the control of experimental visceral leishmaniasis, little is known about the behaviour of B cells in the granuloma microenvironment. Here, we first compared the hepatic B cell population in infected mice, where ≈60% of B cells are located within granulomas, with that of naïve mice. In infected mice, there was a small increase in mlgMlo/ldLo mature B2 cells, but no enrichment of B cells with regulatory phenotype or function compared to the naïve hepatic B cell population, as assessed by CD1d and CD5 expression and by IL-10 production. Using 2-photon microscopy to quantify the entire intra-granuloma B cell population, in conjunction with the adoptive transfer of polyclonal and HEL-specific BCR-transgenic B cells isolated from L. donovani-infected mice, we demonstrated that B cells accumulate in granulomas over time in an antigen-independent manner. Intravital dynamic imaging was used to demonstrate that within the polyclonal B cell population obtained from L. donovani-infected mice, the frequency of B cells that made multiple long contacts with endogenous T cells was greater than that observed using HEL-specific B cells obtained from the same inflammatory environment. These data indicate, therefore, that a subset of this polyclonal B cell population is capable of making cognate interactions with T cells within this unique environment, and provide the first insights into the dynamics of B cells within an inflammatory site.

Introduction

B cells have many diverse roles in immunity extending far beyond their ability to produce antibodies. B cell-derived LTαβ2 plays a role in i) the differentiation of follicular dendritic cells (FDCs), mesenchymal cells that subsequently produce B cell survival factors and follicle-homing chemokines to maintain follicular architecture and function in lymphoid tissues [1] and ii) in infection-induced remodelling of this compartment [2]. During the early stages of an immune response, B cells function as APC and interact in a cognate manner with CD4⁺ T cells to facilitate their own activation and differentiation into antibody secreting cells [3], and within the follicle, the interaction between B cells and follicular helper T cells is essential for survival and to rescue and expand high affinity class-switched B cells [4,5,6]. During this cross-talk, B cells also contribute cytokines or other factors that facilitate CD4⁺ T cell functional differentiation [7]. In addition, a growing number of studies also implicate B cells, directly or indirectly, as negative regulators of T cell immunity through IL-10-dependent and IL-10-independent mechanisms [8,9,10,11,12,13]. A number of recent studies have exploited intravital 2-photon microscopy to dissect the dynamic behaviour of B cells in the follicular and extra-follicular environments of peripheral lymph nodes, defining many of the dynamic features associated with cognate B cell-T cell and B cell-DC interactions [14,15,16,17,18,19]. However, in spite of the common occurrence of B cells at inflammatory sites induced by infection and autoimmune disease [15,20,21], there is little known about the dynamic behaviour of B cells outside of peripheral lymphoid tissues.

The role of B cells in regulating immunity to leishmaniasis has been most extensively studied in models of cutaneous leishmaniasis, but with conflicting findings. Treatment with anti-IgM showed that susceptibility to L. major in BALB/c mice is B-cell dependent, possibly mediated by regulatory T cell induction by B cells [22,23], whereas in resistant C3H/HeN mice, B cells contribute to resistance [24]. Ronet et al. used B cell transfer experiments to show that the impact of B cells on susceptibility to L. major infection was parasite strain specific [25] and studies in T cell-reconstituted scei mice indicated that co-transfer of B cells could promote disease in a non-cognate manner [26]. Other studies however showed that B cell-deficient μMT mice did not exhibit any defects in T cell priming and polarisation [27].
addition, in another study, IL-7 treatment lead to B cell expansion, accelerated lesion growth and reduced survival [28]. Most recently, IL-10-producing B cells have been found in BALB/c mice infected with L. major and proposed to shape Th2 development [12]. In a different model of cutaneous leishmaniasis, infection of B cell-deficient JbD mice with L. amazonensis led to delayed lesion development, and these mice displayed decreased CD4+ T cell recruitment to skin granulomas and decreased CD4+ T cell responses, suggesting that B cells may play a role in priming the immune response against L. amazonensis infection [29].

