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**Recommended Citation**

Kannan, Yashaswini; Perez-Lloret, Jimena; Li, Yanda; Entwistle, Lewis J.; Khoury, Hania; Papoutsopoulou, Stamatia; Mahmood, Radma; Mansour, Nuha R.; Huang, Stanley Ching-Cheng; Pearce, Edward J.; de Carvalho, Luiz Pedro S.; Ley, Steven C.; and Wilson, Mark S., "TPL-2 regulates macrophage lipid metabolism and M2 differentiation to control TH2-mediated immunopathology." *PLoS Pathogens*. 12,8. e1005783. (2016).

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RESEARCH ARTICLE

TPL-2 Regulates Macrophage Lipid Metabolism and M2 Differentiation to Control T\textsubscript{H}2-Mediated Immunopathology

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Abstract

Persistent T\textsubscript{H}2 cytokine responses following chronic helminth infections can often lead to the development of tissue pathology and fibrotic scarring. Despite a good understanding of the cellular mechanisms involved in fibrogenesis, there are very few therapeutic options available, highlighting a significant medical need and gap in our understanding of the molecular mechanisms of T\textsubscript{H}2-mediated immunopathology. In this study, we found that the Map3 kinase, TPL-2 (Map3k8; Cot) regulated T\textsubscript{H}2-mediated intestinal, hepatic and pulmonary immunopathology following \textit{Schistosoma mansoni} infection or \textit{S. mansoni} egg injection. Elevated inflammation, T\textsubscript{H}2 cell responses and exacerbated fibrosis in \textit{Map3k8}\textsuperscript{-/-} mice was observed in mice with myeloid cell-specific (LysM) deletion of \textit{Map3k8}, but not CD4 cell-specific deletion of \textit{Map3k8}, indicating that TPL-2 regulated myeloid cell function to limit T\textsubscript{H}2-mediated immunopathology. Transcriptional and metabolic assays of \textit{Map3k8}\textsuperscript{-/-} M2 macrophages identified that TPL-2 was required for lipolysis, M2 macrophage activation and the expression of a variety of genes involved in immuno-regulatory and pro-fibrotic pathways. Taken together this study identified that TPL-2 regulated T\textsubscript{H}2-mediated inflammation by supporting lipolysis and M2 macrophage activation, preventing T\textsubscript{H}2 cell expansion and downstream immunopathology and fibrosis.
Author Summary

Chronic helminth infections can cause significant morbidity and organ damage in their definitive mammalian hosts. Managing this collateral damage can reduce morbidity and preserve vital tissues for normal organ function. One particular consequence of some chronic helminth infections is the deposition of fibrotic scar tissue, following immune responses directed towards helminth material. In this study we tested the role of a particular signalling kinase, TPL-2, and identified that it critically regulated the magnitude of fibrotic scarring following infection. Using several murine models with genetic deletions of TPL-2 in either all cells or specific deletion in subsets of immune cells (Map3k8–/–Map3k8fl/fl) we identified that expression of TPL-2 in myeloid cells was essential to prevent severe immune-mediated pathology. Using genome-wide analyses and metabolic assays, we discovered that TPL-2 was required for normal lipid metabolism and appropriate activation of myeloid cells/macrophages to limit fibrosis. These results revealed a previously unappreciated role for TPL-2 in preventing severe pathology following infection. Thus, activating this pathway may limit immune mediated pathology following chronic helminth infection. More broadly, this pathway is being targeted to treat inflammatory diseases and cancer [1, 2]. This study would suggest that caution should be taken to prevent untoward co-morbidities and fibrosis-related pathologies in patients when targeting TPL-2.

Introduction

Immune-mediated pathologies and fibrotic scarring are a major cause of global morbidity and mortality. This is due, in part, to a shortage of available drugs and a lack of novel therapeutic targets to limit fibrogenesis, highlighting a major unmet medical need [3, 4]. Chronic infection resulting in recurring inflammation and wound repair can lead to tissue remodelling, fibrosis and ultimately organ failure. Infection with the parasitic blood fluke, Schistosoma mansoni, can cause severe intestinal and hepatic pathologies caused by fibrotic lesions surrounding trapped parasite material. Parasite eggs become lodged within vascularised tissue invoking a distinctive eosinophil and macrophage (MΦ)-rich type-2 immune-mediated granuloma [5]. Th2-cell derived IL-4 and IL-13 [6] stimulate IL-4 receptor (IL-4R)-expressing MΦ’s [7, 8] to develop an M2 or alternative activation (AA) state characterised by expression of Arginase (Arg1), Resistin-like molecule alpha (Retnla, Fizz-1) and chitinase-like molecules (Chi3l3, Chi3l4) [9]. Animal models have indicated that IL-4R-dependent M2-MΦ’s are essential to 1) prevent fatal intestinal damage and sepsis following schistosome infection [7]; 2) orchestrate tissue remodelling and fibrotic responses [10–12] and 3) regulate Th2 cell proliferation and activation [13–15]. Despite the clear and well-documented importance of M2-MΦ’s during schistosome infection and the resulting immune-mediated protection, pathology and regulation, the critical regulatory proteins that control M2-MΦ differentiation are poorly understood.

The MAP3 kinase, TPL-2 (also known as COT and encoded by Map3k8) is ubiquitously expressed, phosphorylating and activating MEK1/2 following stimulation of Toll-like receptors (TLRs) and the receptors for TNF and IL-1β, leading to the activation of ERK1/2 MAP kinases [16]. TPL-2 is required for Th1 and Th17-associated inflammation and is essential for the development of autoimmunity and immunity to bacterial and protozoan pathogens [17–20]. In MΦ’s, TPL-2 is required for the synthesis and secretion of a variety of cytokines and chemokine’s following classical activation (CA) with TLR ligands [17, 20–29]. However, it is unclear...
whether TPL-2 controls M2-MΦ function to regulate chronic T\textsubscript{H}2-associated inflammation and immunopathology.

Two distinct inflammatory pathways contribute to fibrogenic responses; classical, pro-inflammatory type-1/17 and TGF-β-mediated fibrosis [30] and type-2 inflammatory pathways leading to IL-4R-dependent fibrosis [4]. It was recently reported that TPL-2–deficient mice, or inhibition of ERK [31], protected mice from type-1/TH17 and TGF-β-mediated pulmonary fibrosis following bleomycin treatment [31] and from hepatic fibrosis following carbon tetrachloride and methionine choline-deficient diet-induced fibrosis [32]. As expected, Map3k8–deficient Kupffer cells had reduced TLR-induced IL-1β and pro-fibrotic gene expression, which the authors suggested was responsible for the reduced hepatic fibrosis in vivo. However this was not directly tested. Nevertheless, this study raised the possibility that targeting TPL-2 may forestall the progression of hepatic fibrosis. Indeed many small molecule inhibitors have been developed that block TPL-2 signalling in vitro [2], but none have yet made it into the clinic. However, it is not known whether TPL-2 contributes to chronic type-2 inflammation and IL-4R-mediated fibrosis.

