

Lung macrophages: old hands required rather than new blood?

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Macrophages are key sentinel cells of the lung, clearing inhaled particulates and micro-organisms from the airway. These cells perform this function without provoking inflammatory responses that could lead to pneumonias. Previous work suggested that the lung macrophage population was maintained by the recruitment of blood monocytes.¹ However, this observation was made in transplanted lungs where macrophages had been depleted, either by hypoxia or chemotherapy. More recent work in murine models using cell fate mapping demonstrated that lung macrophages are generated early in embryonic development initially in the foetal yolk sac and slightly later in the foetal liver.² Macrophages that arise from these sources then seed the lung, where they reside as a self-renewing population with little further recruitment from blood monocytes.

The relevance and translation of these murine findings to the ontogeny of human lung macrophages has remained unclear. The study by Eguíluz-Gracia *et al*³ has provided some intriguing evidence regarding the stability and longevity of the lung macrophage population. This study prospectively followed 10 gender mismatched lung transplant patients for 2 years. Transbronchial biopsies were collected at regular intervals throughout the follow-up period. The macrophage populations in these biopsies were analysed for known macrophage markers, indicators of proliferation and, importantly, in situ hybridisation for the X and Y chromosomes to differentiate between recipient and donor cells. Comparisons to normal lung tissue obtained from cancer resections, suggest that there is no significant difference in the total number of macrophages in

transplanted lung tissue across the 2 years. Moreover, while there was a large amount of inter-individual variation the abundance of donor-derived macrophages was largely unchanged over the follow-up period. At the same time, there was a rapid infiltration of recipient-derived macrophages that, once established, also remained stable.

To investigate whether these infiltrating cells were recruited blood monocytes, Eguíluz-Gracia *et al*³ analysed the co-expression of the macrophage markers CD68 and DC-SIGN (CD209) and the monocyte marker calprotectin. Calprotectin was only detected on CD68+DC-SIGN+ macrophages at 2 weeks post-transplantation but not at 2 years post-transplantation. Taken together these results provided circumstantial evidence that blood monocytes were being recruited initially to the transplanted lung before maturing into macrophages as they became resident in the tissue. The investigators then went a step further to prove that recruited monocytes did indeed differentiate into lung macrophages by injecting human CD34+ cord blood cells into humanised mice that had been sublethally irradiated. The lungs of these animals were then harvested after 10 weeks and macrophages analysed for the presence of human antigens. In line with the human findings, the macrophages in these mouse lungs expressed the human CD45, CD68 and DC-SIGN antigens.

While these experiments demonstrated that monocytes were recruited to the murine lung, the loss of calprotectin over the 2 years suggested that this recruitment was not a common occurrence in the human lung. The authors therefore questioned whether recruited monocyte-derived lung macrophages could go on to proliferate to maintain this population. Eguíluz-Gracia and colleagues observed expression of the Ki67 protein marker of cellular proliferation in a median of 2% of the CD68+ macrophage population and that this proportion of Ki67 expressing cells remained stable over the 2 years. Interestingly, both donor and recipient macrophages were observed to express Ki67, although accurate quantitation of the number of proliferating donor cells was precluded due to low numbers of donor-

derived Ki67+ macrophages. These results suggest that donor cells can persist in the lung for at least 2 years and some of this longevity may be due to proliferation of these macrophages. Furthermore, blood monocytes can replenish the lung if alveolar macrophages are depleted in some way, but once the lung is refilled macrophage numbers may once again be maintained by local proliferation rather than further recruitment from the monocyte pool.

The biggest limitation of this study is that it is impossible to definitively identify whether the persistence of the macrophage population is due to cellular longevity or self-renewal, that is, the 2% of cells identified as proliferating give rise to the other 98% of macrophages. As with any study using human patients rather than murine models, there is substantial inter-individual variation in the sex, underlying disease and chemotherapy of the patients. However, none of these factors appeared to be related to the variation in cell densities. A further possibility is that the lack of significant changes in donor macrophage numbers could be confounded by the small numbers of subjects included. However, given the logistical issues in running such a study, not least persuading patients to return to hospital for transbronchial biopsies five times post-transplant, the authors should be commended on the quality of the data obtained. One further possible contribution to the inter-individual variation, also acknowledged by the authors, is that during counting only 50% of the cells could be called as having two sex chromosomes and thus being identified as a donor or recipient macrophage. However, the authors present data from biopsies sampled at the early time points demonstrating that the proportions of chromosome determined and undetermined cells were close to 50% in all samples examined, so this is unlikely to have skewed the cell counts.

The authors discuss the implications of this work in terms of its applicability for the use of donor monocyte-derived macrophages to replace defective macrophages in patients with alveolar proteinosis.⁴ However, this work also has implications for lung macrophage biology outside of the transplantation field, and COPD in particular. Macrophage numbers are increased in lung tissue taken from patients with COPD and the number of airways containing macrophages increases in line with disease severity.⁵ While numbers are increased, there are reports of macrophages being less mature in COPD in terms of both size⁶ and

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surface marker expression.⁷ Taken together with the increased expression of the chemokine monocyte chemoattractant protein-1 (CCL2) described in the COPD airway,⁸ these observations suggest that monocytes may be recruited to the COPD airway. The question remains what do these newly recruited cells do in the COPD lung and what is recruiting them?

The development of mature lung macrophages is controlled by a number of factors including granulocyte/macrophage colony-stimulating factor as well as interactions with the lung microbiome acquired early in life. The microbiome has been shown to play a role in susceptibility to inflammatory diseases of the gut and recent work implicates dysbiosis of the lung microbiome in a number of conditions including COPD (reviewed in ref. 9). Resident macrophages may be largely tolerant of the normal lung microbiota through recognition of bacterial-derived short chain fatty acids as in the gut.¹⁰ It is possible that newly recruited macrophages have been recruited and 'trained' by the milieu of the dysbiotic airway to either not recognise potentially pathogenic microorganisms in this environment or respond to these bacteria inappropriately which may explain the dysregulation of these cells in COPD. For example, a recent study of the transcriptome of macrophages from the COPD lung revealed a diminution of inflammatory responses.¹¹ Supporting this observation of reduced macrophage responses, we have demonstrated that COPD macrophages do not express the regulatory ligand PD-L1 in response to viral infection,¹² which may indirectly lead to increased inflammation of the COPD airway.

The scenario described above may not be limited to COPD as both macrophages¹³ and microbiome changes⁹ are also evident in many lung pathologies. However, more evidence will be required to support these speculations and, based on the work by Eguíluz-Gracia *et al.*,³ we now have a firm rationale for generating such data. The recent description of markers for macrophage subsets in the human lung may enable us to sort the resident from the recruited cells.¹⁴ When combined with our ability to analyse macrophage gene expression and the lung microbiome to a greater level than ever before, we are on the brink of beginning to understand these complex interactions between macrophage, disease and microbiome and what they may mean for disease pathogenesis and progression.

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