In experimental visceral leishmaniasis (EVL), early studies identified infrequent B220+ B cells in the hepatic granulomas of BALB/c mice infected with L. donovani, as well as B cells lying adjacent to infected Kupffer cells in infected athymic (nu/nu) BALB/c mice [30]. An inhibitory role for B cells in host resistance was indicated by studies of L. donovani infected B cell-deficient B6,µMT mice, which showed that these mice had accelerated granuloma formation and reduced hepatic parasite burden compared to B6 mice. Resistance could not be reversed by serum-transfer [31]. Furthermore, B6,µMT mice also had exaggerated neutrophil recruitment and suffered severe tissue pathology, suggesting that B cells may also play a role in protecting against host-mediated damage. More recently, polyclonally activated B cells appearing early after L. infantum infection and producing IgM have been implicated in disease exacerbation [9]. Although the induction of myeloid cell-derived IL-10 by B cells has been described [32], this study indicated a distinct IL-10-independent role for C5 activation and C5aR (CD88) signalling in IgM-mediated immune regulation [9].

In addition to providing information on the cellular mechanism of host immunity to L. donovani, the hepatic granulomas formed during EVL provide an excellent model system to study the dynamics of cellular interactions outside of peripheral lymphoid tissues, combining a variety of attributes including small size, complex cellular composition and accessibility to intra-vital imaging [33,34,35]. Here, we address whether B cell recruitment to hepatic granulomas is a function of phenotype, time, antigen specificity and/or the development of tertiary lymphoid organ (TLO) characteristics. We show that B cells accumulate over time p.i., but enter and exit granulomas with relative ease independently of antigen specificity and without enrichment of regulatory B cells. By comparing the dynamics of polyclonal B cells and transgenic B cells with an irrelevant specificity, we demonstrate significant differences in their interactions with intra-granulomas T cells and the existence of cognate B:T cell interactions within the granuloma microenvironment. These data provide the first glimpse of the behaviour of B cells in an inflammatory setting and should inform the design of future imaging studies in this and other disease settings.

**Results**

**B cells traffic into L. donovani-induced hepatic granulomas**

In BALB/c mice, intra-granuloma B220+ cells have been described [30]. Similarly, in B6 mice, B220+ B cells were mostly observed in the periphery of granulomas (Figure 1A and B), with number being directly proportional to granuloma cross-sectional area (r = 0.4756; p < 0.05; Figure 1C). Although by flow cytometry, ~90% of B220+ cells co-expressed CD19, B220+ CD19+ cells expressing CD11c and/or γδ TCR were also present (Figure 1D). To more selectively examine B cells, we infected CD19+ xR26eYFP xVaDsRed.A mice (herein called Bgont/Tgont mouse; Figure 2), where eYFP is expressed on all cells that have expressed CD19 or are derived from CD19+ cells [36], and DsRed is expressed on T cells under the hCD2 locus control region [37] (Figure S1). We also employed 2-photon microscopy to capture the entire 3-dimensional cellular composition of individual granulomas (Figure 2; [33]), to avoid the sampling limitations of conventional thin sections. CD19+ cells were readily detected at d14 p.i. (Figure 2A), though outnumbered 12:1 by T cells (Figure 2D). At d21 and day 28 p.i., B cells were more abundant (62.9 ± 1.6% of all hepatic B cells being found within granulomas; n = 95 FOV, n = 3 mice at d21) and the T:B cell ratio within hepatic granulomas was decreased (Figure 2B–D). In some but not all granulomas, B cells aggregated into small clusters (Figure 2E and F). Although B cell aggregation is often associated with the development of TLOs and associated FDCs [38], we could not detect ectopic CD21/CD35+ FDCs within hepatic granulomas (Figure 2G).

