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A cross-circulatory platform for monitoring innate responses in lung grafts

Glorion Matthieu

Foch Hospital

Pascale Florentina

Foch Hospital

Estéphan Jérôme

Université Paris-Saclay, INRAE, UVSQ, VIM

Huriet Maxime

Université Paris-Saclay, INRAE, UVSQ, VIM

Gouin Carla

Université Paris-Saclay, INRAE, UVSQ, VIM

Urien Céline

Université Paris-Saclay, INRAE, UVSQ, VIM

Blanc Fany

Université Paris-Saclay, INRAE, AgroParisTech, GABI

Rivière Julie

Université Paris-Saclay, INRAE, AgroParisTech, GABI

Richard Christophe

Université Paris-Saclay, UVSQ, INRAE, BREED

Gelin Valérie

Université Paris-Saclay, UVSQ, INRAE, BREED

De Wolf Julien

Foch Hospital

Le Guen Morgan

Foch Hospital

Magnan Antoine

Foch Hospital

Roux Antoine

Foch Hospital

Schwartz-Cornil Isabelle (✉ isabelle.schwartz@inrae.fr)

Université Paris-Saclay, INRAE, UVSQ, VIM

Sage Edouard

Foch Hospital

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Abstract

Lung transplantation is the only curative option of end-stage chronic respiratory diseases. However the survival rate is only about 50% at 5 years. Whereas experimental evidences support that innate allo-responses impact on the clinical outcome, the knowledge of the involved mechanisms is limited. Here, we evaluate a cross-circulatory platform for monitoring the early recruitment and activation of immune cells in an extracorporeal donor lung by coupling blood perfusion to cell mapping with a fluorescent marker in the pig, a commonly-used species for lung transplantation. The perfusing pig cells were easily detectable in lung cell suspensions, in broncho-alveolar lavages and in different areas of lung sections, indicating infiltration of the organ. Myeloid cells (granulocytes and monocytic cells) were the dominantly recruited subsets. Between 6 and 10 h of perfusion, recruited monocytic cells presented a strong upregulation of MHC class II and CD80/86 expression, whereas alveolar macrophages and donor monocytic cells showed no significant modulation of expression. Altogether the cross-circulation model permits to monitor the initial encounter between perfusing cells and lung graft, in an easy, rapid, and controllable manner, for generating robust information on innate response and testing targeted therapies for improvement of lung transplantation outcome.

Introduction

Allogeneic lung transplantation (LT) is the sole therapeutic option of terminal chronic respiratory diseases. While LT practice is increasing worldwide, the outcomes are disappointing with a median survival of 6 years¹. The major non-infectious early complication of successful surgery is severe primary graft dysfunction (PGD), a non-cardiogenic pulmonary oedema syndrome that occurs in the first 3 days post LT, in about 11–25% of patients². In most patients, chronic lung allograft dysfunction (CLAD) develops following series of rejection and reparation events and stands as a major impediment to long term survival³.

Several experimental works support that the innate allogeneic response occurring upon surgery strongly impacts on the clinical outcome of LT, i.e. PGD occurrence, rejection events and CLAD development^{4,5}. At the surgical step, the brutal reoxygenation of the hypoxic graft upon de-clamping, so called ischemia-reperfusion, generates huge production of reactive oxygen species, inflammatory cytokines and Damage-Associated Molecular Pattern molecules, leading to ischemia-reperfusion injuries⁵. Additional activation signaling implicates the allo-recognition through innate molecules such as CD47/CD172A and paired immunoglobulin-like receptors (PIR)/MHC class I, as shown in mouse models^{6,7}. The innate responses of alveolar macrophages (AMs), polymorphonuclear (PMNs) cells and monocytes were shown to be instrumental in the PGD pathogenesis in mouse models of allogeneic LT^{8,9}. The combination of inflammation and allo-recognition is proposed to trigger the differentiation of monocytes into inflammatory dendritic cells (DCs) that will then stimulate allogeneic T and B cell responses leading to chronic rejection^{5,6}. Altogether the knowledge of the different cell types and mechanism involved in the

allogeneic innate response of LT is fragmentary and needs to be tackled for designing suitable therapeutics, in preclinical models of better translational relevance than mice.

