T Cell Hypo-Responsiveness against *Leishmania major* in MAP Kinase Phosphatase (MKP) 2 Deficient C57BL/6 Mice Does Not Alter the Healer Disease Phenotype

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Abstract

We have recently demonstrated that MAP kinase phosphatase 2 (MKP-2) deficient C57BL/6 mice, unlike their wild-type counterparts, are unable to control infection with the protozoan parasite *Leishmania mexicana*. Increased susceptibility was associated with elevated Arginase-1 levels and reduced iNOS activity in macrophages as well as a diminished Th1 response. By contrast, in the present study footpad infection of MKP-2−/− mice with *L. major* resulted in a healing response as measured by lesion size and parasite numbers similar to infected MKP-2+/+ mice. Analysis of immune responses following infection demonstrated a reduced Th1 response in MKP-2−/− mice with lower parasite specific serum IgG2b levels, a lower frequency of IFN-γ and TNF-α producing CD4+ and CD8+ T cells and lower antigen stimulated spleen cell IFN-γ production than their wild-type counterparts. However, infected MKP-2−/− mice also had similarly reduced levels of antigen induced spleen and lymph node cell IL-4 production compared with MKP-2+/+ mice as well as reduced levels of parasite-specific IgG1 in the serum, indicating a general T cell hypo-responsiveness. Consequently the overall Th1/Th2 balance was unaltered in MKP-2−/− compared with wild-type mice. Although non-stimulated MKP-2−/− macrophages were more permissive to *L. major* growth than macrophages from MKP-2+/+ mice, reflecting their reduced iNOS and increased Arginase-1 expression, LPS/IFN-γ activation was equally effective at controlling parasite growth in MKP-2−/− and MKP-2+/+ macrophages. Consequently, in the absence of any switch in the Th1/Th2 balance in MKP-2−/− mice, no significant change in disease phenotype was observed.

Citation: Schroeder J, McGachy HA, Woods S, Plevin R, Alexander J (2013) T Cell Hypo-Responsiveness against *Leishmania major* in MAP Kinase Phosphatase (MKP) 2 Deficient C57BL/6 Mice Does Not Alter the Healer Disease Phenotype. PLoS Negl Trop Dis 7(2): e2064. doi:10.1371/journal.pntd.0002064

Introduction

*Leishmania* species are protozoan parasites that are transmitted by infected female sandflies and cause a wide spectrum of diseases ranging from self-healing cutaneous lesions to fatal systemic disease. After entering their vertebrate host, promastigotes are taken up initially by neutrophils and ultimately macrophages and dendritic cells, where they turn rapidly into amastigotes and survive within parasitophorous vacuoles [1]. Resistance against cutaneous infection with *Leishmania (L.) major* typically requires the presence of an antigen-specific type 1 immune response comprising of IFN-γ/TNF-α/IL-2 producing CD4+ T [2,3] cells but also CD8+ T cells have been shown to play an important role in parasite clearance [4,5]. Subsequently, activated T cells migrate to the site of infection where they release IFN-γ and TNF-α which in turn upregulate inducible nitric oxide synthase (iNOS) in infected macrophages, enabling nitric oxide (NO) mediated killing of the intracellular parasites [6,7]. Susceptibility, on the other hand, has been associated with a failure to produce a type-1 response, which may be a consequence of IL-10 production from Fc-γ mediated macrophage uptake of amastigotes [8], or from natural and or type-1 regulatory T cells [9–11], or regulatory B cells [12], or an elevated Th2 response and the excessive production of IL-4 by CD4+ T cells [2,13], or indeed a combination of all these factors (reviewed by [14]). IL-4 in particular has been shown to promote alternative macrophage activation including increased expression of Arginase-1 [15], suppression of iNOS [16] and increased growth of *L. major*.