**Phenotype of hepatic B cells in infected mice**

As hepatic granulomas cannot be isolated in a reproducible way, total hepatic B cell populations (of which ~60% represent intra-granuloma B cells, see above) were studied by flow cytometry. In naive livers, CD19+ mIgMlo/mIgDlo- (transition- al/MZ), mIgM+ mIgDlo+ (T2) and mIgM+ mIgDhi+ (mature) B cells were all found, in addition to a small proportion of mIgMlo- "mIgDhi+" cells, which likely represent class-switched mature B cells. In infected mice, the frequency of T2 B cells was reduced with a commensurate increase in mIgMlo/mIgD+ mature B cells (Figure 3A and B). Indicative of activation, we found a 1.7 ± 0.2-fold increase in MFI for MHCII in CD19+ B cells from infected compared to naive mice. In contrast, CD40 expression was unchanged (Figure 3C and D). Plasma cells (CD138+) were not specifically recruited into the hepatic microenvironment of infected mice and were diluted out in the hepatic B cell pool of infected mice (Figure 3E and F).

Expression of CD1d and CD5 in naive and infected mice was minimal (<3%), again with evidence of dilution by mature B cells in infected mice (Figure 4A–C). After PMA, ionomycin and LPS stimulation [39] negligible levels of IFN-γ and IL-4 were detected in CD19+ cells (Figure 4D, E and H). CD19+ cells produced TNF, but the frequency was lower in infected compared to naive mice (Figure 4F, H and I). A small fraction of CD19+ cells also made IL-10 and accumulated detectable levels of IL-10 mRNA, but hepatic B cells from infected mice did not have an elevated capacity for IL-10 production (Figure 4G, H and I). Collectively, these data suggest that most intra-hepatic B cells in infected mice have a phenotype consistent with recruited T2 and mature re-circulating B2 cells that have been non-specifically activated by the cytokine environment in infected mice.

**Adoptively transferred B cells are recruited to hepatic granulomas**

To confirm that peripheral B cells could be recruited into granulomas from the circulation, we co-transferred CD19+ B cells (>90% purity; data not shown) from infected and naive mice that expressed similar levels of the liver homing receptor CCR6 [40], into d21-infected VaDxRed (herein called Tgont) mice (Figure 5A–D). 12 h following B cell transfer, labelled B cells originating from both naive and infected mice were found in similar numbers (Figure 5E–G). Hence, within 12 h of transfer, B cells from naive as well as infected mice are capable of being recruited from the periphery into pre-existing granulomas.
Dynamic imaging of B cells in hepatic granulomas

As the behaviour of B cells outside of the LN is unknown, we first examined the dynamics of an endogenous B cell population in granulomas of infected B<sup>green</sup>/T<sup>red</sup> mice. B cells moved freely, with average velocity of 4.2±0.3 μm/min (Figure 6A), similar to granuloma T cells [33,35] and B cells in lymphoid tissue [14,17,18,19]. Under static imaging conditions, T-B conjugates were readily observed, sometimes being associated with collagen fibres that traversed the granuloma (Figure 6B and C).

To examine these interactions further, we developed a co-transfer model allowing simultaneous analysis of two independent B cell populations within the same granulomas, thus controlling for heterogeneity in granuloma size, maturation status and/or cellular composition [33]. We used MD-4 transgenic B cells with defined but un-related antigen specificity (hen egg lysozyme; HEL) as a comparator for immune B cells (i.e. polyclonal B cells from infected mice). To ensure that all B cells has been exposed to the same inflammatory environment prior to transfer, we infected B6.CD45.1/2 bone marrow micro-chimeras (Figure 7A) and at d21 p.i., CD19<sup>+</sup> B cells were sort purified (>98% purity) on the basis of CD45.1/2 expression (Figure 7B). MD-4 B and immune B cells were labelled and co-transferred into d21-infected T<sup>red</sup> recipient mice. 12 h post transfer, intra-vital imaging was performed (Figure 7C and D). Similar numbers of immune B cells and MD-4 B cells were detected within granulomas at d21 p.i. (Figure 7E), further confirming that antigen-recognition per se was not required for B cells to be recruited into L. donovani-induced hepatic granulomas.