Pig is a highly pertinent model for lung transplantation and is commonly used in translational research in that field ¹⁰. In order to get more insight into the innate allogeneic response in LT using this species, we developed a model inspired from a cross-circulation platform that was established by a New York team for repairing damaged lungs ^{11,12}, and that was shown to be amenable to a 4 day-duration ¹³. In this platform, an extracorporeal donor lung is connected to the blood circulation of the perfusing pig, and we coupled to it to a cell mapping strategy by a systemic CFSE injection in the perfusing pig. Compared to full transplantation, this relatively simple, rapid and highly controllable model enables repeated sampling of extracorporeal lung fragments to study the cell recruitment and activation. We show here that the innate allo-immune response involves the recruitment of many cells types with the dominant implication of myeloid cells, i.e. PMN and monocytic cells (MoCs). Furthermore the expression of MHC class II and CD80/86 was systematically enhanced on recruited MoCs between 6 h and 10 h and not significantly on donor lung MoCs and AMs. Therefore the cross-circulation platform coupled to cell mapping permits to monitor the cell responses of recruited cells upon innate allo-stimulation and could be used to assess the effect of anti-inflammatory and immunomodulatory drugs.

Results

1. Extracorporeal perfusion of lungs with an allogeneic cross-circulation platform maintains lung functions and structures over 10 h.

For analyzing the initial donor:recipient cell encounter in lung allografts, we adapted the cross-circulation platform that was developed recently in pigs ^{11,12,14}. To generate ischemia-reperfusion in allogeneic conditions, we used outbred donor and perfusing pig of different sex that originated from independent families and the donor lungs were explanted upon circulatory death with about 80 min warm ischemia. The perfusion was done by canulating the superior vena cava of the recipient and connecting the blood circulation to an extracorporeal circuit warmed at 37°C that perfuses the donor lung, as described in Fig. 1 and Supplemental 1. For cell mapping purposes, we proceeded to an i.v. injection of CFSE in the perfusing pig about 30 min before initiating the perfusion in order to label most circulating leukocytes cells ¹⁵. The static compliance, $\Delta PO_2:FiO_2$ and pulmonary vascular resistance laid within effective ranges and remained stable in the graft throughout the whole procedure (Fig. 2a-d). In the perfusing pigs, vital parameters (heart beats, blood pressure, creatinine, lactate, glucose levels) remained within normal values (Supplemental 2). Blood counts slightly varied with time within the normal range (Supplemental 2). Gross appearance remained normal, without signs of pulmonary oedema or hemorrhage (Supplemental 3). Cell viability was evaluated with exclusion of DAPI on dissociated lung cell preparations and indicated that the cross-circulatory process preserved cellular viability over time (Fig. 2e). Conventional blinded histopathological assessment revealed the preservation of airway and parenchymal structures over time, no sign of oedema and a weak score of infiltration with inflammatory polymorphonuclear cells (PMNs), mainly around blood vessels (Fig. 2f).

The ischemia-reperfusion in the allogeneic context of a cross-circulatory platform with CFSE injection induced minimal changes in the lung tissue structure, and maintained the hemodynamic, respiratory, and vital functions.

2. The perfusing CFSE⁺ cells in the ex vivo lung during cross-circulation are recruited to various lung territories.

We next analyzed the distribution of the CFSE⁺ cells in the perfusing pig blood and in different donor lung compartments. Flow cytometry analysis of the blood cells showed that the majority of the cells (> 80%) were CFSE-labelled 1 h after initiation of cross-circulation (Fig. 3a). This systemic staining slowly decreased over time, with a majority of cells being still stained after 8 h (Supplemental 4). Cell suspensions from enzymatically dissociated lung fragments collected at 0, 6, 10 h post-initiation of cross-circulation were analyzed by flow cytometry. Lungs had been flushed with saline at 0 and 10 h but not at 6 h, to avoid perturbing the perfusion. Figure 3b shows that the % of CFSE⁺ lung cells reached 31.14% ± 8.03 at 6 h and 24.78% ± 6.31 at 10 h (no significant difference between 6 h and 10 h). The lack of increase of the CFSE⁺ cell fraction between 6 and 10 h could be related to the omitted flushing at 6 h combined with a decrease in CFSE intensity at later timing. CFSE⁺ cells were also found in the bronchoalveolar lavages (BAL) at 10 h, indicating that some of the recruited CFSE⁺ cells proceeded to diapedesis through lung vascular barriers (Fig. 3c). Note that the vast majority of cells in BAL cells are alveolar macrophages (AMs) which are from the donor and are therefore CFSE⁻. The CFSE signal was sufficiently high for FACS detection but not for detection with microscopy. We used anti-FITC sheep IgG to amplify the CFSE signal (supplemental 5) and we found CFSE⁺ cells dispersed throughout the lung parenchyma on cryosections (Fig. 3d).