Mitogen-activated protein kinase phosphatase 2 (MKP-2) is a dual-specific nuclear phosphatase (DUSP) and is associated with the MAPK signalling pathway, where it has been shown to dephosphorylate and thereby inactivate protein kinases ERK, JNK but not p38 in vitro [17,18]. MKP-2 could therefore have significant effects on *L. major* infection as these parasites have a well recognized ability to subvert the development of Th1 responses partly via effects upon MAP kinase signalling. Studies using *L. major* metacyclic promastigotes indicated that the parasite via lipophosphoglycan (LPG) differentially regulated IL-12 as well as NO production by targeting ERK and p38 MAPK, respectively [19,20]. In order to better understand the role of MKP-2 in immune functions we, and others, have recently created MKP-2 deficient mice on a C57BL/6 background [21–24]. Thus regulatory roles specific for MKP-2 have been demonstrated in the inflammatory response associated with sepsis [22], cell cycle progression and apoptosis [23] and infection [21]. Furthermore, MKP-2−/− macrophages have severe ablation of LPS or IFN-γ -
induced iNOS expression and nitric oxide release and enhanced basal expression of Arginase-1. Given that Arginase-1 competes with iNOS for their common substrate L-arginine, it suggested that MKP-2 could have a regulatory function significant in immune responses involving intracellular pathogens [25]. Indeed, following infection with the intracellular parasite L. mexicana we demonstrated that it was changes in Arginase-1 and iNOS rather than changes in kinase mediated signalling that dictated the subsequent in vivo response to MKP-2 deletion [21]. MKP-2 deficiency mice displayed increased lesion size and parasite burden, and a significantly modified Th1/Th2 bias compared with wild-type counterparts [21]. This was related to a significant down-regulation of specific Th1 activity in MKP-2 deficient animals. However, there was no intrinsic defect in MKP-2/−/− T cell function as measured by anti-CD3 induced IFN-γ production. Rather, MKP-2/−/− bone marrow-derived macrophages, as a consequence of increased Arginase-1 expression, were found to be inherently more susceptible to infection with L. mexicana than MKP-2+/+ derived macrophages.

The immune-regulatory mechanisms controlling L. mexicana and L. major differ significantly and while most mouse strains heal following infection with L. major the vast majority develop chronic infections following infection with L. mexicana (reviewed by [26]). Thus, while C57BL/6 mice can control L. mexicana lesion growth they do not cure and MKP-2 deficiency results in progressive disease following L. mexicana infection [21]. C57BL/6 mice on the other hand are more resistant to L. major and lesions heal following infection. Given the well documented importance of NO killing in controlling L. major infections we addressed the question as to whether MKP-2 deficiency would make C57BL/6 mice more permissive host cells for L. major than wild-type macrophages, if classically activated they were equally effective at controlling parasite growth.

Materials and Methods

Ethics statement

Female DUSP-4 (MKP-2) wild-type and deficient mice were generated as previously described and bred onto C57BL/6 background [21] and female BALB/c mice were bred and maintained under specific pathogen-free conditions in the animal facilities of the Strathclyde Institute for Pharmacy and Biomedical Sciences at the University of Strathclyde. Animals were used at 6–9 weeks of age and were age matched within each experiment. All animal experiments adhered to the UK Animals (Scientific Procedures) Act 1986 and were conducted under Project Licenses to RP (PPL60/3439 “genetic models of cancer and inflammation”) and JA (PPL60/3929 “mechanism of control of parasite infection”) granted by the UK Home Office and with local ethical approval.

Parasite infections

Leishmania major (IR75) promastigotes were grown in HOMEM (Gibco) supplemented with 10% FCS (Biosera, East Sussex, UK) until late stationary phase. In order to enable direct comparison with previous studies on the role of MKP-2 during L. mexicana infections [21] and to Arginase-1/L. major studies [27,28] relevant to the work presented here we used the high dose inoculation model. Mice were given 2×10^6 L. major subcutaneously into the left hind foot pad and lesion development was monitored as the difference in thickness between infected and uninfected foot pads using a dial gauge calliper. For determination of parasite burden, mice were sacrificed by cerebral dislocation and foot pads, popliteal lymph nodes and spleens were removed and homogenized. Single cell suspensions were adjusted to equal volumes and subjected to limiting dilution assay as described elsewhere [29].

T cell assay

Single cell suspensions from spleens and lymph nodes of infected mice were prepared and erythrocytes were lysed using Erythrocyte lysis buffer pH 7.3 (160 mM NH₄Cl, 10 mMKHCO₃ and 100 µM EDTA). Cells were resuspended in RPMI containing 10% FCS, L-Glutamine and Penicillin/Streptomycin and 2×10^6 splenocytes were restimulated either on dendritic cells pulsed with 5 µg L. major antigen over night, PMA (10 ng/ml), ConA (10 µg/ml) or medium alone for 6 h (intracellular staining) or 48–72 h (ELISA) in 24-well plates. For intracellular staining, cytokine release was inhibited by the addition of BrefeldinA (10 µg/ml) at 3 h into re-stimulation. All reagents were supplied by Sigma-Aldrich (St. Louis, USA).

Magnet assisted cell sorting (MACS)

CD4^+ T cells from spleens and lymph nodes of infected mice were isolated on LS magnetic columns (Miltenyi Biotech GmbH, Bergisch Gladbach, Germany) using the negative selection kit (Miltenyi) following manufacturer’s instructions.

Enzyme linked immunosorbant assay (ELISA)

Mouse serum was analysed individually for antigen-specific IgG1 or IgG2b using single sided ELISA as previously described [30]. Resulting titres were expressed as reciprocal values of the half-maximal absorption at 450 nm using a Spectromax 190 plate reader.

