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Insulin-like Growth Factor 1 Supports a Pulmonary Niche that Promotes Type 3 Innate Lymphoid Cell Development in Newborn Lungs

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In the initially published Figure 3, the numbers of ZBTB16^{iDTR} mice (n = 9) or Rosa26^{iDTR} treated with DT (n = 8) in Figure 3I was incorrect. The error occurred because the text in Figure 1I had accidentally been copied to Figure 3I. The revised Figure 3I now appears online and in print here with the corrected number of ZBTB16^{iDTR} mice (n = 7) or Rosa26^{iDTR} treated with DT (n = 7). The statistical analysis in the published manuscript was done using n = 7 for each group. Therefore, this involves no change in the conclusion or statistical relevance. Furthermore, in the originally published Figure S3B, an incorrect panel was copied. This error was made during the preparation of final figures. The corrected figure now appears online and in print here. The authors humbly accept these mistakes and sincerely apologize for any confusion that these errors might have made for the readers.

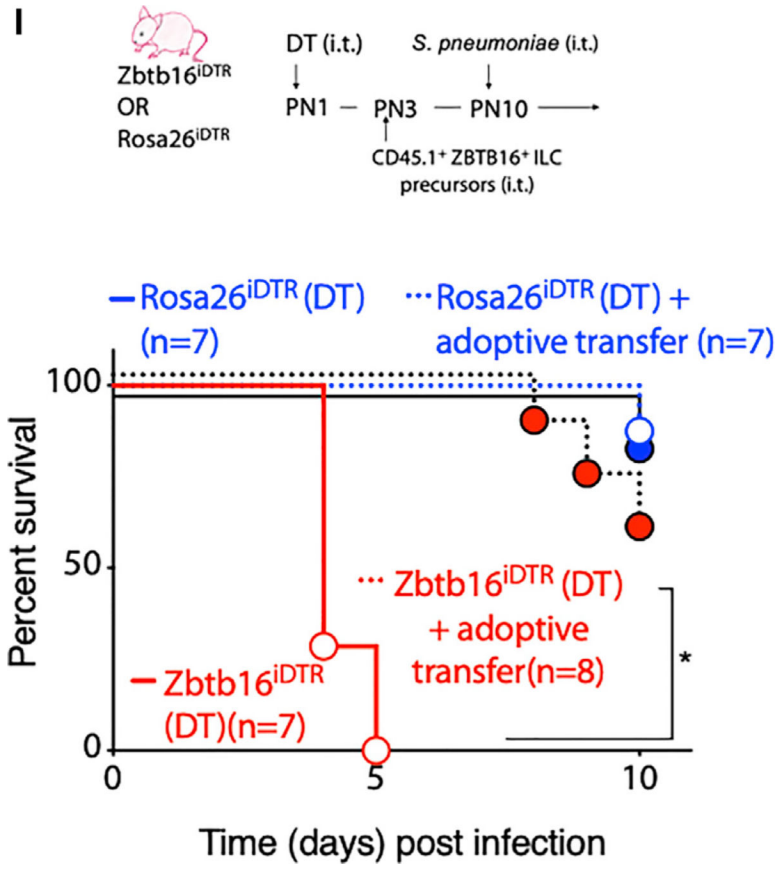

Figure 3I.

Pulmonary ZBTB16^{+} ILC Precursors Contributed to the Homeostatic Pool of Mature ILC3 in the Newborn Lung (corrected)

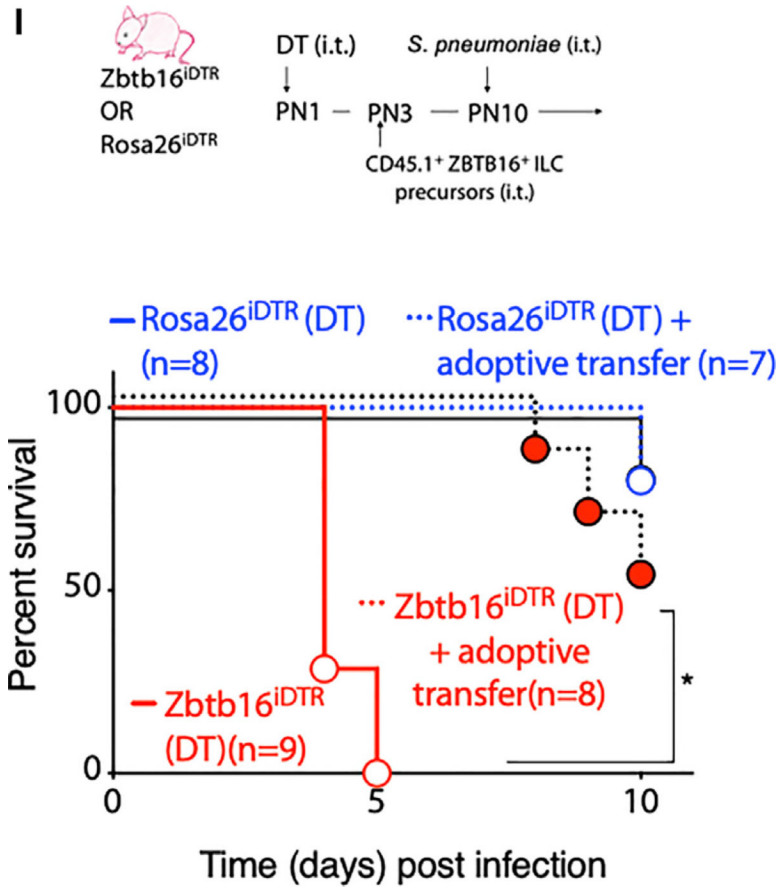

Figure 3I.

