IL-17A both initiates, via IFNγ suppression, and limits the pulmonary type-2 immune response to nematode infection

Jesuthas Ajendra1,2, Alistair L. Chenery1,2, James E. Parkinson1,2, Brian H. K. Chan1,2, Stella Pearson1,2, Stefano A. P. Colombo1,2,3, Louis Boon4, Richard K. Grecnis1,2, Tara E. Sutherland1 and Judith E. Allen1,2

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INTRODUCTION

Innate and adaptive sources of interleukin-17A (IL-17A) are responsible for a range of neutrophil-associated inflammatory conditions as well as protection from many bacterial and fungal pathogens.1,2 In contrast, type-2 immunity is required for effective control of most helminth infections3,4 and is characterised by eosinophilic inflammation and the cytokines IL-4, IL-5 and IL-13. When both type-2 and IL-17 responses are present during helminth infection enhanced pathology is observed, as shown for human schistosomiasis5,6 and onchocerciasis.6 The detrimental relationship between IL-17A and type-2 immunity is illustrated by enhanced IFNγ production and that this suppression was essential for effective control of most helminth infections3 and is characterised by eosinophilic inflammation and the cytokines IL-4, IL-5 and IL-13. When both type-2 and IL-17 responses are present during helminth infection enhanced pathology is observed, as shown for human schistosomiasis5,6 and onchocerciasis.6 The detrimental relationship between IL-17A and type-2 associated inflammation and the cytokines IL-4, IL-5 and IL-13. When both type-2 and IL-17 responses are present during helminth infection enhanced pathology is observed, as shown for human schistosomiasis5,6 and onchocerciasis.6 The detrimental relationship between IL-17A and type-2 immunity is illustrated by enhanced IFNγ production and that this suppression was essential for effective control of most helminth infections3 and is characterised by eosinophilic inflammation and the cytokines IL-4, IL-5 and IL-13. When both type-2 and IL-17 responses are present during helminth infection enhanced pathology is observed, as shown for human schistosomiasis5,6 and onchocerciasis.6

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We and others have demonstrated a prominent role for IL-17A during infection with the lung-migrating nematode Nippostrongylus brasiliensis,11 a well-defined pulmonary model of type-2 immunity. After entering the host via the skin, N. brasiliensis larvae migrate through the lung, causing tissue damage and haemorrhage. IL-17A-dependent neutrophil recruitment is largely responsible for the lung damage in this model.11 We previously found that the chitinase-like protein Ym1 induces expansion of IL-17A-producing γδ T cells and Ym1 blockade or IL-17A-deficiency protects mice from peak lung damage.8 More surprising was our finding that Ym1 neutralisation or IL-17A-deficiency prevents the development of a full type-2 response during N. brasiliensis infection.8

The notion that IL-17A is required for development of a type-2 response appears counter to the evidence that type-2 cytokines suppress IL-17A production.9,10 However, previous studies using murine models of allergic inflammation also show impaired type-2 immunity in the face of IL-17A-deficiency12,13 or blockade.14 In an infection or injury context, the specific tissue as well as timing might all play decisive roles in whether IL-17A augments or suppresses type-2 responses. We therefore used N. brasiliensis infection to address the contribution of γδ T cell-derived IL-17A to the development of a subsequent type-2 immune response in the lung. We found that IL-17A suppressed early IFNγ production and that this suppression was essential for the optimal development of a type-2 response. Once the type-2 response was established, IL-17A acted as a negative regulator, revealing distinct roles during innate and adaptive stages of the response. Notably, Trichuris muris, a nematode restricted to the gastro-intestinal tract also induced a lung type-2 response that was IL-17A-dependent. However, we found no evidence that IL-17A regulated the intestinal type-2 response. Thus, IL-17A serves as a lung-specific regulator of the type-2 immune response.

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RESULTS

IL-17A-deficient mice mount a diminished type-2 response

In keeping with the known ability of *N. brasiliensis* to induce a strong pulmonary type-2 immune response on day 6 post infection (d6pi), we found the bronchoalveolar lavage (BAL) and lungs of C57BL/6 mice to be dominated by eosinophils (Supplementary Fig. 1a, b). This response was accompanied by elevated numbers of CD4^+^ T cells as well as induction of Group 2 Innate lymphoid cells (ILC2, Supplementary Fig. 1c, d). The establishment of a type-2 response was further confirmed by increased type-2 cytokine expression by CD4^+^ T cells and gene expression in whole lung (Supplementary Fig. 1e, f). As we and others previously reported,^8^ infected mice exhibited increased IL-17A production within the first 48 h post infection (Fig. 1a) and consistent with previous reports, the main source of IL-17A was γδ T cells. On d2pi the BAL consisted mainly of neutrophils (Supplementary Fig. 1a), which, together with *N. brasiliensis* larval migration, is known to cause acute lung injury.^11^