Therefore, cell tracking with CFSE staining shows that recipient cells are recruited to various lung territories during the cross-circulation.

3. Myeloid subsets are the main cell types recruited upon cross-circulation.

We proceeded to multiparameter FACS analyses of the recruited CFSE⁺ cells in order to analyze their composition in different myeloid and lymphoid cell subsets (Fig. 4, Supplemental 6). The dominant subset among CFSE⁺ cells in both groups was PMNs (44.02 ± 5.02% at 6 h and 42.92 ± 11.33% at 10 h) followed by SSC-A^{lo}CD172A^{hi} cells (9.62 ± 2.34% at 6 h and 9.23 ± 2.38% at 10h) that are monocytic cells (MoCs) in the pig¹⁶. Upon their recruitment in lung, monocytes from blood could start their differentiation into macrophages, a status which cannot be appreciated with the current staining, therefore we will keep the designation of these cells as MoCs¹⁷. Other subsets represented less than 10% of the CFSE⁺ cells and included SSC^{lo}CD172A^{int} cells (not known population), CD13⁺ cells previously shown to represent conventional dendritic cell type 1 (cDC1,¹⁸), NKp46⁺ cells (natural killer), CD21⁺ cells (mature B-cells¹⁹), and CD4⁺CD3⁺ and CD8⁺CD3⁺ T-cells.

The cross-circulation platform reveals that PMNs and MoCs are the main populations of recruited cells in lung at the early 6 h and 10 h time points.

4. The recruited MoCs from the perfusing pig upregulate their expression of MHC class II and CD80/86, whereas donor MoCs and AMs showed no significant modulation of expression.

Expression of MHC class II and CD80/86, which are major molecules driving antigen presentation, can be upregulated on monocytes upon ischemia-reperfusion²⁰. In addition monocytes and AMs have been shown to play an important role in the ischemia-reperfusion response and PGD in a mouse model^{9,21} and these cells are expected to play a role in antigen-presentation leading to rejection^{5,6,22}.

In order to test whether the cross-circulation platform could reveal MoC and/or AM activation, we analyzed MHC class II and CD80/86 modulation of expression on MoCs and AMs at 6 h and 10 h in the perfused lung. When focusing on the recruited CFSE⁺ MoCs, we found that the percent of MHC class II⁺ cells increased from 21.8 ± 7.1% at 6 h to 38.6 ± 3.04% at 10 h (p = 0.008) (Fig. 5a left, and Supplemental 7). Similarly, the percent of CD80/86⁺ cells in CFSE⁺ MoCs cells increased from 11.41 ± 5% at 6 h to 24.26 ± 5% at 10 h (p = 0.002, Fig. 5b left and Supplemental 7).

We performed a similar analysis on the CFSE⁻ MoCs. CFSE⁻ MoCs originate from the donor and possibly from some CFSE⁻ recruited cells. On CFSE⁻ MoCs, the upregulation of MHC class II and of CD80/86 was non-statistically significant between 6 h and 10 h (Fig. 5, middle panels, and Supplemental 7), although some non-statistically significant upregulation related to inter-animal variability was observed. This finding indicates that donor MoCs do not consistently upregulate the antigen presentation molecules, at the difference with the recruited CFSE⁺ MoCs. In the AMs (SSC^{hi}CD172A^{hi}CD163^{hi}, Supplemental 8) all express MHC class II and CD80/86, therefore geometric mean intensities were used for illustration and their expression was not modified by cross-circulation (Fig. 5, right panel).

Finally, we checked that the upregulation of the MHC class II and CD80/86 molecules on the perfusing pig SSC^{lo}CD172A^{hi} MoCs was well induced by the cross-circulation process in the lung and not by a non-specific effect of the non-biological surfaces of the circuit. Indeed, in a pig blood connected to the circuit without lung, expression of MHC class II and CD80/86 was not increased, neither on CSFE⁺ and nor on CFSE⁻ MoCs, at 6 and 10 h (Supplemental 9).