In this study, we used the well-established Schistosoma mansoni infection model to test whether TPL-2 regulated chronic type-2 associated inflammation, immunopathology and fibrosis. In contrast to the reduced fibrosis observed in Map3k8\textsuperscript{−/−} mice following chemical and diet-induced fibrosis [32], Map3k8\textsuperscript{−/−} mice had significantly increased type-2 immune responses with concomitant elevated inflammation and fibrosis surrounding trapped parasite eggs. Using genome-wide transcriptional analysis and metabolic assays we found that TPL-2 was required for lipid oxidative metabolism and M2-MΦ activation. Specifically, TPL-2 was required for expression of immunoregulatory molecules (Retnla and Arg1) and regulated pro-fibrotic genes (Col genes and Ctgf). Consequently, myeloid cell-specific deletion of Map3k8 resulted in increased type-2 inflammation and significantly increased fibrosis in vivo, phenocopying Map3k8\textsuperscript{−/−} mice. Collectively, our study identifies a novel and previously unappreciated role for TPL-2 as a molecular regulator of lipolysis in M2-MΦ’s, regulating type-2 inflammation, immunopathology and hepatic fibrosis.

Results

Map3k8-deficient mice develop increased T\textsubscript{H}2-mediated immunopathology and fibrosis following infection with Schistosoma mansoni

Following maturation and worm pairing, gravid worms release hundreds of eggs, many of which traverse the wall of the intestine and are released into the environment via the fecal route. However, many eggs do not successfully reach the intestinal lumen but instead become trapped in the intestinal wall or within vascularised organs, particularly the liver. An eosinophil and MΦ-rich fibrotic granuloma forms around trapped eggs causing significant tissue damage, orchestrated by CD4\textsuperscript{+} T\textsubscript{H}2 cells and a highly polarised type-2 immune response [5].

To test whether TPL-2 contributed to S. mansoni-associated intestinal and hepatic pathology and fibrosis, we infected Map3k8\textsuperscript{−/−} mice with 50 S. mansoni cercariae. Histological analysis indicated that S. mansoni-infected Map3k8\textsuperscript{−/−} mice had more fibrosis in the liver with larger hepatic granulomas (Fig 1A and 1B), despite a similar egg burden (S1 Fig) and serum LPS level as S. mansoni-infected WT mice (S1 Fig). Similarly, intestinal inflammation was also significantly increased in Map3k8\textsuperscript{−/−} mice (Fig 1C and 1D). Consistent with the increased collagen staining observed in Map3k8\textsuperscript{−/−} mice, collagen-synthesising genes, Col3 and Col6, were both significantly elevated in the liver and small intestine of Map3k8\textsuperscript{−/−} mice, compared to WT controls (Fig 1E), with significantly more hydroxyproline in the liver of Map3k8\textsuperscript{−/−} mice (Fig 1F).
Fig 1. Map3k8<sup>−/−</sup> mice develop increased hepatic and intestinal inflammation and fibrosis following S. mansoni infection. WT and Map3k8<sup>−/−</sup> mice were infected percutenously with 50 S. mansoni cercariae and analysed at 8 weeks post-infection. A & C) Perfused tissue was fixed and embedded in paraffin before sectioning and staining with Masson’s trichrome. B) Granuloma size was determined from 10–20 individual granulomas per sample measured using Image J. Scale bars are 1000μm (top), 200μm (middle) and 100μm (bottom). D) Intestinal pathology score, as described in
Map3k8\textsuperscript{-/-} mice had elevated expression of Il13 in the liver, but not Il1b, Tgfβ, Il17a, Ifng, Tnfa or Il6 (S1 Fig), suggesting that IL-13-driven fibrosis was exacerbated in Map3k8\textsuperscript{-/-} mice [33] rather the development of other inflammatory mechanisms of fibrosis [30].

CD4\textsuperscript{+} T\textsubscript{H}2 cell-derived IL-4 and IL-13 are essential for granuloma formation [6], mobilising and activating a suite of innate immune cells, including MΦ's and eosinophils, and promoting local collagen deposition. T\textsubscript{H}2 cell-mediated inflammatory responses are controlled by Foxp3\textsuperscript{+} regulatory T (T\textsubscript{REG}) cells [34], which restrain T\textsubscript{H}2 cell expansion. It was previously suggested that T cell intrinsic TPL-2 regulates T\textsubscript{H}2 [35] and Foxp3\textsuperscript{+} T\textsubscript{REG} cell differentiation [36]. However, these conclusions were based on in vitro experiments and were not tested in vivo. To determine whether Map3k8\textsuperscript{-/-} mice had dysregulated T\textsubscript{H}2 and Foxp3\textsuperscript{+} T\textsubscript{REG} responses following S. mansoni infection, we crossed Map3k8\textsuperscript{-/-} mice with Il4\textsuperscript{GFP} and Foxp3\textsuperscript{RFP} reporter mice, generating dual-reporter Map3k8\textsuperscript{-/-} mice (Map3k8\textsuperscript{-/-}Foxp3\textsuperscript{RFP}Il4\textsuperscript{GFP}). These reporter mice allowed us to accurately and simultaneously monitor T\textsubscript{H}2 (Il4\textsuperscript{GFP}+) and Foxp3\textsuperscript{+} T\textsubscript{REG} (Foxp3\textsuperscript{RFP}+) cells in Map3k8\textsuperscript{-/-} mice without the requirement for re-stimulation or intra-nuclear staining. Map3k8\textsuperscript{-/-} deficiency did not alter CD4\textsuperscript{+}CD25\textsuperscript{+}Foxp3\textsuperscript{+} T\textsubscript{REG} cell frequencies in the spleen, mesenteric lymph node (MLN) or in the local liver tissue, indicating that TPL-2 was not required for T\textsubscript{REG} cell development or recruitment following S. mansoni infection (Fig 1G, top row). However, CD4\textsuperscript{+}CD44\textsuperscript{+}T\textsubscript{H}2 cells in both lymphoid tissues and the liver were significantly increased in Map3k8\textsuperscript{-/-} mice compared to WT mice (Fig 1G, middle row). Map3k8\textsuperscript{-/-} deficiency also increased the frequency of Il4\textsuperscript{GFP}Foxp3\textsuperscript{RFP}+ cells in the MLN.

Pharmacological inhibition of MEK1/2, a downstream target of TPL-2, protected mice from bleomycin induced fibrosis [31]. We have previously reported that bleomycin-induced fibrosis is mediated by a pro-inflammatory type-1/type-17 and TGFβ driven response, distinct from type-2 mediated pulmonary fibrosis[30]. It therefore remained unclear whether TPL-2 contributed to type-2 driven pulmonary fibrosis. To test this we treated mice intravenously with S. mansoni eggs to invoke type-2 inflammation in the lungs leading to the development of pulmonary fibrosis, as previously described [30]. Similar to responses in the liver, Map3k8\textsuperscript{-/-} mice had increased collagen staining in the lung and increased hydroxyproline levels, compared to WT mice given S. mansoni eggs (S2 Fig). In the lung tissue and local draining thoracic lymph nodes (TLN), Map3k8\textsuperscript{-/-} mice had increased Th2 cell frequency (S2 Fig) promoting increased Il13, Col6 and Mmp12 expression in the lung (S2 Fig). Collectively, these data indicate that TPL-2 is an important negative regulator of type-2 inflammation, immunopathology and fibrosis following S. mansoni infection or S. mansoni egg induced pulmonary fibrosis in vivo.