To investigate the role of IL-17A during the development of type-2 immune responses, we infected Il17a^−/−^KO mice and WT C57BL/6 controls with *N. brasiliensis* L3. Larvae leave the lung within 48 h and are expelled from the gut within 6–8 days. Consistent with our previous findings^8^ Il17a^−/−^KO mice were significantly more susceptible to infection exhibiting an intestinal worm burden almost two-fold higher than WT controls on d4pi (Fig. 1b). As expected, the early d2 neutrophilia in response to *N. brasiliensis* infection was muted in the Il17a^−/−^KO mice relative to the WT controls (Fig. 1c). Between d2pi and d6pi, there was a switch from neutrophilic to eosinophilic responses in the lungs (Supplementary Fig. 1a). Whilst increased eosinophil numbers were observed in all d6 infected mice relative to naïve animals, this increase was less evident for Il17a^−/−^KO mice (Supplementary Fig. 1g). ILC2s displayed a similar pattern, as cell numbers significantly increased with infection in WT but not in Il17a^−/−^KO mice (Fig. 1d). In addition, increased expression of the hallmark type-2 cytokine Il4 was observed at d6pi in WT mice, whilst Il17a^−/−^KO mice had significantly reduced Il4 expression compared with WT controls (Fig. 1e). Secretion of IL-13 protein levels at d6pi by total lung cells restimulated with α-CD3 were significantly higher in WT mice compared with naïve controls, whilst Il17a^−/−^KO mice failed to secrete detectable amounts of IL-13 (Fig. 1f). We also measured expression levels of the major mucins in the lung because host mucin production is another feature of protective type-2 responses,^15^ whilst *N. brasiliensis* infection drove an early increase in mucins Muc5ac and Muc5b expression in the lungs of WT mice at d2pi corresponding to a timepoint when the larvae are transitioning from the lungs. Expression of both Muc5ac and Muc5b were increased in WT mice compared with naïve controls, but reduced in Il17a^−/−^KO mice (Supplementary Fig. 1h).

While we observed an impairment in innate type-2 features, we next aimed to determine whether the impact of IL-17A-deficiency on type-2 immunity was due to changes in the adaptive immune response, mainly T cell activation or polarisation during *N. brasiliensis* infection. Using flow cytometry, we observed a failure to induce CD4^+^ T cells numbers on d7pi in the lungs in Il17a^−/−^KO mice compared with WT controls (Fig. 1g). In contrast, there were no significant differences in CD4^+^ T cell numbers in the lung-draining lymph nodes (Fig. 1g). We also examined expression of the Th2 transcription factor GATA3. While WT mice showed a significant increase in absolute numbers and frequency of GATA3^+^ CD4^+^ T cells upon infection, Il17a^−/−^KO mice had a significantly lower frequency of GATA3^+^ CD4^+^ T cells and failed to upregulate these cells on d7pi compared with WT controls (Fig. 1h). Not only were there fewer GATA3^+^ CD4^+^ T cells in the lungs of Il17a^−/−^KO mice, the CD4^+^ T cells in Il17a^−/−^KO mice produced significantly less IL-13 and IL-5 (Fig. 1i). Strikingly, by d7pi, IL-5^+^ and IL-13^+^ CD4^+^ T cell numbers did not increase in Il17a^−/−^KO mice in response to infection (Fig. 1j). At the same time point post-infection, we also found that expression of the activation marker CD69 was upregulated on CD4^+^ T cells in the lungs of WT but not Il17a^−/−^KO mice (Fig. 1k). However, CD69 did not differ between all tested groups in the lung-draining lymph nodes (Fig. 1l). Recently, Minutti et al. showed that epidermal growth factor receptor (EGFR) in complex with ST2 on T cells allows for IL-33-induced IL-13 production at the site of *N. brasiliensis* infection.^16^ We therefore analysed surface expression of EGFR and ST2 on lung and lung-draining lymph node T cells. *N. brasiliensis* infection increased the number of CD4^+^ T cells expressing these markers in the lung, but this increase was significantly reduced in Il17a^−/−^KO mice (Fig. 1k). No significant changes to ST2 and EGFR expression between WT and KO mice were observed in the lung-draining lymph nodes (Fig. 1l). We also measured PD-1 expression, an important regulator of T cell function during helminth infection.^17,18^ *N. brasiliensis* infection in WT mice led to increased numbers of CD4^+^ T cells expressing PD-1 in the lung and the lung draining lymph node of WT mice (Fig. 1k, l). However, there were significantly fewer PD-1^+^ CD4^+^ T cells in the lungs (Fig. 1k) of infected Il17a^−/−^KO mice but not the lymph node (Fig. 1l). Overall, these data demonstrated a significant impairment of the type-2 response in the absence of IL-17A during helminth infection, with lung CD4^+^ T cells failing to become fully activated and produce type-2 cytokines.