Overall, these data show that perfusion in lung grafts induces a significant upregulation of antigen presentation molecules on the recruited MoCs and not on donor MoCs and AMs.

Discussion

We show here that the cross-circulatory platform using pig lung *ex vivo* is a potent method for dissecting the cellular recruitment and response to the initial encounter between donor and host in the lung, during extended periods of time, in the pertinent pig preclinical model of LT. In particular we showed that myeloid

cells, namely PMNs and MoCs, are the main recruited cell types. Furthermore, we found that recruited MoCs strongly and rapidly upregulated the MHC class II and CD80/86 molecules upon the reperfusion in the allogeneic lung, and not so on donor MoCs and AMs. Thus recruited MoCs from the recipient rapidly acquire antigen-presentation molecules and therefore could play an important role in generating subsequent T- and B-cell allo-responses. The effectiveness of treatments controlling this activation could be evaluated in future studies using this cross-circulatory platform.

The pig preclinical model is commonly used in LT for different reasons: i) pig and human share similar lung anatomy (similar number of bronchial generations, histology, size) ²³, ii) they share similar physiology (digestion, nictemeral rhythm) and iii) the porcine immune system resembles human for more than 80% of analyzed parameters in contrast to the mouse that shares about 10% ¹⁰. Rodent models, despite their incomparable advantages offered by the targeted gene mutants for mechanistic studies, present anatomic limitations (size and specificities) as well as immunological properties leading to tolerance of allografts and less need for immunosuppressive drugs, therefore they do not generally serve as appropriate models for LT ²⁴. The first surgical technique of LT was established in the pig model as well as improvements of the surgical practice and evaluation of immune suppressive therapies ²⁴, therefore the cross-circulatory platform is highly pertinent for translational research in LT. Furthermore the cross-circulatory platform in pigs reveals to be a robust method due to the experimental controllability of the system. The tight regulation of the vascular flow during the process and the controlled ventilation probably explains the high degree of reproducibility regarding cell recruitment and activation parameters across experiments. Furthermore, it offers a direct and continuous accessibility to the perfused lung for regular sampling. It is also simple and fast to perform as compared to a complete LT, with little to no suffering of the perfusing animal that could be re-used for other purposes after the experiment, provided that the exposure to allogeneic lung would not generate biases. All these qualities and advantages would not be obtained with complete LT in rodent or pig models that induce important surgical stress, hemodynamic constraints, and require high levels of technicality and man-power support, especially in case of pig LT. Cross-circulation has been initially demonstrated to repair damaged lung tissue and maintain extracorporeal lungs healthy for extended periods of time up to several days by a team at Columbia university, NY, USA ^{11,13,14}. Therefore the use of this platform for studying the first donor:recipient encounter could be extended to much longer durations. The settings of the Columbia team slightly differed from ours regarding pig strain, donor death (beating heart vs circulatory death in our case), heating method of the circuit, anesthetic regimen (halogenated volatile versus propofol maintenance in our case) and blood reservoir. We checked that the circuit and blood reservoir had no effect on the antigen-presenting molecule expression on the perfusing pig MoCs (Supplemental 9). The Columbia team also systematically treated the recipient pig with corticosteroids, what we did not do, and in absence of corticosteroids, we found that the lung respiratory function and structures, global hematological parameters and cell viability were not altered at least for 10 h.

The cross-circulatory platform coupled to cell mapping permitted to document that many cell types were recruited to the allogeneic lung, mainly PMNs and MoCs and also NK cells, T-cells, B-cells and cDC1. We

proceeded to cell labeling exclusively on freshly isolated cells in order to avoid bias related to cell freezing. Therefore our antibody panel was adjusted to feasibility and for instance, it did not include the complex mAb combination for cDC2 detection¹⁶. Importantly, the single cell RNA-seq technique could be particularly suitable to further study the cell recruitment and activation in the first encounter between donor and recipient cells. Indeed, as we used female perfusing pigs and male donor, the donor origin can be deduced from genes expressed by the Y chromosome. Using scRNA-seq, discrete cell types could be identified and well characterized with much larger signatures than what can be done with classical cytometry. In addition modulated functions and pathways in the recruited cells and donor cells could be inferred from differentially expressed genes, thereby providing a powerful system to finely investigate ischemia-reperfusion response in the allogeneic context not only in immune cell types but also in epithelial, fibroblastic, lymphatic and endothelial cells of the graft.