Cytokines from T cell supernatants were determined with Sandwich-ELISA using cytokine specific capture and detection
antibodies (BD Bioscience) and recombinant cytokines for standard curves (R&D Systems, Minneapolis, USA). IL-4 was determined using the mouse IL-4 QuantiKine kit (R&D Systems) following manufacturer’s instructions. To allow detection of IL-4, 5 μg/ml IL-4 receptor blocking antibody (anti-mouse CD124, BD Pharmingen, USA) was added to the cultures.

Flow cytometry

Cells were harvested and passed through a nitex mesh to remove clumps. For isotype control, 5 × 10^5 cells of medium-, antigen- and PMA/IONomycin-reconstituted splenocytes were pooled for each individual. After blocking unspecific binding with 10% mouse serum and Fc receptor blocking antibodies (anti-mouse CD16/32, eBioscience, UK), cell surface was stained with conjugated antibodies for CD3e (PerCP, CD4 (APC-H7), both BD Pharmingen (USA) and CD8b (Alexa Fluor 488, eBioscience UK) for 45 min at 4°C. Cells were fixed for 15 min and permeabilized using the Fix and Perm kit (Invitrogen, Paisley, UK). Intracellular staining was carried out simultaneously with permeabilisation for 1 h at room temperature using conjugated antibodies for IFN-γ (Allophycocyanin (APC)) and TNF-α (Phycerythrin (PE)) or their respective isotypes anti-rat IgG1 (APC) and anti-rat IgG1 (PE), all eBioscience. After washing steps, cells were resuspended in 200 μl PBS and subsequently run and analysed on the FACS Canto flow cytometer (BD Bioscience) using FACS Diva software.

Cell culture

Bone marrow-derived macrophages (BMDM) and dendritic cells were derived from mice of 6 to 8 week old mice. Bone marrow was flushed, and in order to obtain macrophages, cells were resuspended in DMEM containing 10% foetal calf serum (FCS), 30% L929-conditioned medium, 2 mM L-Glutamine (Gibco, Invitrogen, Paisley, UK), 1% Penicillin/Streptomycin (Gibco) and seeded into Petri dishes. After 10 days at 37°C, adherent macrophages were harvested with cold/warm PBS, washed and resuspended in complete RPMI (10% FCS, 2 mM L-Glutamine, 1% Penicillin/Streptomycin). Dendritic cells were generated from bone marrow precursors by culturing in RPMI containing 10% FCS, 2 mM L-Glutamine, 1% Penicillin/Streptomycin and 2.5–10% GM-CSF conditioned medium (X63). Non-adherent cells were harvested at day 7, washed and resuspended in complete RPMI.

Macrophage infections

BMDMs were seeded onto cover slips at 2 × 10^5 cells/500 μl in 24-well plates and left to adhere at 37°C over night. For infection L. major promastigotes (IR75) were added at a multiplicity of infection (MOI) of 5 and plates were briefly spun at 300 xg to allow close proximity of parasite and macrophages. After 1 h at 34°C, supernatants were removed and macrophages were washed in PBS to remove any free parasites. Fresh complete RPMI supplemented with or without 100 ng/ml LPS, 100 U/ml IFN-γ or 100 U/ml IL-4 was added and cells were incubated at 34°C for different periods of time.

To determine number of parasites and infection rates, macrophages were fixed in methanol and stained with Giemsa solution. Cover slips were mounted onto glass slides and intracellular parasites were counted in a total of 200 macrophages across each cover slip using a bright field microscope.

Immunoblotting

Arginase-1 expression was determined from whole cell lysates of 1 × 10^6 L. major-infected BMDM. SDS-PAGE analysis and detection was carried out as described elsewhere [21].

Statistics

Statistical analysis was performed using GraphPad Prism Program (Version 4.0, GraphPad Software, San Diego, California). P values below or equal to 0.05 were considered significant.

Results

MKP-2 deficiency in C57BL/6 mice did not alter the course of infection with L. major

MKP-2 deficient and MKP-2 wild-type C57BL/6 mice were given 2 × 10^5 late-stationary phase L. major promastigotes subcutaneously into the left hind footpad and lesion development was monitored over 12–13 weeks. Surprisingly, given that MKP-2−/− mice were previously found to be more susceptible to L. mexicana than their wild-type counterparts [21], no difference in lesion growth between MKP-2−/− and MKP-2+/+ mice was detected throughout the course of infection with L. major. Both MKP-2−/− and MKP-2+/+ mice developed lesions which healed spontaneously after several weeks and in a manner typical for L. major resistant C57BL/6 mice (Figure 1A). At two time points, the peak of infection (Figure 1B) and after onset of healing (Figure 1C), five mice were sacrificed and the parasite burdens of infected footpads, popliteal draining lymph nodes and spleens were determined. Consistent with the lesion development, parasite numbers were high at the peak of infection and consequently dropped with the onset of healing. We could not observe a difference in phenotype between wild-type and MKP-2−/− mice at either peak lesion growth or after healing in any of the tissues examined. This observation was made in several independent experiments.