IL-17A leads to a downregulation of early IFNγ during *Nippostrongylus* infection

Rapid early IL-17A production is critical for protective immune responses in different settings of lung immunity. To better understand the early events unfolding in the lung during *N. brasiliensis* infection, we performed a Nanostring gene expression array using a myeloid immunity panel (700 genes). In whole lung, differentially expressed (DE) genes between naïve WT mice and infected WT and Il17a^−/−^KO mice at d2pi were assessed in total unamplified RNA (Fig. 2a). IL-17A deficiency led to a distinct gene expression profile compared with WT mice in response to *N. brasiliensis* infection. Notably, when analysing all DE genes (Fig. 2a) using the Ingenuity pathway analyser (Qiagen), Ifng was predicted as the most significantly increased upstream regulator in *N. brasiliensis* infected Il17a^−/−^KO compared with WT mice (Fig. 2b). This led us to hypothesize that IL-17A may suppress IFNγ, which would facilitate Th2 cell development and explain why mice deficient in IL-17A cannot induce a full type-2 immune response. This hypothesis was consistent with our unpublished and published findings that γδ T cells were the main producers of IFNγ during infection, supporting the hypothesis that IL-17A plays an important role in downregulating IFNγ at the site of infection during the lung migratory phase of *N. brasiliensis* infection.
IFNγ neutralization in Il17a-KO mice rescues the impaired type-2 immune response

We next asked whether global suppression of early IFNγ by IL-17A was required for the full development of type-2 immunity in the lung. IFNγ was neutralised at day −1 and 1 of infection in Il17a-KO and WT mice, and responses examined at d8pi, a time point when the type-2 response should be fully developed (Fig. 3a). The significant defect in eosinophilic responses in Il17a-KO mice compared with WT mice was still evident at d8pi. However, blocking IFNγ in Il17a-KO mice enhanced eosinophil numbers...
Fig. 1. Mice deficient in IL-17A mount a diminished type-2 response at the site of infection. C57BL/6 (WT) and il17a-KO mice were infected with 250 N. brasiliensis L3s and cell frequencies and cytokines were measured at different time points post infection compared with WT naive mice. Frequencies of IFN-γ producing γδ T cells on d1pi and d2pi and representative flow-plots at d1pi (a). Worm burden in small intestine assessed in WT and il17a-KO mice on days 2, 4, and 6 post N. brasiliensis infection (b). Absolute numbers of neutrophils (Ly6G+ CD11b+) (c) and lung ILC2s (Lineage− KLREP CD127− CD90.2+ ST2+ ) as measured via flow cytometry (d). Relative mRNA expression of cytokine Il4 in whole lung as quantified by qRT-PCR (e). To test this hypothesis we neutralized IL-17A in WT mice and assessed single-suspension lung cells (f). Absolute numbers of live CD4+ T cells in lung tissue and lung draining lymph nodes (LdLNs) (g). Frequency and absolute numbers of GATA3+ CD4+ T cells in the lung (h). Representative flow-plots showing the frequency of IL-5 and IL-13 production by CD4+ T cells in lung from WT and il17a-KO infected mice (i). Absolute numbers of IL-5− and IL-13+ CD4+ T cells in the lung (j). Expression of CD69 on CD4+ T cells and absolute numbers of EGFR CD4+ CD8+ T cells and PD-1+ CD4+ T cells in lung (k) and LdLNs (l). Data are representative (mean ± s.e.m.) of at least 3 individual experiments (a, c–l) or pooled data from three experiments (b). Data were tested for normality using Shapiro-Wilk test and analysed using one-way ANOVA followed by Sidak’s multiple comparisons test for selected groups. *P < 0.05, **P < 0.01, ***P < 0.001, ****P < 0.0001.

(Fig. 3b). The same pattern was observed for numbers of CD4+ T cells in the lungs (Fig. 3c). To determine whether IFNγ neutralisation altered the type-2 response, we assessed expression of the key type-2 cytokines Il4 and Il13 and the type-2 marker Chil3. As expected, based on our results thus far, expression of these genes in the lungs was significantly reduced in infected il17a-KO mice compared with WT mice (Fig. 3d, e). Notably, IFNγ depletion completely recovered expression of these cytokines in il17a-KO mice compared with isotype-treated animals, with Il4 expression surpassing the levels seen in WT infected mice (Fig. 3d). Similarly, analysis of numbers of IL-5 and IL-13 producing CD4+ T cells at d4pi showed complete restoration of the type-2 response in il17a-KO mice that received the neutralising IFNγ antibody (Fig. 3f). Consistent with the ability of IFNγ to regulate type-2 cytokines, IFNγ depletion also restored the activation status of CD4+ T cells in the il17a-KO mice as shown by CD69 expression (Fig. 3g) and increased the numbers of CD4+ T cells expressing type-2 markers EGFR, PD1 and ST2 (Fig. 3h). Again, no effect was observed in these parameters in IFNγ-depleted WT mice. Together these data demonstrate that an initial reduction in IFNγ levels during N. brasiliensis infection mediated by IL-17A, allows the subsequent development of type-2 immunity in the lung.

IL-17A suppresses an established type-2 response in the lung IFNγ depletion in il17a-KO mice not only restored the type-2 response, but in some cases exceeded WT levels. Therefore, we hypothesized that although innate IL-17A promotes the establishment of type-2 immunity in N. brasiliensis infection, once the adaptive response is in place, IL-17 can act to negatively regulate the type-2 pulmonary response. To test this hypothesis, we neutralized IL-17A at d4pi, d5pi and d6pi in WT mice and assessed immune responses at d7pi (Fig. 4a). Blocking of later stage IL-17A led to a significant increase in both ILC2 numbers and frequencies in the lung (Fig. 4b), as well as the numbers of ILC2s producing IL-5 and IL-13 (Fig. 4c). Although CD4+ T cell numbers in the lung were comparable between isotype-treated and anti-IL-17A-treated WT mice (Fig. 4d), the ability of CD4+ T cells to produce type-2 cytokines may partly rely on IL-17A, as mice administered anti-IL-17A showed a slight increase in numbers of CD4+ T cells producing IL-5 and IL-13 (Fig. 4e). This data demonstrated that IL-17A can have differential effects depending on the time and status of infection. While early IL-17A promotes the type-2 response, later in infection IL-17A acts to suppress and limit excessive type-2 immunity, particularly in ILC2s.