T cell-intrinsic TPL-2 does not regulate T\textsubscript{H}2-mediated immunopathology following S. mansoni infection

It has previously been reported that T cell-intrinsic TPL-2 regulates T\textsubscript{H}2 cell differentiation in vitro and acute type-2 inflammation in the airways [35], however it has remained unclear whether T cell-intrinsic TPL-2 regulates T\textsubscript{H}2 cell differentiation and function in vivo. To formally test whether T cell-intrinsic TPL-2 contributed to the enhanced inflammation and fibrosis observed in Map3k8\textsuperscript{-/-} mice (Fig 1) we restricted Map3k8 deficiency to T cells using Cd4\textsuperscript{Cre}Map3k8\textsuperscript{GFP/GFP} mice. Deletion of Map3k8 in T cells (Cd4\textsuperscript{Cre}Map3k8\textsuperscript{GFP/GFP}) had no impact on
granuloma development in the liver (Fig 2A and 2B) or small intestine (Fig 2C) following S. mansoni infection. Similarly, fibrosis (Fig 2A and 2C) and expression of collagen synthesising genes, Col3 and Col6, were not affected following the deletion of Map3k8 in CD4+ cells (Fig 2D). IL-5 and IL-10 production was significantly increased in re-stimulated MLN cells from Map3k8−/− mice, compared to WT cells; however production of these cytokines was not affected when Map3k8 was deleted in T cells only (Fig 2E). IL-17 production was low and unchanged between all groups, however IFNγ secretion from lymph node cells was reduced in Map3k8−/− mice and Cd4CreMap3k8floflo mice, in line with a previous report [18].

To further test whether T cell intrinsic TPL-2 was required for Th2 cell differentiation, we isolated naïve T cells (TCRβ+ CD4+CD44−) from WT and Map3k8−/− mice and polarised them under Th1 or Th12 conditions in vitro. Similar frequencies of IFNγ+ or IL-4+ cells were observed between WT and Map3k8−/− T cells, respectively (Fig 2F), suggesting that T cell-intrinsic TPL-2 does not contribute to Th1 or Th12 differentiation in vitro. Taken together, these data indicate that T cell-intrinsic TPL-2 is required for optimal IFNγ secretion in vivo, but does not contribute to Th12 cell differentiation in vitro or in vivo, and that T cell-intrinsic TPL-2 does not contribute to Th12 cell-mediated immunopathology following S. mansoni infection.

**Myeloid cell-intrinsic Map3k8 critically regulates Th12-mediated immunopathology**

Alternatively activated macrophages (AA or M2-Mφ) contribute significantly to inflammation, immunopathology and fibrosis following S. mansoni infection [12]. TPL-2 has a well-defined role in classically activated Mφ’s (M1 or CA-Mφ) [17, 20–29], however it is unclear whether TPL-2 contributes to M2-Mφ following S. mansoni infection. Firstly, to test whether myeloid cell-intrinsic TPL-2 contributed to the exacerbated immunopathology observed in Map3k8−/− mice, we restricted Map3k8 deletion to Lysozyme M-expressing cells using LysMCreMap3k8floflo mice (S3 Fig). Mice with myeloid cell-specific deletion of Map3k8 had significantly more inflammation with larger hepatic (Fig 3A and 3C) and intestinal (Fig 3B) granulomas and more severe intestinal pathology (Fig 3D), without any appreciable change in serum LPS (S3 Fig). Of note, a distinct collagen-rich fibrotic ring surrounded hepatic granulomas in LysMCreMap3k8floflo mice, which was absent in mice with WT myeloid cells. Increased collagen staining in the liver was supported by increased expression of collagen-synthesising genes, Col3 and Col6 (Fig 3E) and increased hydroxyproline (Fig 3F). Similar to Map3k8−/− mice, mice with myeloid cell-specific deletion of Map3k8 had elevated type-2 cytokine secretions (IL-13, IL-5 and IL-10) following lymph node re-stimulation without any appreciable change in IFNγ or IL-17A secretion (Fig 3E). Similarly, elevated expression of Il13 but not Il1b, Tgfβ, Il17a, Ifng, Tnfα or Il6 was observed in LysMCreMap3k8floflo mice, compared to control mice (S3 Fig). These data clearly indicated that macrophage/myeloid cell intrinsic-TPL-2 contributed significantly to the regulation of Th12-mediated inflammation and fibrosis following S. mansoni infection.

**Map3k8 regulates M2 macrophage activation**

Th12-cell derived IL-4 and IL-13 [6] activates IL-4 receptor (IL-4R)-expressing Mφ’s [7, 8] to prevent lethal pathology following S. mansoni infection. To determine whether myeloid cell-intrinsic TPL2 contributed to M2-Mφ’s activation in vivo, we isolated Mφ’s ex vivo from infected mice. Chimeric mice were generated following the observation that Map3k8−/− mice and LysMCreMap3k8floflo mice had significantly elevated type-2 inflammation and fibrosis, compared to WT controls (Figs 1 and 3). Generating 50:50 chimeric mice by reconstituting lethally irradiated WT mice with 50% bone marrow from CD45.1+ WT mice and 50% bone marrow from CD45.2+ Map3k8−/− mice (Fig 4A), normalised and controlled for these environmental differences allowing
Fig 2. T cell-intrinsic \( \text{Map3k8} \) does not contribute to exacerbated inflammation and pathology following \( \text{S. mansoni} \) infection.

\( \text{Cd4}^{\text{Cre}} \text{Map3k8}^{+/+} \) and \( \text{Cd4}^{\text{Cre}} \text{Map3k8}^{fl/fl} \) mice were infected percutaneously with 50 \( \text{S. mansoni} \) cercariae and analysed at 8 weeks post-infection. A–C) Perfused tissue was fixed and embedded in paraffin before sectioning and staining with Masson’s trichrome. B) Granuloma size was determined from 10–20 individual granulomas per sample measured using Image J. D) Expression of \( \text{Col3} \) and \( \text{Col6} \) was determined from RNA extracted from liver or small intestinal tissue. Data is expressed relative to \( \text{HPRT} \) and shown as a fold-change relative to uninfected mice. E) Mesenteric lymph node cells were re-stimulated with anti-CD3 for 3 days. Cytokines were measured in supernatants, by ELISA. F) Naive T cells (CD4\(^{+}\)CD44\(^{-}\)CD25\(^{-}\)CD62L\(^{+}\)) were FACS purified from WT and \( \text{Map3k8}^{-/-} \) mice and cultured under TH1 and TH2 conditions. Frequencies of CD4\(^{+}\)IFN\(\gamma\) and CD4\(^{+}\)IL-4\(^{+}\) cells were determined by intracellular FACS analysis on day 7. All experiments are representative of 2–3 independent experiments with 5–10 mice/genotype. * \( p < 0.05 \) as assessed by two-tailed Mann-Whitney test.

doi:10.1371/journal.ppat.1005783.g002
Fig 3. Myeloid cell (LysM⁺) expression of Map3k8 regulates Th2-mediated immunopathology and fibrosis following *S. mansoni* infection. LysMCreMap3k8+/+ and LysMCreMap3k8fl/fl mice were infected percutenously with 50 *S. mansoni* cercariae and analysed at 8 weeks post-infection. A–B) Perfused tissue was fixed and embedded in paraffin before sectioning and staining with Masson’s trichrome. Scale bar is 200μm. C) Granuloma size was determined from 10–20 individual granulomas per sample measured using Image J. D) Intestinal pathology score, as described in methods. E) Expression of Col3 and Col6 was determined.
Hydroxyproline was quantified in liver tissue from naive and S. mansonii-infected WT mice. Data is expressed relative to HPRT and presented as a fold-change relative to HPRT levels of uninfected WT mice. F) Cytokines were measured in supernatants, by ELISA. All experiments are representative of 2–3 independent experiments with 5–10 mice/genotype. * p<0.05 as assessed by two-tailed Mann-Whitney test.

