IL-4 mediated resistance of BALB/c mice to visceral leishmaniasis is independent of IL-4R α signalling via T cells'.

SUPPLEMENTARY INFORMATION

Page

- 7 **Table S1:** *L. donovani* parasite burdens in the spleen, liver and bone marrow of CD4⁺ T cell-specific IL-4R α -deficient (Lck^{cre}IL-4R α ^{-/lox}) mice, wild-type littermate control (IL-4R α ^{-/lox}) and global IL-4R α ^{-/-} BALB/c mice at days 14 or 15 post-infection.
- 7 **Table S2:** Granuloma maturation in *L. donovani* infected CD4⁺T cell-specific IL-4R α -deficient (Lck^{cre}IL-4R α ^{-/lox}) mice, wild-type littermate control (IL-4R α ^{-/lox}) and global IL-4R α ^{-/-} BALB/c mice at day 14 and 15 post-infection.
- **Figure S1:** Representative photomicrographs of the hepatic granuloma response in an *L. donovani* infected wild-type (IL-4R $\alpha^{-/lox}$), CD4⁺ T cell-specific IL-4R α -deficient (Lck^{cre}IL-4R $\alpha^{-/lox}$), and global IL-4R $\alpha^{-/-}$ mice at days 15 post-infection.

Table S1: *L. donovani* parasite burdens in the spleen, liver and bone marrow of CD4⁺ T cell-specific IL-4R α -deficient (Lck^{cre}IL-4R α ^{-/lox}), wild-type littermate control (IL-4R α ^{-/lox}) and global IL-4R α ^{-/-} BALB/c mice at different times post-infection. Mice were infected with *L. donovani* on day 0 and parasite burdens were determined on day 14 or 15 in separate experiments.

Treatment	Mean parasite burden ± SE		
	Spleen	Liver	Bone marrow
Day 14			
WT (IL-4R $\alpha^{-/lox}$)	194 ± 16	4144 ± 256	1038 ± 31
CD4 ⁺ T cell-specific			
IL-4Rα-deficient			
$(Lck^{cre}IL-4R\alpha^{-/lox})$	223 ± 45	3597 ± 560	947 ± 108
Global IL-4Rα ^{-/-}	214 ± 31	4223 ± 337	788 ± 181
Day 15			
WT (IL-4R $\alpha^{-/lox}$)	91 ± 16	2349 ± 192	387 ± 106
CD4 ⁺ T cell-specific			
IL-4Rα-deficient			
$(Lck^{cre}IL-4R\alpha^{-/lox})$	143 ± 19	2312 ± 192	528 ± 99

Table S2: Granuloma maturation in *L. donovani* infected CD4⁺ T cell-specific IL-4R α -deficient (Lck^{cre}IL-4R α ^{-/lox}), wild-type littermate control (IL-4R α ^{-/lox}) and global IL-4R α ^{-/-} BALB/c mice at day 14 and 15 post-infection in separate experiments. ND – not determined.

Granuloma stage	Strain		
	WT control	CD4 ⁺ T cell-specific	Global IL-4Rα ^{-/-}
	$(\mathbf{IL}-4\mathbf{R}\alpha^{-/\mathbf{lox}})$	IL-4Ra-deficient	
		(Lck ^{cre} IL-4Rα ^{-/lox})	
Day 14			
Sterile	1.5 ± 1.1	0.4 ± 0.7	0.6 ± 1.1
Mature	6.8 ± 1.1	3.1 ± 1.7	2.7 ± 2.7
Immature	29.6 ± 7.1	22.7 ± 6.4	17.4 ± 5.6
Kupffer cells	61.9 ± 6.7	73.3 ± 7.9	67.2 ± 8.3
Day 15			
Sterile	2.00 ± 0.8	1.03 ± 0.9	ND
Mature	8 ± 2.5	5.83 ± 0.9	ND
Immature	28.18 ± 3.3	30.30 ± 3.6	ND
Kupffer cells	59.32 ± 9.0	62.5 ± 3.3	ND

Fig. S1: Representative photomicrographs of the hepatic granuloma response in *L. donovani* infected wild-type (IL-4R $\alpha^{-/lox}$), CD4⁺T cell-specific IL-4R α -deficient (Lck^{cre}IL-4R $\alpha^{-/lox}$) and global IL-4R $\alpha^{-/-}$ mice at day 15 post-infection. Immature granulomas and amastigotes within the Kupffer cells at Day 15 are denoted by (arrows).



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