IL-17A does not regulate type-2 immune responses at the site of T. muris infection

Our data demonstrate an impairment of the type-2 immune response in the lung during infection of il17a-KO mice with the lung-migrating nematode N. brasiliensis. We wanted to investigate whether impairment of type-2 immunity by IL-17A was unique to the pulmonary environment. Initially we examined type-2 cytokine gene expression in the small intestine of N. brasiliensis infected mice at d7pi. However, we did not observe any significant changes in WT infected mice (data not shown). We therefore decided to use Trichuris muris, a nematode that establishes infection solely in the gastro-intestinal tract. Infection with T. muris begins with the ingestion of infective eggs that accumulate in the caecum. L1 larvae hatch and penetrate the caecum and proximal colon wall, undergoing moults to L2 (d9-11pi), L3 (d17pi), L4 (d22 pi) and adults (d29-32). High dose infection in C57BL/6 mice induces a strong type-2 response by d17pi, and subsequent clearance of the adult parasites.22,23 We infected WT and il17a-KO mice with a high dose of 200 T. muris eggs and found worm counts in the caecum were comparable between mouse strains at d19pi and d32pi (Fig. 5a), indicating IL-17A does not alter parasite expulsion rate. Cell numbers in the caecum were analysed and no differences in eosinophil and neutrophil frequency were observed between il17a-KO mice and WT controls on d19pi and d32pi (Supplementary Fig. 2a, b), suggesting an IL-17A-independent recruitment mechanism for both these cell types in the large intestine. CD4+ T cell numbers in the mLN were also comparable on d19pi and d32pi between il17a-KO mice and WT controls (Supplementary Fig. 2c). Although there was an induction of type-2 cytokines in infected mice as measured by intracellular cytokine staining, the numbers of IL-4, IL-5 or IL-13-producing CD4+ T cells in the mLN did not significantly differ between the groups (Supplementary Fig. 2d). Similarly, the relative expression of cytokines Il4, Il5 and Il13 did not differ between il17a-KO mice and WT controls within the caecum (Supplementary Fig. 2e). Secreted levels of IL-5, IL-9 and IL-13 in MLN cells were also not impaired in il17a-KO mice relative to WT controls (Supplementary Fig. 2f). Together, these data failed to provide any evidence that IL-17A is an important regulator of type-2 immune responses in the intestine or draining lymph nodes during T. muris infection.

Previous studies have shown that despite the restriction of the T. muris lifecycle to the gastro-intestinal tract of the mammalian host, evidence of a type-2 immune response can be observed at distant sites, such as the lung.24 Therefore, the immune response in the lung of T. muris infected WT vs il17a-KO mice at d19pi and d32pi was assessed. Neutrophils numbers were increased in infected WT animals at d19pi and d32pi, but this was significantly reduced in il17a-KO mice on d19pi (Fig. 5b). No significant changes were observed for eosinophils (Fig. 5c). Whilst lung CD4+ T cell numbers in infected animals did not change compared with naive controls (Fig. 5d), il17a-KO mice had significantly fewer IL-5+ CD4+ T cells at d19 and d32pi compared with WT controls (Fig. 5e). Although the effect on IL-13+ CD4+ T cells was less evident, infected il17a-KO mice failed to significantly increase numbers of IL-13+ CD4+ T cells compared with uninfected controls (Fig. 5e). Supporting the intracellular cytokine staining, qRT-PCR analysis in whole lung tissue showed an impairment of type-2 cytokines in the il17a-KO mice, with significantly decreased expression of Il4 (d32pi) and Il5 (d19pi) (Fig. 5f).