To determine how TPL-2 was regulating M2-MΦ activation we used the well-described in vitro macrophage activation assay, activating bone marrow-derived MΦs (BMDM) with IL-4 and IL-13. Following 6hrs of exposure to IL-4 and IL-13, Arg1, Retnla, Chi3l3 and Ear11 were all significantly reduced in Map3k8−/− M2-MΦs, compared to WT M2-MΦs (Fig 5A–5D), similar to that observed in ex vivo MΦs. At 24hrs, Retnla, Chi3l3 and Ear11 were still significantly reduced in Map3k8−/− M2-MΦs, demonstrating a non-redundant role for TPL-2 in M2-MΦ activation. The early reduction of Arg1 expression in Map3k8−/− M2-MΦs led to a reduction in arginase activity with reduced ornithine production, as determined by LCMS (S4 Fig).

Inhibition of the kinase activity of TPL-2, using the pharmacological inhibitor, C34 [37], phenocopied Map3k8−/− MΦs indicating that the kinase activity of TPL-2 was responsible for the reduced Retnla expression in M2-MΦ (S4 Fig). Phosphorylated (p)STAT6, pERK, pp38α and pJNK were similar in Map3k8−/− and WT MΦs (Fig 5E), suggesting that TPL-2 did not regulate responsiveness of MΦs to IL-4 and/or IL-13 and was not required for activation of these downstream transcription factors or kinases.

To determine whether TPL-2 regulated the expression of other genes, beyond the characteristic M2-MΦ-associated genes, we profiled the transcriptional landscape of WT and Map3k8−/− MΦs following 24hrs of IL-4 and IL-13 stimulation (Fig 6A). Pathway analysis identified increased inflammatory pathways in Map3k8−/− M2-MΦs, including proliferation, migration and fibrogenesis (Fig 6B). Of the significantly differentially regulated genes (P<0.05, > 2-fold, relative to un-stimulated) (Fig 6C), we identified 351 Map3k8-dependent genes (i.e. genes differentially regulated in WT only, Fig 6C and 6D) and 279 Map3k8-regulated genes (i.e. genes differentially regulated in Map3k8−/− MΦs only, Fig 6C and 6D). Of note, several of these elevated genes in Map3k8−/− M2-MΦs contribute to fibrogenesis, including Adam19 [38], Cxcr4 [39], Mmp13 [40], Cav1 [41], Itgav [42] and Vcam1 [43] (Fig 6F). Of the commonly regulated genes in both WT and Map3k8−/− M2-MΦs (Fig 6E and 6G) Map3k8−/− M2-MΦs had elevated expression of collagen-synthesising genes (Col1a1, Col3a1, Colla5a2) and the connective tissue growth factor, Ctgf (Fig 6H) compared to WT MΦs (Fig 6G). Concurrent with increased expression of pro-fibrotic genes, characteristic M2-MΦ genes (Retnla, Arg1, Ear11 (Rnas2) and Chi3l3) were reduced, compared to WT M2-MΦs (Fig 6H). Taken together, these in vitro gene expression data suggested that Map3k8−/− M2-MΦs had both elevated pro-fibrotic properties and reduced regulatory/inhibitory functions (Arg1 and Retnla (Figs 4 and 5)).

**Map3k8 regulates lipolysis for proficient alternative activation of Macrophages**

Oxidative lipid metabolism is a metabolic programme recently reported to be essential for M2-MΦ activation [44]. It also has previously been reported that TPL-2, MEK 1/2 and ERK 1/2...
Fig 4. TPL-2 is required for M2 activation of Macrophages, in vivo.

WT C57BL/6 mice were lethally irradiated (900rad) and reconstituted with 50% CD45.1+ WT bone marrow and 50% Map3k8−/− bone marrow and left for 6–8 weeks, prior to infection with 50 S. mansoni cercariae. A) After 8 weeks of infection, mice were sacrificed and CD3−CD19−CD11b+F4/80+ Macrophages were FACS-sorted. B-C) Expression of Arg1, Relma, Chi3l3, Col1a1, Col3a1 and Ctgf was determined from RNA extracted from purified macrophages. Data is expressed relative to HPRT and presented as a fold-change relative in genotype-controlled naïve bone marrow derived macrophages. Experiments are representative of 2 independent experiments with 5 mice/genotype. * p < 0.05 as assessed by two-tailed Mann-Whitney test.

doi:10.1371/journal.ppat.1005783.g004
**Fig 5. TPL-2 is required for M2 activation of Macrophages, in vitro.**

A-D) Bone marrow-derived macrophages (BMDM) were stimulated with IL-4 and IL-13 for 6 or 24 hours, as indicated. Cells were harvested, RNA extracted and gene expression was determined by qRT—PCR and expressed relative to un-stimulated genotype control cells. E) BMDM’s were stimulated for 1.5, 3 and 6 hours, as indicated. Total Protein was extracted with phosphorylated and total protein levels of STAT6, ERK, p38a, JNK and α-tubulin determined by western blot. All experiments are representative of 2–3 independent experiments with 3–5 biological replicates and 3 technical replicates in each experiment. *p < 0.05 as assessed by two-tailed Mann-Whitney test.

doi:10.1371/journal.ppat.1005783.g005
[45–48] can regulate lipid metabolism in a variety of different cells. We therefore hypothesised that the compromised M2-ΜΦ activation of Map3k8-deficient ΜΦ’s was due to reduced lipid metabolism. To investigate this possibility, we analysed the expression of 220 genes involved in lipid metabolism from the transcriptional data obtained from WT and Map3k8−/− M2-ΜΦ’s (S1 Table) and identified 16 TPL-2-dependent genes involved in lipid metabolism that were up regulated in WT M2-ΜΦ’s but not in Map3k8−/− M2-ΜΦ’s (Fig 7A). These genes included the LDL receptor, Olr1, which is required for lipid uptake [49] and Adipoq encoding adiponectin, which promotes lipid oxidation [50] and the alternative activation of human ΜΦ’s [51]. In addition, Aldh1a2 (also referred to as Raldh2) which catalyses the synthesis of the lipid metabolism-promoting metabolite, retinoic acid, from retinaldehyde [52] and the NFκB-regulated sialyltransferase, St8sia1 [53], which catalyzes the transfer of sialic acid from CMP-sialic acid to GM3 to produce gangliosides, were reduced in Map3k8−/− M2-ΜΦ’s, compared to WT M2-ΜΦ’s. Several of these genes are downstream of TPL-2 (S5 Fig), supporting our hypothesis that TPL-2 regulates lipolysis in M2-ΜΦ’s. Together these changes in gene expression in Map3k8-deficient M2-ΜΦ’s were consistent with TPL-2 signalling regulating lipolysis.