For 134 immune B cells and 121 MD-4 B cells, we determined individual displacement, track length and average velocity and calculated the displacement rate and meandering index (Figure 8A–E). By none of these criteria could we distinguish the behaviour of immune B cells from MD-4 B cells. This result was expected given the highly motile nature of B cell-T cell conjugates seen in lymphoid tissue [17] and the similar degree of mobility of granuloma B cells (Figure 6A) and T cells [33,35]. We then examined the ability of B cells to engage with endogenous T cells, using the study of Qi et al as a template for our analysis and to allow ready comparison with data obtained using MD-4 B cells in LN [17]. We analysed 1089 individual contacts between MD-4 B cells and endogenous T cells. The average contact duration of 2.8±0.11 min was similar to that observed for non-cognate interactions, mostly between 3 and 10 min of duration but often extending for 30 min (Figure 9A). In contrast to MD-4 B cells, the average contact time for immune B cells was 4.0±0.19 min (n=756 contacts, P<0.0001; Figure 9A), indicating that at a population level, these immune B cells had sustained engagement with T cells compared to MD-4 B cells. Stratifying the data according to contact time per incidence confirmed that there was a significantly greater trend for polyclonal B cells to make contacts of longer duration than those made by MD-4 B cells (Figure 9B; χ<sup>2</sup>=34.9, df=4, p<0.0001). We next analysed each individual B cell for the duration of contacts that it made with individual T cells. A range of behaviour was observed, from repeated short interactions to single long contacts that spanned the entire imaging window. Single frame images taken from the imaging videos, and contact plots (where each period of contact is indicated by a white box) are shown for two MD-4 B cells displaying different behaviour during a 30 min imaging window (Figure 9C and Video S1). For each B cell shown, there were also periods when simultaneous contact was made with multiple T cells (shown as adjacent white boxes in

Figure 1. B cells are present in L. donovani-induced hepatic granulomas. A and B. C57BL/6 mice (n = 3) were infected with 3×10<sup>7</sup> TdTOMato transgenic L. donovani amastigotes (red) and on day 21 p.i., liver sections were prepared and stained using an isotype control mAb or with B220 (green) mAb. Sections were counterstained with DAPI (blue). C. Correlation between granuloma cross-sectional area and number of intra-granuloma B220<sup>+</sup> cells. Each symbol represents a single granuloma. D. Hepatic mononuclear cells were labeled with CD19 and B220, and B220<sup>+</sup>CD19<sup>+</sup> cells (R1) were subsequently analyzed for CD11c and γδ TCR.

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Figure 2. The recruitment of B cells to *L. donovani*-induced hepatic granulomas. $B^{\text{green/Tred}}$ mice (*n* = 6) were infected with *L. donovani* and at d14 (A), d21 (B) and d28 (C) p.i. liver explants were imaged using 2-photon microscopy to visualize T cells and B cells. D. Number of T cells and B cells at d14 (open bars), d21 (grey bars) and d28 (black bars) p.i. derived from 25–35 hepatic granulomas per time point from 2 mice per group. Data are shown as mean ± SEM along with the T cell: B cell ratio. E. B cells aggregate in hepatic granuloma. F. The number of B cell aggregates per granuloma was determined from 25–35 granulomas (mean ± SEM). G. C57BL/6 mice were infected with *L. donovani* (*n* = 3). Frozen sections were labeled with B220 (red), CD21/35 (green) and counterstained with DAPI (blue, except left panel). 60 hepatic granulomas were imaged. Control staining of spleen was performed (left panel).