Similar to infection with N. brasiliensis, we also observed an upregulation of IFNγ in the lung during T. muris infection in il17a-KO mice. Both the number and the frequency of IFNγ‘CD4+ T cells
Fig. 2 Presence of IL-17A leads to a downregulation of early IFNγ during *N. brasiliensis* infection. Whole lung RNA from C57BL/6 (WT) and *Il17a*-KO mice on d2pi with *N. brasiliensis* compared with WT naïve mice, were analysed by Nanostring. Unsupervised, hierarchically clustered heat map showing significant differentially expressed genes between infected WT, *Il17a*-KO mice and uninfected (naïve) WT (a). Top differentially regulated genes from (a) between infected WT and *Il17a*-KO mice were run in Ingenuity pathway analyzer, with top predicted regulators shown in (b). Relative expression of *Ifng* in whole lung of naïve WT and d2 *N. brasiliensis* infected WT and *Il17a*-KO mice (log2 expression relative to *actb* (β-actin)) (c). Frequencies of IFNγ⁺ γδ T cells, CD4⁺ T cells, CD8⁺ T cells and NK cells in WT and *Il17a*-KO mice d2pi compared with WT naïve mice as assessed by flow cytometry (d). Frequency of IFNγ⁺ γδ T cells 16 h post *N. brasiliensis* infection in WT and *Il17a*-KO mice (e). Representative flow plots showing CD44 and CD27 γδ T cell subsets in naïve mice and mice 16 h post *N. brasiliensis* infection as well as frequency of IFNγ⁺ CD27⁺ γδ T cells 16 h post *N. brasiliensis* infection in WT and *Il17a*-KO mice (f). Data (c-f) are expressed as mean ± s.e.m. and are representative of at least 2 individual experiments with at least three mice per infected group. Data were tested for normality using Shapiro-Wilk test and analysed using one-way ANOVA followed by Sidak’s multiple comparisons test for selected groups or student’s t test. *P < 0.05, **P < 0.01.
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in the lung were significantly increased in Il17a-KO compared with WT infected mice on d19pi (Fig. 5g). This data utilising T. muris infection models suggests that IL-17A-dependent suppression of IFNγ allows promotion of the type-2 immune response specifically in the lungs but not the intestine and highlights means of communication between the intestine and the lung involving IL-17A, not previously described.

**DISCUSSION**

IL-17A, the key cytokine of the IL-17 family, is central to barrier immunity, combating fungal infections and inducing antimicrobial proteins as well as neutrophil activating and recruiting chemokines. However, in the context of type-2 immunity, a combination of type-2 cytokines and IL-17A is often a signature for severe disease pathology. For example, IL-17A contributes to asthma...
pathology by enhancing IL-13 activity and a dysregulated balance between IL-17A and type-2 responses exacerbates pathology during schistosomiasis and onchocerciasis. Understanding the relationship between IL-17A and type-2 immune responses is thus critical, and we and others have previously demonstrated that development of a full type-2 response can require IL-17A.

In our effort to understand how IL-17A might be required for full type-2 immunity, we have discovered that IL-17A suppresses early IFN-γ expression in the lung during helminth infection. Although several studies show links between IL-17A and IFN-γ, whether IFN-γ is up- or downregulated in response to IL-17A varies with setting, timing and location. For example, enhanced IFN-γ in Il17a-KO mice has been described in a viral infection, experimental visceral leishmaniasis, and Toxoplasma gondii infection. Evidence also exists in the context of helminth infection, where a lack of IL-17A drives elevated IFN-γ during infection with the filarial nematode Litomosoides sigmodontis or Schistosoma japonicum and Schistosoma mansoni. In contrast, IL-17A can promote IFN-γ production during kidney-ischaemic reperfusion injury or Francisella tularensis infection. Importantly, the consequence of IL-17A-IFN-γ cross-regulations in the context of type-2 inflammation has never been shown and here we reveal IFN-γ downregulation as a new mechanism through which IL-17A establishes a protective type-2 response in the lung.

Another key finding of our study was that the requirement for IL-17A to suppress IFN-γ appears restricted to the lung. The protective type-2 immune response in the gut of Il17a-KO mice was not impaired, and mice were still able to expel N. brasiliensis from the small intestine and T. muris. However, this finding was that even though T. muris does not have a lung stage, the concurrent type-2 response in the lung was impaired in Il17a-KO mice. CD4+ T cells in the lung produced less type-2 cytokines in the Il17a-KO mice and consistent with our findings in N. brasiliensis, CD4+ T cells in Il17a-KO mice produced significantly higher amounts of IFN-γ than in their WT counterparts. These findings raise major questions as to the nature of the insult that induces IL-17 in the lungs of T. muris infected mice. Although our data suggest that the impact of IL-17A on type-2 development might be lung restricted, there may still be a fundamental requirement for suppression of IFN-γ for type-2 immunity to progress. Artis et al. demonstrated that the type-2 immune response during T. muris requires TSLP, and in very similar experiments to those described here, demonstrated that TSLP functions to suppress IFN-γ. Thus, early suppression of IFN-γ may be a general pre-requisite for the development of a type-2 response during helminth infection.
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environment with a requirement for IL-17A in the lung and TSLP (or other factors) in the gut.

We have not yet addressed the full mechanism behind IL-17A-mediated suppression of IFNγ during *N. brasiliensis* infection but it is notable that IL-17A not only impairs type-2 cytokine production, but also alters the cellular activation status and expression of type-2 markers. Interestingly, in our model, we only observe impairment of type-2 immune responses in the lung itself and not in the Th2 cells from lung-draining lymph nodes. Expression of EGFR and ST2, two markers closely associated with type-2 settings, were reduced on the CD4+ T cells of Il17a-KO mice in the lung. EGFR expression on Th2 cells is critical for resistance during GI helminth infection and a signalling complex between EGFR and ST2 can activate Th2 cells to secrete IL-13 in an antigen-dependent manner upon IL-33 exposure. Our data would suggest that this “licensing” of Th2 cells does not occur in the Il17a-KO mice during *N. brasiliensis* infection, indicating that IL-17A is needed for a proper induction of the adaptive Th2 response in the lung.