To formally test whether lipid metabolism was compromised in Map3k8−/− M2-ΜΦ’s, we used extracellular flux analysis and measured the oxygen consumption rate (OCR) and spare respiratory capacity (SRC, the quantitative difference between maximal uncontrolled OCR, and the initial basal OCR, indicative of commitment to oxidative phosphorylation) in un-stimulated and IL-4/IL-13 mediated M2-ΜΦ’s. At baseline, un-stimulated WT and Map3k8−/− ΜΦ’s had a similar OCR and SRC (S6 Fig). However, Map3k8−/− M2-ΜΦ’s had significantly reduced OCR and SRC (Fig 7B and 7C), indicating that TPL-2 is required for lipid metabolism in M2-ΜΦ’s, and providing a mechanistic explanation for reduced M2-ΜΦ’s in Map3k8−/− mice.

Map3k8−/− mice and LysMcCreMap3k8fl/fl mice, which had compromised M2-ΜΦ activation, had elevated Th2 cell responses. It has previously been reported that M2-ΜΦ’s can directly regulate T cell responses [13–15]. We therefore hypothesised that Map3k8−/− M2-ΜΦ’s would not regulate Th2 cell differentiation and proliferation as well as WT M2-ΜΦ’s. To test this hypothesis, we co-cultured WT or Map3k8−/− BMMΦ’s with naïve cell trace violet (CTV)-labelled CD4+CD44+ T cells in the presence of IL-4, IL-13 and OVA and determined the proliferation (CTV dilution) and differentiation (Il4gfp expression) of T cells. After 3 days, 24% of T cells had proliferated and differentiated when co-cultured with WT ΜΦ’s (Fig 7D, top row middle panel). However, co-culture of T cells with Map3k8−/− ΜΦ’s led to significantly more Th2 cell differentiation and proliferation (~38%, Fig 7D, bottom row middle panel). Finally, to determine whether lipid metabolism contributed to ΜΦ-mediated regulation of Th2 cell proliferation and differentiation we pre-treated ΜΦ’s for 6 hours with Orlistat, an irreversible lipase inhibitor, prior to co-culture with T cells. Orlistat treated WT ΜΦ’s led to more Th2 cell proliferation and differentiation (~32%, Fig 7D, top row right panel), phenocopying Map3k8−/− ΜΦ’s and indicating that lipid metabolism in IL-4/IL-13 activated ΜΦ’s was required for optimal ΜΦ-mediated control of Th2 cell proliferation, as previously reported [44]. Of note, Orlistat treated Map3k8−/− ΜΦ’s only led to a small increase in Th2 cell proliferation, suggesting that lipid metabolism was already at a minimum in Map3k8−/− ΜΦ’s.

Taken together this study has demonstrated that TPL-2 is a critical regulator of immune-mediated pathology and fibrosis following S. mansoni infection, functioning as an important metabolic regulator in M2-ΜΦ activation.

**Discussion**

Liver fibrosis and cirrhosis, which is responsible for over 1.5 million fatalities per year [54], can develop following a variety of infectious insults, including chronic infection [4]. Diseases
Fig 6. TPL-2 regulates pro-fibrotic and immuno-regulatory pathways in M2 macrophages. Bone marrow-derived macrophages (BMDM) were stimulated with IL-4 and IL-13 for 24 hours. Cells were harvested, RNA extracted and genome-wide transcriptional expression was determined by microarray analysis using 3 biological replicates. A) Heat map of differentially regulated genes in un-stimulated and IL-4+IL-13 stimulated cells. B) Ingenuity pathways analysis of transcriptional profiles of differentially regulated genes. C-F) Venn diagram and bar graphs of TPL-2 dependent (1), common (2) and TPL-2-regulated genes.
characterized by persistent T_{H2} cytokine responses, such as chronic helminth infections, are associated with the development of significant tissue pathology and fibrotic scarring. Although the molecular pathogenesis of fibrotic diseases are slowly emerging [4], there are very few novel therapeutic candidates progressing through clinical trials [55], highlighting a significant unmet medical need.

Two distinct inflammatory axes contribute to inflammation-driven fibrosis; type-1/ T_{H17} mediated inflammation [30] and type-2 driven fibrosis [56]. In this study we established that the Map3 kinase, TPL-2, is an important negative regulator of chronic type-2 inflammation-driven fibrosis following schistosome infection. These data are in contrast to a previous study testing the role of TPL-2 in three models of pro-inflammatory type-1/17-associated fibrosis (carbon tetrachloride-, methionine-choline-deficient diet- and bile duct ligation-induced fibrosis)[57, 58]. In two of these three models Map3k8^{−/−} mice had significantly reduced fibrosis [32]. These seemingly contrasting results most likely reflect the different inflammatory events contributing to the fibrogenic response. For example, it has been widely reported that TPL-2 is required for pro-inflammatory type-1/17-associated inflammation and immunity [17–29]. It therefore stands to reason that TPL-2 would be required for pro-inflammatory type-1/17-associated fibrosis, as reported by Perugorria and colleagues [32]. In contrast, TPL-2 appears to function as a negative regulator of type-2 inflammation in the lung and liver (Fig 1, S2 Fig), with increased acute [35] and chronic type-2 inflammation in Map3k8^{−/−} mice, as presented here. In this context, the exacerbated type-2 inflammatory response in Map3k8^{−/−} mice resulted in increased fibrosis. If these animal models reflect human disease, focused strategies targeting TPL-2 would benefit from identifying a prognostic biomarker and treating patients with type-1/17-associated fibrosis, rather than patients with type-2-associated fibrosis.

Inflammation-driven fibrosis involves a co-ordinated and often dysregulated wound healing response involving a variety of migratory leukocytes activating local stroma. TPL-2 is expressed in both leukocytes and local stroma and therefore identifying where TPL-2 was regulating the fibrogenic process was essential for us to identify how TPL-2 regulated fibrosis. It has been previously suggested that increased acute T_{H12} responses in Map3k8^{−/−} mice was due to a T cell-intrinsic role for TPL-2 [35], however this was not tested in vivo. Similarly, increased intestinal inflammation and tumorigenesis in Map3k8^{−/−} mice was attributed to a reduced frequency of Foxp3^{+} T_{REG} cells, [36], however again this was not specifically tested in vivo. Using Map3k8^{−/−} Il4^{−/−}Foxp3^{rfp} mice and re-stimulated local lymph nodes we identified that TPL-2 negatively regulated the differentiation, expansion and/or recruitment of TH2 cells, however this was not due to a T cell-intrinsic role for TPL-2, as mice with a T cell-intrinsic deletion of Map3k8 mounted similar T_{H12} responses as WT mice. Furthermore, exacerbated hepatic fibrosis observed in Map3k8^{−/−} mice was not observed in mice with a T cell-specific deletion of Map3k8, indicating that T cell-intrinsic TPL-2 had no impact on type-2-mediated inflammation or fibrosis in these systems.