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Figure 3. Phenotype of hepatic B cells in *L. donovani*-infected mice. A. Hepatic mononuclear cells from naïve and d21 *L. donovani*-infected C57BL/6 mice were stained for CD19 along with membrane IgM and membrane IgD. B. Frequency of T1, T2, mature and class-switched B cells in naïve (grey bars) and infected (black bars) mice. Data represents the mean ± SEM for 5 independent experiments with cells pooled from 3 mice per experiment (**, *p* < 0.01). C. CD19$^+$ cells were stained for MHCII and CD40. D. Data are shown as fold change in MFI compared to naïve (mean ± SEM from 5 independent experiments). E. Expression of CD138 on CD19$^+$ cells from naïve and infected mice. F. Data are shown as % CD19$^+$ cells in naïve (grey bars) and infected (black bars) mice expressing CD138.

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Figure 4. Hepatic B cells in L. donovani infected mice do not display regulatory phenotype. CD19+ hepatic B cells were stained for CD1d (A) and CD5 (B). Data from 6 independent experiments (using pooled cells from 24 mice in total) comparing naive (grey bars) and infected (black bars) mice is shown as mean ± SEM (C); *, p<0.05. Hepatic mononuclear cells from naive and d21-infected C57BL/6 mice were stimulated with PMA, ionomycin and LPS and stained for IFN-γ (D), IL-4 (E), TNF-α (F) and IL-10 (G). Data are from 3 independent experiments, with cells pooled from 3–4 mice per experiment. H. Data from 3 independent experiments comparing naive (grey bars) with infected (black bars) mice is shown as mean ± SEM % of CD19+ cells positive for each cytokine. *, p<0.05. I. Hepatic CD19+ B cells from naive and d21-infected C57BL/6 mice were FACS purified and mRNA accumulation for IL-10, TNF and IL-6 was determined by quantitative RT-PCR. Data are shown as fold change compared to mRNA accumulation in naive mice and represent mean ± SEM for 3 mice assayed in duplicate.

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However, we did not observe any B cell division within granulomas during this study. When summated to provide an accumulated total contact time for each B cell, the total contact time for any individual B cell also varied widely (from less than 1 min to over 80 min). It was difficult, however, to discern any significant difference between the behaviour of the immune and MD-4 B cell populations by this approach (Figure 9D). Nevertheless, when we analysed the percentage of short (<3 min) vs. long (3–40 min) contacts made by each B cell population, immune B cells had a significant bias towards making long term contacts compared to the MD-4 B cells (p<0.01; Figure 9E). Although demonstrating a clear difference in behaviour at the population level, due to the variability in behaviour of the MD-4 B cells it was not possible to determine with any degree of statistical certainty the absolute frequency of immune B cells that engaged in these cognate interactions with granuloma T cells.

Discussion

B cells are a notable feature of inflammation, yet little is known about their behaviour and ability to interact with T cells outside of the lymphoid follicle. Using a model of hepatic granulomatous...
Figure 6. Endogenous B cell behavior in hepatic granulomas. A. B cell velocity in granulomas of d21-infected B<sup>green</sup>/T<sup>red</sup> mice. Each symbol represents an individual B cell. Bar shows median velocity calculated from 71 B cells in 12 granulomas imaged in two mice. Liver explants tissue from B<sup>green</sup>/T<sup>red</sup> mice was imaged using 2-photon microscopy and B cell-T cell contacts (B) identified by static imaging. The second harmonic signal is also visible (blue). Data represents the mean ± SEM % B cells interacting with T cells (C) and were derived from 25 and 35 granulomas from 2 mice per time point.