Fig. 5 Lack of IL-17A impairs concurrent type-2 immune responses in the lung following infection with *Trichuris muris*. C57BL/6 WT and Il17a-KO mice were infected with a high dose of *T. muris* and immune parameters investigated at d19 and d32pi compared with uninfected (naive) C57BL/6 WT and Il17a-KO mice. Worms counts in the caecum (a). Absolute numbers of neutrophils (b), eosinophils (c) and CD4+ T cells (d) per gram of lung at d19pi and d32pi compared with naive mice. Absolute numbers of IL-5+ and IL-13+ CD4+ T cells per gram of lung tissue (e). Relative mRNA expression of cytokines *Il4* and *Il5* from whole lung (log2 expression relative to *actb* (β-actin)) of infected mice (f). Absolute numbers and frequency of IFNγ+ CD4+ T cells per gram of lung tissue (g). Data are expressed as mean ± s.e.m. and are representative of 3 individual experiments with at least four mice per infected group and one mouse per control group. Data were tested for normality using Shapiro-Wilk test and analysed with one-way ANOVA followed by Sidak’s multiple comparisons test for selected groups. Data in (f) were log2 transformed to achieve normal distribution and statistical tests were performed on transformed data *P < 0.05, **P < 0.01.
It is well documented that type-2 responses are essential to limit excessive IL-1735–37 but a novel finding from our study is that the reverse is also true. While the early γδ T cell-derived IL-17A supported the type-2 response, late IL-17A, derived from both Th17 cells and γδ T cells, negatively regulated type-2 cytokines. To our knowledge, IL-17A suppression of type-2 cytokines has not previously been described in vivo and illustrates a major cross-regulatory axis between type-2 cytokines and IL-17A, each required to contain the excessive production of the other. In the numerous situations in which a combination of type-2 cytokines and IL-17A results in severe disease pathology,4,6,25 it is apparent that this cytokine balance has failed. Together our data demonstrate that early events in the lung shape the protective type-2 immune response, with IL-17A as a critical regulator of type-2 immunity. IL-17A, as a driver of tissue damage,8 may itself be needed to establish a subsequent type-2 repair response. However, the ability of IL-17A to then suppress type-2 responses, reveal an important feedback loop that must go awry during severe asthma and other type-2 conditions in which IL-17A plays a damaging and pathogenic role. Finally, in combination with previous data,34 suppression of IFNγ at barrier sites may be a central paradigm for type-2 immunity.

MATERIALS AND METHODS

Mice and ethics statement

For experiments using only WT mice, C57BL/6 J mice were obtained from Charles River. C57BL/6 Il17a−/− mice were originally provided by Dr Brigitta Stockinger.38,39 For Il17a-KO experiments C57BL/6 WT mice and C57BL/6 Il17aCre/Rosa26EGFP homozygote mice were bred at the University of Manchester. Mice were age- and sex-matched and all mice were housed in individually ventilated cages. Both males and females were used. Mice were not randomized in cages, but each cage was randomly assigned to a treatment group. Mice were culled by asphyxiation in a rising concentration of CO2. Experiments were performed in accordance with the United Kingdom Animals (Scientific Procedures) Act of 1986.

N. brasiliensis infection

N. brasiliensis was maintained by serial passage through Sprague-Dawley rats, as described.40 Third-stage larvae (L3) were washed ten times with PBS (Dulbecco’s PBS, Sigma) before infection. On day 0, mice were infected subcutaneously with 250 or 500 larvae (L3). At various time points mice were euthanised, BAL was performed with PBS containing 1% BSA and lungs were taken for further analysis. For worm counts, the small intestines of infected mice were collected in PBS. Small intestines were then cut longitudinally along the entire length, placed in a 50 ml Falcon and incubated at 37 °C for 4 h. Settled worms were then counted with the aid of a dissecting microscope.

Flow cytometry

Single-cell suspensions of the lung were prepared by digesting minced lung lobes for 30 min at 37 °C with 0.2 U/ml Liberase TL (Roche) and 80 U/ml DNase (Life Tech) in Hank’s balanced-salt solution before forcing tissue suspensions through a cell strainer (70 μm, Greiner). Red blood cells were lysed using Red Blood Cell Lysing Buffer Hybri Max (Sigma) for 3 min at RT and reaction was halted by diluting samples in PBS. Total live cells were counted with AO/PI dye on an automated cell counter (Auto2000, Nexcelom). Cells were stained for live/dead (Life Technologies) with AO/PI dye on an automated cell counter (Auto2000, Nexcelom). Cells were stained for live/dead (Life Technologies) and incubated with Fc-block (1:500 CD16/CD32 and 1:50 mouse serum) and were then stained with fluorescence-conjugated antibodies. Cells were identified by expression of surface markers as follows: neutrophils Ly6G/CD11b+; eosinophils CD11b+CD11c−SigF−; CD4 T cells CD4+; TCRβ+CD11b−, γδ T cells TCRβ+, TCRγδ+, CD11b− and ILC2s Lineage− (CD11b, TCRβ, TCRγδ, CD90.2, CD44, CD49, CD127 A7R34 Invitrogen; EGFR EGFR1 Abcam). T. muris infection

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Ly6G, F4/80, CD11c, SigF, CD19) CD90.2+KLRG1+CD127+. Antibody clones used are listed in Table 1. For staining of intracellular cytokines, cells were stimulated for 4 h at 37 °C with cell stimulation cocktail containing protein transport inhibitor (eBioscience), then stained with live/dead. After surface antibody staining, cells were fixed for 10 min at 4 °C using IC fix (Biolegend) and cells were then incubated in for 20 min at RT in Permeabilization buffer (biolegend). Intracellular staining was performed for cytokines using antibodies for IL-5, IL-13, IL-17A and IFNγ as well as for Gata3 and Ym1. Samples were analysed by flow cytometry with LSR Fortessa or LSR II (Becton-Dickinson) and data analysed using FlowJo v10 software.