TPL-2 has been extensively studied in TLR-mediated classical macrophage activation (CA, M1-MΦ) [20]. However, it was unclear whether TPL-2 contributes to M2-MΦ differentiation. M2-MΦ’s are central regulators of inflammation, wound-healing and fibrosis following schistosome infection [12]. Specifically, Arginase (Arg-1) in M2-MΦ’s catalyses the cleavage of arginine to ornithine and urea, depleting extracellular arginine and depriving local leukocytes of this essential amino acid. Consequently, M2-MΦ’s limit T cell proliferation, by starvation, in an Arg-1-dependent manner [14]. Similarly, production of Retnla by M2-MΦ’s can suppress T
Fig 7. TPL-2 regulates lipolysis in M2 macrophages and regulation of Tp2 cell differentiation and proliferation. Bone marrow-derived macrophages (BMDM) were stimulated with IL-4 and IL-13 for 24 hours. A) Analysis of genes involved in lipid metabolism was performed by Ingenuity pathways analysis (S1 Table) with Map3k8-dependent genes depicted in a heat map. B) After 24hrs of stimulation with IL-4 and IL-13 oxygen consumption rates (OCR) were determined in M2 macrophages using an XF-96 Extracellular Flux Analyzer (EFA) during sequential treatments with oligomycin, FCCP, and rotenone/antimycin. Spare
respiratory capacity (SRC), the quantitative difference between maximal uncontrolled OCR, and the initial basal OCR, is depicted in the plot. C) Basal oxygen consumption rates (OCR) and spare respiratory capacity (SRC) in WT and Map3k8−/− M2 macrophages. D) WT or Map3k8-deficient bone marrow-derived macrophages (BMDM) were generated from 3 individual mice and co-cultured with a pool of cell trace Violet (CTV)-labelled naïve OTII CD4+CD44hiCD44+ T cells and stimulated with IL-4 and IL-13 for 3 days. TH2 cell differentiation (i4ββP expression) and proliferation (CTV dilution) was determined by FACS after 3 days. In some wells BMDM were pre-treated with Orlistat for 6 hours and washed, prior to co-culture. Data is representative of 2–3 independent experiments with a minimum of 3 biological replicates per experiment. * p<0.05 as assessed by two-tailed Mann-Whitney test.

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microenvironment. In summary, this study has identified that TPL-2 is an important metabolic regulator in M2-MΦ's in vitro and that myeloid cell intrinsic TPL-2 critically controlled chronic type-2-mediated inflammation and fibrosis in vivo.

Materials and Methods

Animals

All mice were bred and maintained under specific pathogen-free conditions at The Francis Crick Institute. Strains used included: WT C57BL/6, Map3k8<sup>−/−</sup>[20], OTII [63], Il4<sup>grp</sup> [64], Foxp3<sup>rfp</sup>[65], Cd4<sup>Cre</sup>Map3k8<sup>flfl</sup>R26<sup>YFP</sup> (B6.Cd4<sup>Cre</sup>[66] crossed with Map3k8<sup>−/−</sup> and B6. R26<sup>YFP</sup>), LysM<sup>Cre</sup>Map3k8<sup>flfl</sup>R26<sup>YFP</sup> (B6.LysM<sup>Cre</sup>[67] crossed with Map3k8<sup>−/−</sup> and B6.R26<sup>YFP</sup>).

*S. mansoni* infection and egg-induced pulmonary inflammation

Mice were infected percutaneously via the tail with 50 cercariae of a Puerto Rican strain of *S. mansoni* (NMRI) obtained from *Biomphalaria glabrata* snails, kindly provided by Dr. Quentin Bickle, LSHTM. Infection intensity was determined following perfusion and granuloma size was determined from 10–20 individual granulomas per tissue sample, measured using Image J. Tissue pathology was analysed following Masson’s trichrome (Collagen, Blue; Nuclei, black/dark blue; Muscle, cytoplasm, Red) staining of 5μm sections from paraffin embedded samples. Intestinal pathology was determined using a comprehensive scoring system taking into account the level of infiltration, disruption and severity of the intestinal architecture [68]. Hydroxyproline content was quantified in liver tissue using a hydroxyproline assay kit according to the manufacturers recommendations (Cambridge Biosciences, UK). Tissue eggs were quantified by digesting a known weight of liver tissue with collagenase and liberase and isolating eggs on a discontinuous percoll gradient, as previously described [69]. For intravenous delivery of *S. mansoni* eggs, eggs were washed extensively in PBS, with 5000k delivered in 200μl of sterile PBS.

Bone marrow derived macrophage culture

Bone marrow cells were plated to a density of 5 x 10<sup>6</sup> cells per 90-mm bacterial Petri dish (Sterilin) in 10ml of DMEM/F-12 Dulbecco’s Modified Eagle Medium/Nutrient Mixture F-12 with GlutaMAX supplement (Gibco) supplemented with 10% FBS, antibiotics, 20% L-cell conditioned medium, L-Glutamine (1%), HEPES (1%), Sodium Pyruvate (1%) and β-mercaptoethanol (1%). After 4 days of culture, 10ml of additional medium was added and cells were cultured for a further 3 days. Non-adherent cells were washed away and the adherent cells were collected in 5ml PBS with 5% FBS and 2.5mM EDTA. For experiments the cells were re-plated in medium with 1% FBS without L-cell supplement and were incubated overnight before stimulation. Cells were stimulated with IL-4 and IL-13 (20ng/ml) (R&D systems) or LPS (100ng/ml) (Alexis Biochemicals). In some experiments, BMDM were generated from three individual mice and co-cultured with a pool of naïve OTII CD4<sup>+</sup>CD44<sup>−</sup>Il4<sup>grp</sup> T cells during stimulation with OVA peptide (323–339) (Invivogen). In some of these co-culture experiments, BMDM were pre-treated for 6 hours with Orlistat (100μM; Cayman), prior to co-culture.

Isolation of hepatic macrophages

Liver tissue was perfused and the organ was collected in gentleMACS columns (Miltenyi Biotech) in incomplete RPMI 1640 (Gibco). The tissues were dissociated in incomplete RPMI 1640 with Liberase TL (0.5mg/ml) (Roche), Collagenase (4μg/ml) (Roche) and DNase (7.5μg/ml) using the gentleMACS dissociator (Miltenyi Biotech). The partly digested tissue fragments were incubated at 37°C for 45 min, following which the tissues were completely dissociated. The
cellular fraction was run through a 100μm filter and the cells were centrifuged at 50g for 3 min, to pellet the non-parenchymal cells and the supernatant fraction was centrifuged at 320g for 5 min. The cellular fractions from both steps were pooled and collected in 1X HBSS Hanks balanced salt solution and mixed with OptiPrep (Sigma) to get a 17% w/v solution and overlayed with 1X GBSS Gey’s balanced salt solution. The samples were centrifuged at 400g for 15 min at room temperature with no brakes and the enriched layer of cells were collected from the interface of the GBSS and the 17% solution. The cells were stained and FACS sorted as Live/CD45+/CD3−/CD19−/NK1.1−/Ly6G−/SiglecF−/CD11b+/F4/80+.