Enhanced susceptibility of MKP-2−/− macrophages to infection with L. major is ablated following classical but not innate or alternative activation

Our previous studies demonstrated that the susceptibility of MKP-2 deficient mice is to a significant degree due to MKP-2−/− macrophages being inherently more susceptible to the growth of L. mexicana than wild-type macrophages a dichotomy that was maintained even following classical activation [21]. We therefore examined the infectivity of L. major metacyclic promastigotes for MKP-2−/− and MKP-2+/+ naïve, innately, classically or alternatively activated bone marrow derived macrophages (Figure 2A, B and C). Over several experiments we found that infected non-stimulated MKP-2−/− BMDM cultures maintained significantly increased parasite burdens and increased frequency of infected macrophages than cultures using their wild-type counterparts (Figure 2A). Our previous studies with L. mexicana had demonstrated the increased susceptibility of non-stimulated MKP-2−/− macrophages to infection to be related to elevated Arginase-1 levels compared with MKP-2+/+ macrophages and similarly enhanced levels of Arginase-1 were found in MKP-2−/− macrophages following infection with L. major promastigotes (Figure 2E). Innate macrophage activation with 100 ng/ml LPS greatly decreased the number of intracellular parasites as well as the frequency of infected cells in both MKP-2−/− and MKP-2+/+ macrophages (Figure 2A). However, despite activation with LPS, MKP-2 deficient macrophages still presented higher parasite survival than their wild-type counterparts. As nitric oxide (NO) is essential for killing of intracellular parasites, we determined the production of NO by measuring nitrite levels in the supernatants of infected macrophages 24 and 48 h after activation with LPS. Consistent with our previous observations with L. mexicana, MKP-2−/− macrophages showed a reduced NO production at both time points in response to LPS activation (Figure 2D). However,
whereas MKP-2−/− macrophages classically activated with LPS and IFN-γ failed to reduce *L. mexicana* parasite burdens to the level of similarly treated MKP-2+/+ macrophages [21] such treatment sufficed following infection with *L. major* promastigotes to ablate any differences in susceptibility between MKP-2−/− and MKP-2+/+ host cells (Figure 2B) and corresponded to LPS and IFN-γ treatment abating the deficiency in NO production in MKP-2−/− macrophages infected with *L. major* (data not shown). The addition of recombinant IL-4 to macrophages infected with *L. major* doubled the numbers of intracellular parasites in both wild-type and MKP-2−/− macrophages (Figure 2C), highlighting the disease-promoting attributes of IL-4 and alternative macrophage activation in *L. major* infections characterized by elevated Arginase-1 and reduced iNOS expression [16,31].

**Figure 1.** MKP-2 deficiency does not alter the course of infection with *L. major*. Ten MKP-2−/− (open symbols) and ten MKP-2+/+ mice (closed symbols) were given with 2×10^6 *L. major* promastigotes subcutaneously into the left hind foot pad. The right uninfected foot pad was used for reference. Lesion development was monitored by measuring the footpad thickness (A) over time using a dial gauge calliper. At peak of infection (day 42, B) and after the onset of healing (day 88, C) five mice were sacrificed at random and parasite burdens in foot pad, popliteal lymph nodes and spleens were determined using a limiting dilution assay. Detection limit was 65 for spleens and 6500 for lymph node and foot pads. Error bars are shown as standard error of the mean (SEM). The data shown is representative of at least three independent experiments.

doi:10.1371/journal.pntd.0002064.g001

MKP-2 deficiency results in a diminished Th1 response following infection with *L. major*

As no discernible change in the healing disease phenotype was observed between MKP-2−/− and MKP-2+/+ mice infected with *L. major*, and given that classically activated macrophages from MKP-2−/− mice were equally as effective as their wild-type counterparts at killing parasites we compared the type-1 response generated in MKP-2+/+ and MKP-2−/− following infection. After re-stimulation with soluble *L. major* antigen-pulsed DCs, flow cytometric analysis of CD4+ T cells from the spleens of infected animals demonstrated that MKP-2−/− mice had a significantly lower frequency of IFN-γ single producing and IFN-γ/TNF-α double producing CD4+ and CD8+ T cells (P<0.05) than their wild-type counterparts (Figure 3A, B and C). Similarly significantly reduced IFN-γ production (P<0.05) was measured in the supernatants of MKP-2−/− derived T cell cultures after stimulation with *L. major* antigen-pulsed DC, PMA/Ionomycin or ConA (Figure 3D). Finally we analysed the serum of *L. major* infected mice for the presence of antigen-specific IgG2b, a surrogate marker for Th1 responses, as IFN-γ is essential for IgG class switch to 2a and 2b [32,33]. At both stages of infection, MKP-2 deficient mice showed lower IgG2b titres, being statistical significant (P<0.05) at day 42 post-infection (Figure 3E). Overall, our data suggest that MKP-2 deficiency results in a diminished Th1 response during infection with *L. major*, similar to our previous observations using *L. mexicana* infections [21].