Quantification of cytokines

Single-cell suspensions of splenocytes, lung-draining lymph nodes or whole lung were stimulated ex vivo with N. brasiliensis excretory secretory product (E/S) antigen51 (1 µg/ml) or anti-CD3 (1 µg/ml). Cell supernatants were measured by cytokine bead array (CBA, BD Biosciences, UK) and cells were not randomized in cages, but each cage was randomly assigned to a treatment group. Mice were culled by asphyxiation in a rising concentration of CO2. Experiments were performed in accordance with the United Kingdom Animals (Scientific Procedures) Act of 1986.

Flow cytometry

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Table 1. List of flow cytometry antibodies used.

| Antigen | Clone | Manufacturer |
|---------|-------|--------------|
| CD11b  | M1/70 | BioLegend    |
| CD11c  | N41B  | BioLegend    |
| Ly6C   | HK1.4 | BioLegend    |
| CD4    | GK1.5 | BioLegend    |
| F4/80  | BM8   | eBioscience  |
| CD90.2 | 30-H12 | Biolegend |
| CD127  | A7R34 | Invitrogen   |
| KLRG1  | 2F1   | BioLegend    |
| TCRβ   | H57-597 | BioLegend |
| TCRγδ  | GL3   | BioLegend    |
| ST2    | DIH9  | BioLegend    |
| IL-5   | TRFK5 | BioLegend    |
| IL-17A | TC11-18H10.1 | BioLegend |
| Ly6G   | 1A8   | BD Biociences |
| Siglec-F | E50-2440 | BD Biociences |
| F4/80  | BM8   | ThermoFisher |
| IL-13  | eBio13A | ThermoFisher |
| Ym1    | Polyclonal | R&D Systems |
| RELM-α | Polyclonal | Peprotech    |
| IFNγ   | XMG1.2 | Biolegend |
| CD69   | H1.2F3 | Biolegend    |
| GATA3  | 16E10A23 | Biolegend    |
| EGFR   | EGFR1 | Abcam        |

Ly6G, F4/80, CD11c, SigF, CD19) CD90.2+KLRG1+CD127+. Antibody clones used are listed in Table 1. For staining of intracellular cytokines, cells were stained for 4 h at 37 °C with cell stimulation cocktail containing protein transport inhibitor (eBioscience), then stained with live/dead. After surface antibody staining, cells were fixed for 10 min at 4 °C using IC fix (Biolegend) and cells were then incubated in for 20 min at RT in Permeabilization buffer (biolegend). Intracellular staining was performed for cytokines using antibodies for IL-5, IL-13, IL-17A and IFNγ as well as for Gata3 and Ym1. Samples were analysed by flow cytometry with LSR Fortessa or LSR II (Becton-Dickinson) and data analysed using FlowJo v10 software.

Quantification of cytokines

Single-cell suspensions of splenocytes, lung-draining lymph nodes or whole lung were stimulated ex vivo with N. brasiliensis excretory secretory product (E/S) antigen51 (1 µg/ml) or anti-CD3 (1 µg/ml). Cell supernatants were harvested 72 h later and were stored at −20 °C until further analysis. Mouse IL-13 DuoSet ELISA kit (R&D Systems) was used for measurement of IL-13 levels. Mesenteric lymph node (MLN) cells from T. muris infected or uninfected mice were collected, cultured and restimulated ex vivo for 36 h with E/S as previously described.22 The concentrations of IL-5, IL-6, IL-9, IL-10, IL-13, IL-17A, TNFα and IFNγ in the MLN culture supernatant were measured by cytokine bead array (CBA, BD Biociences, UK) as per the manufacturer’s protocol.

Antibody depletion experiments

IFNγ was depleted using an anti-IFNγ monoclonal antibody (clone XM1G1.2) and injected intraperitoneally (500 µg/mouse/day) on days −1 and 1 of infection with N. brasiliensis. Control mice were injected with an equal amount of corresponding isotype control (GL113). IL-17A was depleted using an anti-IL-17A (17F3) or IgG1 isotype (both Invivo mAB) injected intraperitoneally (100 µg/mouse/day) on days 4, 5 and 6 post-infection with N. brasiliensis.
Extraction of RNA and quantitative real-time PCR
A fragment of the right lung lobe was stored in RNAlater (Ambion) before homogenization of tissue in Qiazol reagent with a TissueLyser (Qiagen). RNA was prepared according to manufacturer's instructions. RNA was quantified using a ND-1000 Spectrophotometer (NanoDrop Technologies). Reverse transcription of 1 µg of total RNA was performed using Tetro reverse transcriptase (Bioline). For reverse transcription, total RNA was treated with 50 U Tetro reverse transcriptase (Bioline), 40 mM dNTPs (Promega), 10 mM MgCl2 (Agilent) and specific primer pairs. PCR amplification was analysed by the second-derivative maximum algorithm (LightCycler 480 Sw 1.5; Roche), and expression of the gene of interest was normalized to that of the housekeeping gene Actb (beta-actin). A list of primer sequences used are shown in Table 2.