Metabolism assays
Day 7 bone marrow-derived monocytes (BMDM) were cultured at a density of 5x10^5 cells in an XF24 plate (Seahorse Bioscience) over night. On day 8 cells were stimulated with 20ng/ml of recombinant IL-4 and IL-13 (R&D systems). On day 9 media was replaced with XF base medium (Seahorse Bioscience) supplemented with 100x Glutamax (Gibco), 100X Sodium Pyruvate (SIGMA) and 25mM glucose and the plate incubated for 10–30 minutes in a non-CO2 incubator at 37°C. For analysis of basal oxygen consumption rate (OCR) and extracellular acidification rate (ECAR), cells were analysed with XF-24 Extracellular Flux Analyzer (Seahorse Bioscience) under standard Seahorse running protocol. The Seahorse cartridge was loaded at a final concentration of 10μM Oligomycin (SIGMA O4876-5mg), 15μM FCCP (triflourocarbonylcyanide phenylhydrazone, SIGMA C2920-10mg), 1μM Rotenone (SIGMA R8875) plus 10μM Antimycin A (A8674-25mg) in ports A, B and C respectively. The bioenergetics profile consisted of basal OCR measurements in the absence of drugs and OCR/ECAR following the injection of drugs. All OCR/ECAR/SRC analyses were obtained from 5 replicates in 3 independent repeats.

LC-MS for arginase assay. At day 7, non-adherent cells were washed away and the adherent cells were collected in 5ml PBS with 5% FBS and 2.5mM EDTA. Cells were re-plated in medium with 1% FBS without L-cell supplement in a 6-well plate. Cells were kept for 12–18 hours, then stimulated with IL-4 and IL-13 (20ng/ml) (R&D systems) for 24hrs. Cells were washed twice with PBS, metabolites were extracted with ice cold mixture of acetonitrile/methanol/water (2:2:1, v/v/v). Plates were kept on dry ice for 5–10 min. Cells were collected, vortexed vigorously then spun for 20 min at 16,000g at 4°C. The supernatant was stored at -80°C until analysis. Residual protein content in the cell lysates was determined with Bicinchoninic acid method (BCA assay, Pierce). Prior to the LC-MS analysis, 100μL of the cell lysates were acidified with an equal volume of acetonitrile containing 0.2% acetic acid. The mixture was spun for 10 minutes at 16,000g, at 4°C. 2μL of the acidified mixture were used for the LC-MS analysis. Chromatography was performed on Agilent 1200 LC system, including a solvent degasser, binary pump and temperature-controlled auto-sampler. Samples were inject to a Cogent Diamond Hydride Type C silica column (150mm×2.1mm i.d, 4μm particle size and 100A pore size) and eluted with flow rate of 0.4 mL/min. The gradient employed is based on the number 3, according to [70]. Metabolites were detected using an Agilent Accurate Mass 6230 TOF apparatus, as previously described [71]. Detected m/z 175.11895 at 16.8 min and 133.0972 at 16.6 min were identified as arginine and ornithine, respectively, on the basis of unique accurate mass-retention time identifiers and spectral data. The reported abundances were normalized to the residual protein content in each corresponding sample.

T_{H2} polarization and re-stimulation of lymph node cells ex-vivo
Naive CD4+ T cells (CD4+TCRβ+CD44+CD25- P1 fueron FACS-purified from spleens of WT or Map3k8−/− mice. Naive T cells were cultured for 6 days in vitro with 10ng/mL IL-4 (R&D), 5ng/mL IL-2 (R&D), 10 μg/mL anti-IFNγ (XMG1.2, BioXcell), and CD3 (0.1–4.0μg) (145-
2C11, BioXcell) and CD28 (10μg/ml) (37.51, BioXcell) in complete IMDM (cIMDM, 10% fetal calf serum (FCS),100 U/mL Penicillin and 100 μg/mL Streptomycin (Gibco), 8mM L-glutamine (Gibco), and 0.05mM 2-mercaptoethanol (Gibco)). For re-stimulation of lymph node cells ex-vivo, lymph nodes were disrupted into a single cell suspension with 2x10^5 cells cultured in a 96-well, round bottom plate with 10μg/ml of anti-CD3 (145-2C11, BioXcell). Supernatants were harvested after 3 days for analysis by ELISA.

**qRT-PCR, ELISA, western blotting and serum LP**

**qRT-PCR.** Tissue samples were frozen in RNAlater (Sigma) and homogenized in QIAzol (Qiagen). Cell pellets were lysed in Buffer RLT (Qiagen). Total RNA was isolated as per manufacturer’s protocol. 100ng-1μg of total RNA was reverse transcribed using the QuantiTect Reverse Transcription Kit (Qiagen). cDNA produced was used for real time quantitative PCR with Power SyBrGreen (Applied Biosystems). The expression levels of different genes were normalised to hypoxanthine-guanine phosphoribosyl transferase (HPRT) expression and expressed as fold change relative to naïve or PBS-treated WT samples.

**ELISA.** IL-5, IL-13, IL-10, IFNγ and IL-17 were measured using DuoSet ELISA kits, according to the manufacturer’s instructions (R&D systems—Biotechne).

**Western blotting.** For immunoblotting, cell lysates were normalized to equal total protein content and resolved on 10% Criterion TGX Gels (Biorad). Separated proteins were transferred onto Trans-Blot Turbo PVDF transfer (Biorad) membranes (Immobilon). Specific bound antibodies were visualized by chemiluminescence (Merck Millipore). Endotoxin levels in serum samples were measured using an LAL chromogenic assay, according to manufacturer’s recommendations (Pierce, Thermo fisher).

**Cell sorting and flow cytometry**

Cell sorting was performed using a FACS Aria II (BD Biosciences) cell sorter. For sorting, cell suspensions were stained for 20 minutes with antibodies in PBS with 2% fetal calf serum (FCS) and then diluted in phenol-red free IMDM (Gibco) (with 1% FCS, 2mM EDTA (Invitrogen), 100U/mL Penicillin and 100μg/mL Streptomycin (Gibco), 8mM L-glutamine (Gibco), and 0.05mM 2-mercaptoethanol (Gibco)). Propidium iodide (PI) or LIVE/DEAD fixable blue dead cell stain (Life Technologies) was used to determine cell viability. Cells were stained for surface antigens by incubation with antibodies in PBS with 2% FCS (20 minutes at 4°C). Intracellular cytokine staining was performed following 6 hours re-stimulation with 50ng/mL phorbol 12-myristate 13-acetate (PMA, Promega) and 1μg/mL ionomycin (Sigma) and BD Golgi Stop and BD Golgi Plug (diluted 1:1000, BD Biosciences). After staining for surface antigens, cells were fixed and permeabilized (Fixation/Permeabilization diluent; eBioscience), prior to incubation with cytokine antibodies in Permeabilization buffer (eBioscience) for 20 min at 4°C. Cells were analyzed using a BD LSRII flow cytometer (BD Biosciences) and data processed using FlowJo software (Version X 10.0.7r2, Treestar Inc). Antibodies used were purchased with eBioscience, Biolegend or BD Pharmingen. They included: CD45 (30-F11), CD3 (17A2), CD4 (RM4-5, GK1.5), CD11b (M1/70), CD19 (6D5, eBio1D3), CD25 (PC61), CD44 (IM7), NK1.1 (PK136), Ly6G (1A8), SiglecF (E50-2440), and F4/80 (BM8), TCR β chain (H57-597). Staining was performed in the presence of FcR Blocking Reagent (Miltenyi Biotec).