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Figure 7. Role of antigen specificity in B cell recruitment to granulomas. A. Protocol for mixed adoptive transfer. B. Pre and post sort purity of adoptively transferred B cells. C and D. Maximum projection image of transferred labeled MD-4 (green) and immune (blue) B cells in granuloma of T<sup>infl</sup> mouse 12 h post transfer, shown with (C) and without (D) DsRed channel switched on. E. Number of recruited MD-4 (black bars) and immune (grey bar) B cells found in granulomas 12 h post transfer. Data represent mean ± SEM B cell per granuloma derived from 28 granulomas from two mice.
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A  B  C  D  E

Figure 8. Dynamic behavior of MD-4 and immune B cells in the granuloma microenvironment. Labeled MD-4 and immune B cells were transferred into d21-infected Tm+ mice as described in Figure 7. At 12–30 h post adoptive transfer, time lapse z-stack videos were prepared, and each B cell was manually tracked in Velocity software to determine (A) displacement, (B) velocity, (C) track length, (D) displacement rate and (E) meandering index. Data are shown for each individual B cell analyzed within 28 granulomas from two mice.
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with CD4+ T cells lasting longer than 3 min were indicative of antigen-specific interactions. We observed a similarly short average contact time per incidence when following the behaviour of HEL-specific MD-4 cells in granulomas, in spite of a limitation of our model being that the identification of endogenous T cells was based on DsRed expressed from the hCD2 LCR, which includes both CD4+ and CD8+ T cells (and some NK cells) [44]. B cells also make relatively short non-cognate interactions with DC in LN, whereas cognate interactions between these two cells last an average of 10 min and may extend to >45 min [18]. Although DC are relatively sparse in granulomas and do not appear to contain parasites at an appreciable frequency or abundance [33], we have not directly examined whether B cells interact with DC in this environment. Our data do indicate, however, that many of the characteristics of B cell dynamics observed in LN appear to hold true when examining these cells in an inflammatory and non-lymphoid setting.

McMahon-Pratt and colleagues [9] have recently re-examined the well-known phenomenon of polyclonal B cell activation in EVL, with important bearings on the interpretation of our data. First, as a comparator B cell population for our dynamic imaging studies on immune B cells, we used HEL-specific MD-4 B cells that had been parked for 21 d in infected B6 mice. Given that the frequency of OVA, chromatin and TNP-specific antibody-secreting B cells increased 5–8-fold over 14 d in infected BALB/c mice [9], it might be expected that MD-4 cells might also be subject to similar polyclonal activation signals. This could provide an explanation for why we found that although at a population level, the contact time for MD-4 cells with endogenous T cells was similar to that expected for naïve MD-4 cells [17], our data showed a much greater degree of spread of contact time per incidence (up to 30 min) than previously observed, where no non-cognate contacts lasted more than 5 min. Alternatively, and with potentially similar consequences, it has also been proposed that bystander B cells may acquire BCR from antigen-activated B cells through membrane transfer [45]. The greater spread of contact times observed between granuloma T cells and MD-4 B cells in our studies made it difficult to obtain and apply a definitive threshold for cognate and non-cognate interactions when analyzing the immune B cell population. In addition we could...
Figure 9. B cells in granulomas make contacts with endogenous T cells. A. MD-4 and immune B cells were evaluated for contacts with DsRed+ cells in 28 granulomas imaged from two L. donovani-infected Tred mice. The duration of all individual contacts is shown as a separate symbol. B. The frequency of contacts of different duration is shown for MD-4 (open bars) and immune (grey bars) B cells; P<0.001. C. Single frames from Video S1 showing two immune B cells making short (top images) or long (bottom images) contacts with DsRed+ cells. Contact plot is shown below, where each box represents a contact period and adjacent boxes indicate where multiple DsRed+ cells are bound simultaneously. D. The total time each individual B cell (grey bars) spends interacting with any granuloma T cell during the imaging period (i.e. the accumulated total contact time) is shown. Note that due to interactions with multiple T cells, some B cells show an accumulated total contact time longer than the imaging time of 30 min. E. Distribution of contact times for each individual MD-4 (open bars) or immune (grey bars) B cell. Data are expressed as frequency of total incidences for each B cell that involved short (<3 min) or long (>3 min) contact with an endogenous DsRed+ cell. ***, p<0.001.