**MKP-2 deficiency results in a reduced Th2 response in *L. major* infected mice**

In stark contrast to experimental infection with *L. mexicana*, MKP-2 deficient mice did not show an increased susceptibility to infections with *L. major* compared with wild-type mice. This was despite a clearly diminished Th1 response in MKP-2−/− compared with MKP-2+/+ mice following *L. major* infection (Figure 3). However, MKP-2 is a generalized negative regulator
of Arginase-1 tissues rich in phagocytes (Figure S1) and it is well
documented that Arginase-1 can induce hypo-responsiveness in a
pan-T cell fashion [27,28] [34]. We therefore examined whether
the lack of a parasite-specific Th1 response could be counterbal-
ced by concomitant impairment of a Th2 response, leaving the
relative Th1/Th2 balance undisturbed. Over a number of
experiments we found MKP-2<sup>−/−</sup> mice infected with
<i>L. major</i> produced less IgG1 but not significantly less than their similarly
infected wild-type counterparts (Figure 4A). To directly measure
the parasite induced production of IL-4, we infected MKP-2 wild-
type and deficient mice as well as susceptible BALB/c mice with
<i>L. major</i> promastigotes. At the peak of infection, mice were sacrificed,
spleens and draining lymph node cells were removed and analyzed for
the production of IL-4. Whole spleen and lymph node cells were re-
stimulated with soluble <i>L. major</i> antigen-pulsed DC, PMA/
Ionomycin or ConA for 72 h in the presence of IL-4 receptor
blocking antibody to prevent immediate uptake of freshly
produced IL-4 in an auto- or paracrine fashion. As expected,
susceptible BALB/c mice produced high levels of IL-4 compared
with mice from the C57BL/6 background (Figure 4B). However,
while antigen induced IL-4 production was reduced in MKP-2<sup>−/−</sup>
splenocytes and draining lymph node cells (Figure 4C and D)
compared with wild-type equivalent cell populations this was not
clearly significant because of small sample sizes. For a more precise
examination of the Th2 response we therefore isolated CD4<sup>+</sup>
T cells from spleen suspensions by negative selection and re-
stimulated these as described above. The levels of IL-4 produced
by CD4<sup>+</sup> T cells in response to <i>L. major</i> infection were indeed
clearly and significantly reduced in MKP-2 deficient mice when
compared with wild-type mice (Figure 4E), thus confirming a
reduced Th2 response in infected MKP-2<sup>−/−</sup> mice.

The Th1/Th2 balance in MKP-2 deficient mice compared
with wild type mice is sustained by general T cell hypo-
responsiveness against <i>L. major</i>

Previous studies have suggested that it is not the quantity
of IFN-γ or IL-4 production that is important in determining