**RNA, qRT-PCR and microarray**

RNA was isolated from tissues and cells using RNAeasy mini spin columns according to manufacturers’ instructions (Qiagen). cDNA was generated from 5ng of total RNA using WT-ovation Pico system (version 1) RNA Amplification System followed by double stranded cDNA
synthesis using WT-Ovation Exon Module. cDNA quality was determined using an Agilent BioAnalyzer and through hybridization performance on Affymetrix GeneChip mouse Genome 430A 2.0 microarray (Affymetrix) by the Systems Biology Unit at The Francis Crick Institute. Microarray data were quantile-normalized and analysed using GeneSpring software (Agilent). Differentially expressed genes were determined using ANOVA and t-tests. Genes with false discovery rate corrected p-values less than 0.1 and fold change values ≥1.5 were considered significant, and as indicated in figure legends. Three biological replicates of each subset were used. Pathways analysis was performed using Ingenuity Pathways Analysis (IPA, Ingenuity Systems, www.ingenuity.com).

Statistical analysis
Data sets were compared by Mann Whitney test using GraphPad Prism (V.5.0). Differences were considered significant at *p ≤ 0.05 using one or two-tailed tests.

Ethics statement
All animal experiments were carried out following UK Home Office regulations (project license 80/2506) and were approved by The Francis Crick Institute Ethical Review Panel.

Supporting Information
S1 Fig. TPL-2 regulated parasitology and pathology. WT and Map3k8−/− mice were infected percutenously with 50 S. mansoni cercariae and analysed at 8 weeks post-infection. A) S. mansoni eggs were quantified in the liver tissue as previously described [69]. B) Endotoxin levels (LPS) in serum was determined using an LAL assay kit at necropsy. C) Expression of Il13, Il1b, Tgfb, Il17a, Ifny, Tfna and Il6 was determined from RNA extracted from liver tissue. Data is expressed relative to HPRT and presented as a fold-change relative to genotype-controlled naïve mice. (TIFF)

S2 Fig. Map3k8−/− mice develop increased S. mansoni egg-induced pulmonary fibrosis. WT or Map3k8−/− mice were given 5000 S. mansoni eggs intravenously before necropsy at day 21. A) Lung tissue was fixed and embedded in paraffin before sectioning and staining with Mason’s trichrome. B) Hydroxyproline was quantified in liver tissue from naïve and S. mansoni egg treated mice. C) Frequency of TREG (CD4+ Foxp3RFP+) and TH2 (CD4+ Il4GFP+) cells in the thoracic lymph nodes (top row) and lung (bottom row) were determined by FACS on day 21. D) Expression of Il13, Col6 and Mmp12 was determined in RNA extracted from lung tissue. Data is expressed relative to HPRT and presented as a fold-change relative to genotype-controlled naïve mice. All experiments are representative of 2 independent experiments with 5 mice/genotype. * p < 0.05 as assessed by two-tailed Mann-Whitney test. (TIFF)

S3 Fig. Myeloid cell (LysM+) expression of Map3k8 regulates T12-mediated immunopathology and fibrosis following S. mansoni infection. LysMCreMap3k8+/+ and LysMCreMap3k8ββ/ββ mice were infected percutenously with 50 S. mansoni cercariae and analysed at 8 weeks post-infection. A) Detection of TPL-2 protein in macrophages (Live/Dead− CD45+F4/80+”LysMCreR26eYFP”) from LysMCreMap3k8+/+ and LysMCreMap3k8ββ/ββ mice. B) Endotoxin levels (LPS) in serum was determined using an LAL assay kit at necropsy. C) Expression of Il13, Il1b, Tgfb, Il17a, Ifny, Tfna and Il6 was determined from RNA extracted from liver tissue. Data is expressed relative to HPRT and presented as a fold-change relative to genotype-controlled naïve mice. (TIFF)
**S4 Fig. TPL-2 regulated macrophage activation.** A) WT and Map3k8−/− bone marrow-derived macrophages (BMDM) were stimulated with IL-4 and IL-13 for 24 hours with cell lysates used for arginine metabolism profiling using LC-MS. B) WT and Map3k8−/− bone marrow-derived macrophages (BMDM) were stimulated with IL-4 and IL-13 for 24 hours in the presence of a specific TPL-2 inhibitor, C34. Cells were harvested, RNA extracted and Retnla expression was determined by qRT-PCR and expressed relative to un-stimulated genotype control cells.

(TIFF)

**S5 Fig. TPL-2 regulated lipid metabolism pathways in M2 macrophages.** Ingenuity pathways analysis of lipid metabolism pathways (S1 Table) from bone marrow-derived macrophages (BMDM) stimulated with IL-4 and IL-13 for 24 hours, as in Figs 5 and 6. Elevated genes involved in lipid metabolism in WT, but not Map3k8−/− macrophages, are displayed, with their relationship with Map3k8 highlighted via intermediate genes.

(TIFF)

**S6 Fig. Lipolysis in un-stimulated WT and Map3k8−/− macrophages.** Bone marrow-derived macrophages (BMDM) were generated and left un-stimulated for 24 hours. A) Oxygen consumption rates (OCR) were determined using an XF-96 Extracellular Flux Analyzer (EFA) during sequential treatments with oligomycin, FCCP, and rotenone/antimycin. Spare respiratory capacity (SRC), the quantitative difference between maximal uncontrolled OCR, and the initial basal OCR, is depicted in the plot. B) Basal oxygen consumption rates (OCR) and (C) spare respiratory capacity (SRC) in WT and Map3k8−/− macrophages. Data is representative of 3 independent experiments.

(TIFF)

**S1 Table. TPL-2 regulates lipolysis in M2 macrophages.** Bone marrow-derived macrophages (BMDM) were stimulated with IL-4 and IL-13 for 24 hours, as in Fig 4. Analysis of genes involved in lipid metabolism was performed by Ingenuity pathways analysis. Gene expression is indicated, relative to un-stimulated macrophages. Highlighted genes are either absent in Map3k8−/− M2 macrophages (Map3k8-dependent) or absent in both WT and Map3k8−/− M2 macrophages.

(PDF)

**Acknowledgments**

We would like to thank Nicolaos Mathioudakis for his hard work and contribution to this study. We would also like to thank The Francis Crick Institute Flow Cytometry Facility, and in particular Bhavik Patel, Graham Preece, Wayne Turnbull, and Dr. Phil Hobson for the provision of cell sorting services in the production of this work. We are indebted to The Francis Crick Institute Procedural Service Section for production of GA lines and Biological Services, especially Anna Sullivan, Trisha Norton, Keith Williams and Adebambo Adekoya for animal husbandry and technical support; to Dr. Radma Mahmood and Dr. Radhika Anand for preparation of histology samples. We are extremely grateful to Dr. Laurent Dupays and Dr. Anotella Spinozza (The Francis Crick Institute) for help with the Seahorse assays, to Dr. Quentin Bickle (LSHTM) for providing S. mansoni cercariae. We also thank Dr. Steven Ley and members of the Ley laboratory who provided reagents, technical advice and discussion throughout these studies. The majority of this work was conducted at the Medical Research Council (MRC), National Institute for Medical Research (NIMR) at Mill Hill, which became The Francis Crick Institute in April 2015.
Author Contributions
Conceived and designed the experiments: YK JPL YL HK SP MSW. Performed the experiments: YK JPL YL HK SP LJE RM MSW. Analyzed the data: YK JPL YL HK SP MSW. Contributed reagents/materials/analysis tools: NRM SCCH EJP LPSdC SCL. Wrote the paper: YK MSW.

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