Pulmonary ZBTB16^{+} ILC Precursors Contributed to the Homeostatic Pool of Mature ILC3 in the Newborn Lung (original)

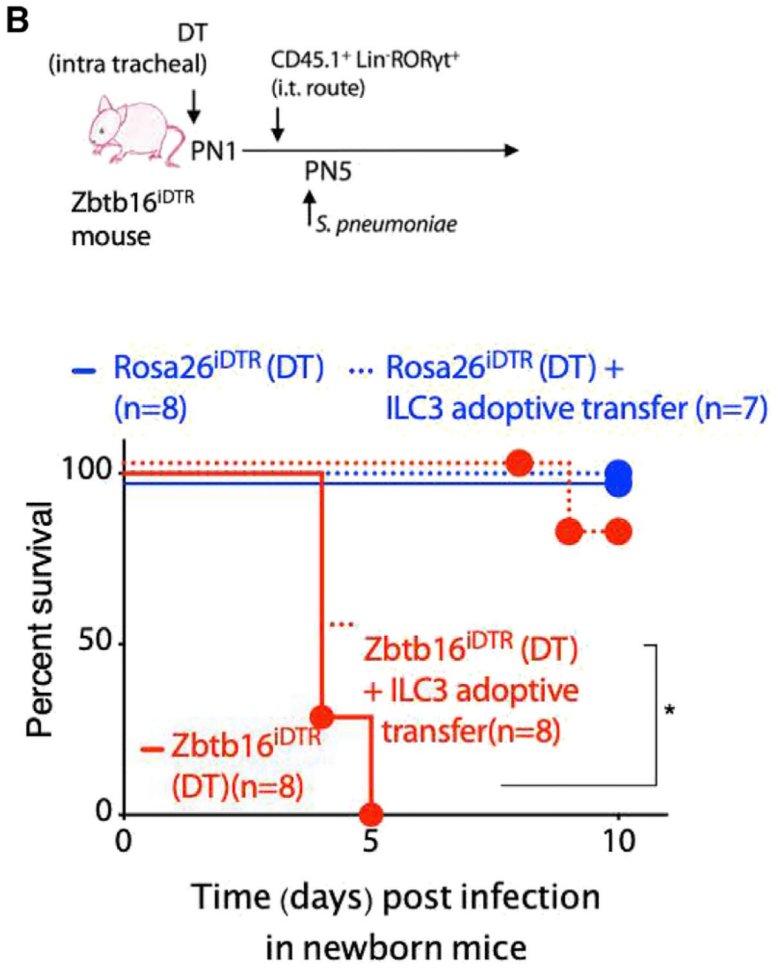

Figure S3B.

Pulmonary ZBTB16^{+} ILC precursors contribute to the homeostatic pool of mature ILC3 in the newborn lung (corrected)

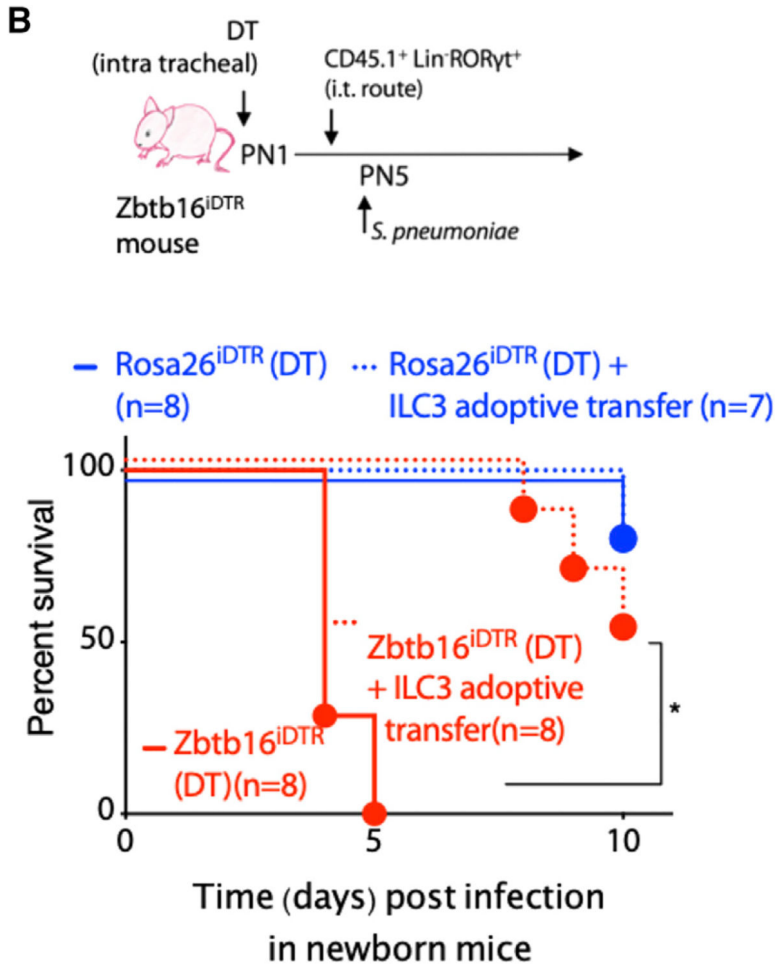

Figure S3B.
Pulmonary ZBTB16^{+} ILC precursors contribute to the homeostatic pool of mature ILC3 in the newborn lung (original)