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not retrospectively confirm whether B cells engaging in longer contacts displayed a different phenotype or activation status. It is likely that this technical limitation will also apply to any study of B cells in the environment of the *Leishmania* infected mouse, irrespective of additional manipulations that might be employed to further define the transferred populations of B cells. The extent of polyclonal activation is thus an important consideration for future imaging-based studies in this or other models of infection where polyclonal activation may occur.

In conclusion, we have shown that mature B cells are non-selectively recruited into hepatic granulomas during *L. donovani* infection, with no enrichment of ‘regulatory’ B cells, and that some engage in long-lasting interactions with T cells. Our data therefore provide the first illustration of the dynamic nature of B cells at an inflammatory site, whilst also suggesting caution in the interpretation of histopathologic data on lymphocyte recruitment in the absence of measures of antigen-specificity.

**Methods**

**Ethics statement**

All experiments were approved by the University of York Animal Procedures and Ethics Committee and performed under UK Home Office license (‘Immunity and Immunopathology of Leishmaniasis’ Ref # PPL 60/3708).

**Mice and Infections**

6–12 week old female C57BL6 mice (Charles River, Margate, UK), congenic B6.CD45.1, CD19<sup>−/−</sup>TR26R<sup>−/−</sup>V<sub>α</sub>DSRed<sub>−/−</sub> mice (B<sup>green</sup>/Tred<sup>−/−</sup>), National Institute for Medical Research, UK) and VaDsRed.B (<sup>Tred<sup>−/−</sup> NIMR, UK) were used. Femurs from MD-4 mice [46] backcrossed onto B6.μMT background were a kind gift of Dr G. Kassiotis (NIMR, UK), *Leishmania donovani* (LV9) and a transgenic line expressing TdTOMATO [47] were maintained in B6.RAG1<sup>−/−</sup> mice. 2.3×10<sup>5</sup> amastigotes were injected intravenously (i.v.) to initiate infection [48].

**Intra-vital imaging**

Intra-vital imaging was performed as described [33]. Briefly, mice were anaesthetised with ketamine (100 mg/kg), xylazine (10 mg/kg) and acepromazine (1.7 mg/kg), the left liver lobe was exteriorised and the animal placed on a custom made imaging platform. Images were acquired on an inverted LSM 510 multiphoton microscope (Carl Zeiss Microimaging) using a 36°C blacked-out environmental chamber (Solenet Scientific, UK). For 4D analysis, 20–40 μM Z stacks were acquired with a Z distance of 2–3 μM approximately every 15–30 sec. Data were rendered and analysed using Volocity software (Improvision) and cell tracking performed manually, or automatically with manual checking. Cell-cell interactions were measured manually by determining the presence of cell-cell contacts in a 3D rendering on a frame-per-frame basis for each movie. The times of initial contact and the time that contact ended was recorded and the difference was calculated manually.

**Analysis of explant tissue**

Freshly removed liver tissue was placed in 35 mm cover-slip bottom Petri dishes (MatTek Corporation), and imaged with a 40×1.1 water immersion objective and fluorescence excitation provided by a Chameleon XR Ti:Sapphire laser (Coherent) tuned to 872 nm. Granuloma volumes were calculated by drawing regions of interest in Velocity to obtain a 3D volume measurement in μm<sup>3</sup>. Blocks mounted in OCT were stored at −80°C freezer until use. 10 μm cryosections of liver and spleen were stained using B220, CD35 and CD21 antibodies, and analysed using a Zeiss LSM 510.