Figure 2. Macrophages of MKP-2<sup>−/−</sup> mice are inherently more susceptible to <i>L. major</i> parasites. Bone marrow-derived macrophages of
MKP-2<sup>−/−</sup> (open symbols) and MKP-2<sup>+/+</sup> (closed symbols) grown on cover slips were infected with <i>L. major</i> promastigotes at a multiplicity of infection
(MOI) of 5. One hour after infection free parasites were removed by washing and medium was replaced with complete RPMI supplemented with or
without 100 ng/ml LPS or a combination of 100 ng/ml LPS and 100 U/ml IFN-γ, or 100 U/ml IL-4 and left for 24 or 48 h at 34 °C (A, B and C
respectively). Macrophages were fixed in methanol and stained with Giemsa. Intracellular parasites were counted in a total of 200 macrophages using
a bright field microscope and are shown as number of parasites per 100 macrophages. The mean percentage of infected macrophages is also
expressed as numbers inside the columns. At time points indicated supernatants from LPS-stimulated macrophages were removed and tested for
nitric oxide production using Griess reagent (D). Error bars show standard error of the mean (SEM). The data shown is representative of at least three
independent experiments. For analysis of Arginase-1 expression 1 x 10<sup>6</sup> BMDM were infected at an MOI of 5 and whole cell lysates were run on 12% gels, blotted on nitrocellulose membranes and stained for Arginase-1. The top cross-reactive band serves as loading control (E).
doi:10.1371/journal.pntd.0002064.g002
Figure 3. MKP-2 deficient mice infected with *L. major* have a reduced Th1 response. Splenocytes from *L. major* infected MKP-2−/− (open columns) and MKP-2+/+ (closed columns) mice were re-stimulated either with antigen pulsed DC, PMA/Ionomycin, ConA or with unpulsed DC (Medium control) for 6 h (FACS) or 48 h (ELISA). Cells were stained for T cell surface markers CD3, CD4 and CD8 as well as intracellular cytokines IFN-γ and TNF-α or their respective isotypes (A). Percentages of IFN-γ single and IFN-γ/TNF-α double producing parasite-specific CD4+ (B) and CD8+ (C) T cells are shown after normalization to medium and isotype controls. Supernatants of 48 h T cell re-stimulations were analyzed for IFN-γ in a sandwich ELISA and values were normalized to medium controls (D). Serum of infected mice was collected and analysed for parasite-specific antibody isotype IgG2b. The titres were calculated as reciprocal dilution of the half-maximal absorption at 450 nm (E). Error bars are shown as standard error of the mean (SEM). *P<0.05, **P<0.01; one-tailed Mann Whitney U test. The data shown is representative of two independent experiments.
doi:10.1371/journal.pntd.0002064.g003
protective immunity to *L. major* but rather the overall Th1/Th2 balance [35,36]. Consequently we calculated the relative Th1/Th2 balance between *L. major* infected MKP-2<sup>−/−</sup> and MKP-2<sup>+/+</sup> mice. In the first instance we calculated the ratio of IgG2b to IgG1 for each individual mouse, as this is a strong indicator of Th1/Th2 balance [37–39]. However no difference in IgG2b/IgG1 levels between *L. major* infected MKP-2<sup>−/−</sup> and MKP-2<sup>+/+</sup> mice were observed throughout infection (Figure 5A). Not surprisingly, the ratio increased after onset of healing toward an IgG2b bias and thus a Th1 response in both MKP-2<sup>−/−</sup> and MKP-2<sup>+/+</sup> mice. We further calculated the percentage reduction in the levels of type-1 and type-2 cytokine responses induced in MKP-2<sup>−/−</sup> compared with wild-type mice following infection. MKP-2<sup>−/−</sup> splenocytes from *L. major* infected mice produced, upon restimulation with *L. major* antigen-pulsed DCs, 32.6% less IFN-γ (Figure 5B) than MKP-2<sup>+/+</sup> splenocytes (20.79±5.53 (SD) and 30.87±5.39 (SD) ng/ml, respectively). Under the same experimental conditions a 37.8% reduction in IL-4 production (Figure 5B) was generated by
MKP-2$^{-/-}$ splenocytes compared with MKP-2$^{+/+}$ splenocytes (35.46±9.32 (SD) pg/ml compared with 56.99±12.13 (SD) pg/ml, respectively). Moreover, when comparing the percentage of IFN-γ/TNF-α double producing CD4$^+$ T cells, as measured by flow cytometry, we found a 28.7% reduction (Figure 5B) in infected MKP-2$^{-/-}$ compared with MKP-2$^{+/+}$ mice. Thus MKP-2 deficiency results in a T cell hypo-responsiveness following L. major infection, which effects both the protective T$_H1$ and the disease-exacerbating T$_H2$ response to a similar degree (Figure 5C).

Consequently, MKP-2$^{-/-}$ mice have a healing phenotype similar to their wild-type counterparts.

![Figure 5. A sustained T$_H1$/T$_H2$ balance in MKP-2 deficient mice is due to general T cell hypo-responsiveness against L. major.](image)

**Discussion**

In a previous study we identified a major role for MKP-2 in protecting C57BL/6 mice from infection with L. mexicana as MKP-2 deficiency resulted in uncontrolled lesion growth with massively increased parasite burdens [21]. Increased susceptibility was associated with MKP-2$^{-/-}$ macrophages being inherently more susceptible than wild-type macrophages to parasite infection as a result of increased Arginase-1 expression and also reduced NO production [21]. In addition, while no specific direct defect in T cell function could be attributed to MKP-2 deficiency, a diminished parasite-specific T$_H1$ and an enhanced T$_H2$ response developed in MKP-2$^{-/-}$ mice infected with L. major [21]. Surprisingly therefore, in the present study, no differences whatsoever in the normal healing phenotype were observed between MKP-2$^{-/-}$ and MKP-2$^{+/+}$ mice infected with L. major. This was despite firstly, naive and innately activated MKP-2$^{-/-}$ macrophages being more permissive to L. major infection than wild-type macrophages which, as with L. mexicana infections, was associated with increased Arginase-1 and reduced iNOS activities, and secondly, a clearly reduced parasite-specific T$_H1$ response in infected MKP-2$^{-/-}$ mice.