T. muris infection and E/S products
*T. muris* eggs were prepared from chronically infected stock mice as described previously. Mice were infected by oral gavage with adult excretory secretory product antigen *T. muris* eggs suspended in ddH2O. At day 19 post infection, *T. muris* burden was assessed by removing the caecum and proximal colon, opening them longitudinally and scraping the contents out with fine forceps. Individual worms were then counted by eye under a binocular dissecting microscope. *T. muris* adult excretory secretory product antigen (E/S) was prepared as described by. In brief, adult *T. muris* were cultured ex vivo at 37 °C, the culture supernatant was collected and centrifuged to remove eggs and worms. The resultant supernatant was then filter sterilised and stored at −20 °C until use for in vitro re-stimulation of MLN cells.

Nanostring RNA profiling
Extracted RNA was run on an Agilent 2200 Tape Station system to ensure high quality lung RNA; samples with a RIN value of <6.5 were excluded. Suitable RNA was then diluted to 20 ng/µL in RNase free H2O, measured using Qubit™ RNA HS Assay Kit (ThermoFisher) and run on a NanoString nCounter® FLEX system using the Myeloid Innate Immunity v2 panel (XT-CSO-MMII2-12) 220 as per manufacturer’s instructions. Raw data were loaded into nSolver version 4.0 using default settings. Non-normalised counts were then exported and subsequent analyses were performed in R (version 3.6) using RStudio Version 1.2.1335 Build 1379 – ©2009–2019 RStudio, Inc. Positive controls were analysed to ensure there was clear resolution at variable expression levels and negative controls were used to set a minimum detection threshold which was applied to all samples. Data were then normalised with EdgeR using the TMM method and differential expression between *N. brasiliensis*-infected WT and Il17a-KO mice was calculated via linear modelling with Empirical Bayes smoothing using the limma R package. Genes with an absolute fold change of greater than one and a significance value of under 0.05 after correction for multiple comparisons using the Benjamini-Yekuteili method were defined as “differentially expressed” and taken forward for further analysis. Heatmaps were then generated from normalized counts of DE genes using the ComplexHeatmaps R package. The networks and functional analyses of DE genes were generated with Ingenuity pathway analyser (QIAGEN Inc., https://www.qiagenbio-informatics.com/products/ingenuity-pathway-analysis). Data were then imported into R for visualisation.

**Table 2.** List of primer sequences used.

| Primer     | Sequence (5′-3′) |
|------------|------------------|
| Ccl8 forward | TTCTTGGCTGTCGTCATA |
| Ccl8 reverse | AGCAAGCTTGACCTAGTGA |
| Il5 forward | ACAAATTGCGCAAAAAAGAG |
| Il5 reverse | CACATTGAAGCAAGGCTAG |
| Chi3 forward | ACCTGCCCCGTTCAGTGCCAT |
| Chi3 reverse | CTTGGAAGTCTTCTTCCACAG |
| Il4 forward | CCTGCTCTTCTTCTCGAAAGT |
| Il4 reverse | CACATCTACTCTCGGCTAG |
| Retnla forward | TATGAAACAGTGGGCTCTCCT |
| Retnla reverse | GAGCAAGTTGCAAGATCTCTCCAC |
| Il13 forward | CTTGCACTAGGGGAGGCTT |
| Il13 reverse | CTTGACTCCTTAAGGAGCTTAT |
| Ifng forward | GAGAAGACTGCGAAGAGG |
| Ifng reverse | TTCAAGACTCTAAGGAGCTGAG |
| Actb forward | GCCGGACTCATCGTACTCC |
| Actb reverse | GTGACCCTGATCGTACTCC |
| Muc5ac forward | GCACTCAATAACAGCGAAT |
| Muc5ac reverse | CGAGTCACCCCCCTGAGTC |
| Muc5b forward | GAGTCACACTACCTCCTG |
| Muc5b reverse | TCTCATGTGTCGTTGTCAG |

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**AUTHOR CONTRIBUTIONS**
JA, ALC, J.P., and S.A.P.C. executed experiments, B.H.K.C. and S.P. provided experimental assistance. J.A., R.K.G., T.E.S. and J.E.A. designed experiments and analysed raw data. J.B. produced reagents. J.A., T.E.S. and J.E.A. wrote the original draft of the manuscript and all co-authors reviewed and edited the manuscript.

**ADDITIONAL INFORMATION**
The online version of this article (https://doi.org/10.1038/s41385-020-0318-2) contains supplementary material, which is available to authorized users.

**Competing interests:** The authors declare no competing interests.

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