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respective murine PAS positive cells are shown (G). ALT-803-induced IL-10 in BALF of Aspergillus challenged mice (H). Data is expressed as mean \pm SD, n=8 mice/group, *p<0.001.

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869 Legends for supplementary figures:

870 Supplementary figure 1. Relative mRNA expression of Th2 cytokines, quantification of eosinophils and goblet cells in rlL-15 pretreated mice exposed to 871 saline or Aspergillus challenge. A diagrammatic representation of rIL-15 treatment 872 and induction in the employed experimental asthma protocol is shown (A). A 873 representative photomicrograph of eosinophil accumulation in lung tissue of rIL-15 874 875 treated saline- or Aspergillus-challenge mice is shown (B-E, original magnification x200). Real time PCR for mRNA of IL-4 (F), IL-5 (G), IL-13 (H) and IL-10 (I) are shown in rIL-876 15 treated saline- or Aspergillus-challenge mice. A representative photomicrograph of 877 goblet cell hyperplasia in the lungs of rIL-15 treated saline or Aspergillus challenged 878 mice is shown (J-M). Data are expressed as mean ± SD, n=12 mice/group. *p<0.05 879 **p<0.001, *** p<0.0001. 880

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Supplementary figure 2. Analysis airway obstruction in rIL-15 treated dust mite and cockroach allergen challenged mice. A diagrammatic representation of rIL-15 treatment and allergen challenged protocol is shown (A). Airway resistance (RI) and compliance (Cydn) in response to various concentrations of methacholine were measured in saline and dust mite (B, C) and saline and cockroach allergen challenged Upparahalli et al

rlL-15 treated and not treated mice are shown (D, E). Data are expressed as mean \pm SD, n = 3 mice/group. *p<0.001.

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Supplementary figure 3. Relative mRNA expression for Th2 and eosinophil
cytokines and chemokines in Aspergillus-challenged DOX and non-DOX diet CC10 IL-15 transgenic mice. Real time PCR analysis for relative mRNA of IL-4, IL-5, IL13, eotaxin-1 and eotaxin-2 in saline- or Aspergillus challenged DOX and non-DOX diet
CC-10-IL-15 bitransgenic mice are shown (A-E). Data are expressed as mean ± SD,
n=12 mice/group. **p<0.001, *** p<0.0001, NS, not significant.

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Supplementary figure 4. Analysis of IL-10 producing regulatory T cells in 897 Aspergillus challenged non DOX and DOX exposed IL-15 bitransgenic mice. Flow 898 cytometry analysis showed that IL-10 producing regulatory T cells are increased in the 899 900 spleen and mediastinal lymph nodes in DOX exposed Aspergillus challenged mice compared to non DOX exposed Aspergillus challenged mice. The absolute number of 901 regulatory T cells and IL-10 producing regulatory T cells are shown in non DOX and 902 903 DOX exposed IL-15 bitransgenic mice (B-E). Data is the representative of 4 mice/group analyzed, *p<0.05. 904

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Supplementary figure 5. Analysis airway resistance and compliance in rIL-10
treated Aspergillus challenged mice. Airway resistance (RI) and compliance (Cydn)
in response to various concentrations of methacholine were measured in rIL-10 treated,
saline and Aspergillus challenged mice. Resistance (A) and compliance (B) is shown for

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Aspergillus challenged mice with and without rIL-10 treatment. . Data are expressed as mean \pm SD, n = 6 mice/group. *p<0.001.

Supplementary figure 6. Analysis of regulatory T cells in rIL-15 treated WT and IL-15^{-/-} mice and anti-IL-15 treated mouse model of experimental asthma. Flow cytometer analysis was performed using anti-CD4-PE, anti-CD25-PECy7 and anti-Foxp3-APC antibodies. The Foxp3 anti-IgG isotype match is used to identify Foxp3⁺ regulatory T cells (A). To establish that IL-15 association to regulatory T cells, we examined Splenocytes of saline and Aspergillus challenged WT and $IL-15^{-/2}$ mice. The decrease levels of regulatory T cells is observed at baseline and following allergen challenge in the IL-15^{-/-} mice compare to WT mice (A-E). Further, the flow cytometer analysis indicated that rIL-15 given WT and $IL-15^{-/2}$ mice show increase levels of regulatory T cells in the spleen compare non treated WT and $IL-15^{-1}$ mice (F-J). Data is the representative of 4 mice/group analysed.