Infection of MKP-2$^{-/-}$ mice and their host cells with L. major differed to that of infection with L. mexicana in two significant ways that undoubtedly had profound effects on disease outcome. Firstly, in contrast to infection with L. mexicana, infection of MKP-2$^{-/-}$ mice with L. major resulted not only in a reduced type-I response, but also in an equally reduced T$_H2$ response compared with wild-type mice. Consequently there was no discernible difference in the T$_H1$/T$_H2$ bias between MKP-2$^{-/-}$ and MKP-2$^{+/+}$ mice infected with this parasite. It is well established in C57BL/6 mice, that once parasite peptide reactive-CD4$^+$ and CD8$^+$ T cell populations reach the proper balance in draining lymph nodes and the sites of infection, there is rapid healing, and immunity is maintained by a persistent small amastigote population in equilibrium with both effector and regulatory T cell populations [10,40]. Furthermore studies in mice have demonstrated that the absolute amount of IFN-γ generated following infection with L. major did not correlate with protection or cure [35,36] and rather, it was the balance in the T$_H1$/T$_H2$ cytokine profile that was important in determining disease outcome. Secondly, classically activated MKP-2$^{-/-}$ macrophages were equally as effective as MKP-2$^{+/+}$ macrophages in controlling the growth of L. major although not, as previously demonstrated, L. mexicana [21]. Therefore, as the T$_H1$/T$_H2$ balance remained unaltered, albeit equally diminished, in L. major infected MKP-2$^{-/-}$ compared with MKP-2$^{+/+}$ mice, the comparative polarisation of classical macrophage activation would remain the same. Consequently healing takes place in MKP-2$^{-/-}$ mice infected with L. major in a similar manner to infected resistant wild-type mice.

Among the intriguing questions regarding the differential outcome of L. major and L. mexicana infections in MKP-2 deficient mice is why T$_H2$ responses are maintained following L. mexicana infection [21] but down-regulated following L. major infection compared with their infected wild-type counterparts? It has been demonstrated previously that during L. major infections high local Arginase-1 levels at the site of infection mediate L-arginine depletion, which results in impaired local CD4$^+$ (and CD8$^+$) T cell function, particularly IFN-γ production but also to a lesser extent IL-4 and IL-10 [28,41]. Investigations carried out by Kropf et al. [34] using Arginase-1 expressing placenta cells and Jurkat T cells showed that Arginase-1-mediated T cell hypo-responsiveness is a consequence of the down-regulation of the CD3$\gamma$ chain, a crucial signal transducing component of the TCR [34] and significantly.

**Figure 5. A sustained T$_H1$/T$_H2$ balance in MKP-2 deficient mice is due to general T cell hypo-responsiveness against L. major.** Antibody titres for IgG2b were divided by respective titres for IgG1 for each individual mouse and for both time points. Higher values are indicative for higher IgG2b levels and lower values reflect a higher IgG1 response. The bar indicates the median of each group (A). The percent reduction of the mean for each measured parameter (IFN-γ, IL-4 and IFN-γ/TNF-α CD4$^+$ T cells) of MKP-2$^{-/-}$ mice was calculated against the mean values of MKP-2$^{+/+}$ mice (=100%) (B). If both immune responses (T$_H1$, IFN-γ/TNF-α CD4$^+$ T cells and IFN-γ) versus T$_H2$ (IL-4) are reduced to a similar extend (B), the balance between disease promoting T$_H2$ and the protective T$_H1$ response is not disturbed. Since the mice are on a C57BL/6 background the T$_H1$ response will outweigh the T$_H2$ response and consequently result in a healing phenotype (C).

doi:10.1371/journal.pntd.0002064.g005
CD3ε has recently been found to be down-regulated on CD4+ and CD8+ T cells along with increased Arginase-1 activity in the lesions of patients infected with L. amazonensis [42]. Depletion of L-arginine, as a result of the generalised elevated Arginase-1 levels in MKP-2−/− mice would explain the general T cell hypo-responsiveness observed in these mice following infection with L. major. However, why should the T olig response be maintained if not enhanced following infection of MKP-2−/− mice with L. mexicana complex and provide one possible explanation as to why these parasites, unlike L. major (reviewed by [43]) tend to induce chronic lesions in the majority of mouse strains such as the C57BL/6 strain used in this study. Not only are L. mexicana CPBs potent inhibitors of T olig responses [44] as a consequence of disrupting macrophage signalling pathways [45], but they have also been shown to be potent inducers of IL-4 production and T olig responses [46]. Consequently C57BL/6 mice infected with L. mexicana CPB null mutants, unlike infection with wild-type parasites, develop a healing response with reduced IL-4 production and T olig responses along with elevated T olig responses [47]. Thus, as a result of their highly expressed T olig promoting and MKP-2−/− mice with L. mexicana, unlike infection with L. major, is able to rescue and promote the T olig component of the parasite specific T cell response which is less subject to Arginase-1 induced hypo-responsiveness than T olig responses [27].