**Flow cytometry and cytokine analysis**

Livers were perfused with ice-cold saline via the hepatic portal vein, and placed into ice-cold RPMI. Tissue was passed through a 100 μm strainer, washed twice (1400 rpm, 7 min), and hepatic mononuclear cells isolated on a 33% Percoll gradient. Hepatic mononuclear cells were resuspended in RPMI with 10% FCS, plated into 96-well plates (1–2×10<sup>5</sup> cells/well) and incubated in FACS buffer containing CD16/32 antibodies for 10 min on ice. After washing, cells were stained in 100 μl FACS buffer containing fluorochrome-labelled antibodies specific for IgM (R6-60-2), IgD (11-26c.2a), B220 (RA3-6B2), CD19 (1D3), CD11c (N418), TCR γδ (GL3) CD5 (53-7-3), CD1d (1B1), CD40 (1C10) or MHCII (M5/114.15.2) or appropriate isotype controls (all from eBioscience or BD Biosciences). For cytokine analysis, cells were plated into a 24-well plate (5×10<sup>5</sup> cells/well) with 50 ng/ml PMA, 500 ng/ml ionomycin and 10 μg/ml LPS at 37°C in 5% CO<sub>2</sub> for 3 hr. 10 μg/ml of Brefeldin A was added for a further 4 hr, before fixation in 2% PFA (10 min on ice), permeabilization (0.5% saponin) and staining IFNγ (XMG1.2), IL-2 (BVD6-24G2), IL-10 (JES5-16E3) and TNF (MP6-XT22). Labelled cells were analysed on a Cyan flow cytometer with Summit software (Beckman Coulter).

As required, B cells were isolated by positive selection using an LS MAGS column according to manufacturer’s instruction (Miltenyi), after incubation with biotin-conjugated anti-CD19 antibody and streptavidin-conjugated magnetic beads.

**B cell adoptive transfer**

MACS purified spleen cells or hepatic mononuclear cells were stained with FITC-conjugated anti-CD45.2 (as required) and PE-conjugated anti-CD19 and sorted on a MoFlo (Beckman Coulter). Purified B cells were resuspended (1–5×10<sup>5</sup> cells/ml) in either CFDA-SE (5 μM) or Hoechst 33342 (6 μM) for 8 min at 37°C followed by quenching with cold medium. After washing labelled B cells were resuspended in RPMI (8×10<sup>5</sup>–1×10<sup>6</sup> in 200 μl) and injected i.v., alone or in a 1:1 ratio.

**Real time –RT-PCR**

mRNA accumulation for cytokines was performed as described elsewhere [43]. Briefly, 1×10<sup>6</sup> sort-purified cells were re-suspended in RLT buffer and RNA isolated with RNeasy spin columns according to the manufacturer’s instructions (Qiagen). Eluted RNA was converted to cDNA and cDNA stored at −20°C or used immediately. The RT-PCR mix consisted of SYBR Green, forward primer (1 μM) and reverse primer (1 μM) and reactions were performed in MicroAmp Optical 96-well reaction plates using an ABI Prism 7000 RT-PCR system. RT-PCR data was analysed using the 7000 system SDS software.

**Statistical analysis**

Parametric or non-parametric tests were selected according to the distribution of the raw data. Comparisons between experimental groups were performed using students t test or Mann-Whitney. Correlations were determined using the Pearson correlation test and distribution data analyzed by chi-squared test. All analyses were conducted using GraphPad InStat (version 3) software (GraphPad software, San Diego, California, US).
Dotted line in histograms represents the B220/CD3 staining of the DsRed anti-CD3 and anti-B220 Ab. Dot plot shows gated eYFP + populations within the spleen and histograms their respective expression of B220 and CD3. Grey population in histogram represents eYFP + dsRed - cells from left hand panel. Dotted line in histograms represents the B220/CD3 staining of the total population shown in left hand panel. (TIF)

**Video S1** Dynamic behavior of adoptively transferred CFSE-labelled MD-4 cells (green) in granulomas of L. donovani-infected Tred mice, showing example of short and long contact (indicated by arrows) with endogenous T cells. Second harmonic is shown in blue. (MOV)

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