Alternative activation of both MKP-2−/− and MKP-2+/+ macrophages resulted in increased susceptibility to L. major. This would support the view that the T olig/T olig balance is important and that IL-4 and IFN-γ activities regulate each other not just at the T cell level but also at the level of the macrophage by modulation of iNOS and Arginase-1 expression. In agreement IL-4Rα signalling via macrophages/neutrophils has been shown to promote early lesion growth in L. major infected BALB/c mice and macrophage/neutrophil specific (LysM−/−IL-4Rα+/lox) IL-4Rα−/− mice display delayed lesion growth [16]. The control of L. major early in infection and MKP-2−/− mice has been identified as being due to enhanced macrophage microbial NO mediated activity in the absence of alternative macrophage activation. Paradoxically we failed to identify any disease promoting contributory role for IL-4Rα signalling via macrophages/neutrophils during L. mexicana infection [46]. In agreement in the present studies we also failed to identify a disease promoting role for IL-4 in either MKP-2−/− or MKP-2+/+ macrophages infected with L. mexicana (Figure S2). Thus the interaction of L. major and L. mexicana with their host macrophages must be significantly different. What may be critical in this regard is that L. amazonensis parasites, which belong to the “mexicana” complex of parasites, have been shown to be more resistant to macrophage-mediated control than L. major requiring higher levels of NO to induce killing [49,50]. The evidence would suggest that macrophage killing of “mexicana” complex parasites unlike L. major requires NO and additionally superoxide [51]. Furthermore, recent evidence indicates that, unlike L. major, there is in fact enhanced replication of the amastigote stage of L. amazonensis in IFN-γ-stimulated murine macrophages despite higher NO production [52], reportedly due to the induction of a novel L-arginine pathway independent of iNOS or host Arginase-1 [53]. Induction of Arginase-1 by L. amazonensis has also been shown to enhance replication of the amastigote stage of the parasite [52,53] while inhibition studies have shown that the enhanced susceptibility of MKP-2−/− macrophages for L. mexicana is associated with enhanced Arginase-1 expression [21]. Thus given the relatively higher sensitivity of L. major to NO matched with the increased importance of Arginase-1 to infection with L. mexicana it is perhaps not surprising that classical macrophage activation ablates MKP-2 deficiency mediated differences in infectivity with the former but not the latter parasite.

Overall the present study confirms our previous observation that MKP-2 is a major factor in determining the immune response against intracellular parasites and potentially the outcome of infection. Naive MKP-2 deficient macrophages are inherently more susceptible to L. major than wild-type macrophages and following infection there is a generalised T cell hypo-responsive-ness. However, despite these apparent deficiencies the disease phenotype of MKP-2−/− mice following L. major infection does not differ from wild-type mice. Our results suggest that this is a consequence of the T olig/T olig balance remaining unaltered in MKP-2−/− mice infected with L. major and that classical macrophage activation suffices to ablate the innate permisiveness of MKP-2−/− macrophages to this species.

Supporting Information

Figure S1 Arginase-1 levels are elevated in spleen and intraperitoneal exudates of MKP-2 deficient mice. Equal concentrations of whole cell lysates of intraperitoneal (IP)-washes (left panel) and spleen (right panel) of three mice were run on 12% gels, blotted on nitrocellulose membranes and stained for Arginase-1. Naïve MKP-2 deficient mice were run in the same gel. The control of MKP-2−/− and MKP-2+/+ bone marrow-derived macrophages from C57BL/6 mice infected with L. major was used as a coomparative standard of Arginase-1 expression. A Table S1 listing all the primers used in the study is provided in the supplementary information. (TIF)

Figure S2 External addition of IL-4 does not drastically increase intracellular growth of L. mexicana. Bone marrow-derived macrophages of MKP-2−/− (open columns) and MKP-2+/+ (closed columns) mice grown on cover slips have been infected with L. mexicana promastigotes at a multiplicity of infection (MOI) of 5. One hour after infection free parasites have been washed off and medium was replaced with complete RPMI supplemented with or without 100 U/ml IL-4 and incubated for 48 h at 34°C. Macrophages were fixed in methanol and stained with Giemsa. Intracellular parasites were counted in a total of 200 macrophages using a bright field microscope and are shown as number of parasites per 100 macrophages. The mean percentage of infected macrophages is also expressed as numbers inside the graphs. Error bars show standard error of the mean (SEM). (TIF)

Acknowledgments

The authors would like to thank Carol Whitehouse and Dr. Owain Millington for their help with the animal studies.

Author Contributions

Conceived and designed the experiments: JS RP JA. Performed the experiments: JS HAM SW JA. Analyzed the data: JS HAM JA. Wrote the paper: JS RP JA.
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