Here we particularly focused on the induction of the expression of MHC class II and costimulatory CD80/86 on monocytes and macrophages, given the place of these cell types in the initiation of acute and chronic rejection of allografts^{22 25}. Of note, we could not detect upregulation of MHC class II on the recruited PMNs (data not shown). This activation of recruited MoCs may lead to their differentiation in MPs and/or Mo-derived DCs, becoming potent antigen-presenting cells (APCs) for initiation of the T- and B-cell allo-responses²⁶. Therefore the cross-circulation platform could be used to test the effectiveness of induction drugs -which are administered at the surgical step- on this recruited monocyte activation. Indeed these drugs which have not been designed to target innate immunity, include corticosteroids, calcineurin inhibitor, and/or mycophenolate, and their effects on mononuclear cells are not known and could be investigated with this platform²⁷. Indeed a recent publication demonstrates in mice that targeting calcineurin inhibitor to myeloid phagocytes induced better protection against skin graft rejection than the conventional administration that aims at suppressing lymphocytes²⁸. New therapeutic avenues such as administration of mesenchymal stem cells²⁹ or photopheresis³⁰ could be similarly tested.

Finally, cross-circulation shall be used to monitor the cellular effects of surgical practices. As said above, at the difference with LT, ventilation and perfusion are highly controllable upon cross-circulation; therefore the detrimental effects on the initial donor:host cell response of forced ventilation as ventilator-induced lung injury³¹ and high flow reperfusion rate upon de-clamping³² could be analyzed in that model and thereby lead to adjustments of practices.

Overall we report on an original, relatively easy, rapid and controllable in vivo technique to study the response to the first encounter between donor lung cells and host immune cells in the pertinent pig model for LT. We show the robustness of the technique and its suitability to monitor innate cell response in an allogeneic context over time. It opens the way to a use for evaluation of anti-inflammatory and immunomodulatory regimen and for testing surgery practice adjustment, with the ultimate objective to improve LT outcome and decreasing the burden of long-term immunosuppressive treatments.

Material And Methods

Animals.

Twenty Large-White pigs (ten donor – recipient pairs) were hosted in the Animal Genetics and Integrative Biology unit (GABI-INRAE, France). Matched pairs of male donors and female recipients from different sibling were used. Animals were 3–5 months of age, with a mean weight of 50.8 ± 4.4 kg.

Donor lung harvest and lung cannulation.

The lung harvest from donor pigs ($n = 10$) was performed in the Animal Surgery and Medical Imaging Platform (CIMA-MIMA2-BREED-INRAE, Jouy en Josas, France). Heart-lung monobloc harvests were performed using a non-heart-beating donor swine model³³. Anesthesia was induced by a combination of 1 mg/kg Rompun® 2% (Elanco, Heinz-Lohmann-Strasse 4, Cuxhaven, Germany) and 15 mg/kg Imalgene® 1000 (Boehringer Ingelheim Animal Health, Lyon, France) and pursued with 6% isoflurane. A 25,000 U heparin bolus (Sanofi, Paris, France) was administered i.v.. Pigs were euthanized with 50 mg/kg Dolethal® (Vetoquinol, Magny-Vernois, France). After cardiorespiratory arrest, a period of 10 min of no touch was applied. After median sternotomy, a cannula was placed and secured in the main pulmonary artery (PA). Lungs were flushed with 3 L of Perfadex® (XVIVO Perfusion, Göteborg, Sweden) administered in a cold anterograde perfusion. The heart-lung block was explanted with inflated lungs to a sustained airway pressure of 15 cmH₂O, and the trachea was stapled (Endo GIA device, Medtronic, Dublin, Ireland). The block was placed on ice. The heart was removed, leaving behind a circumferential left atrial cuff. Dedicated cannula (XVIVO Perfusion) were sutured to the left atrium with 5 – 0 Prolene® (Ethicon, Somerville, NJ, USA). The arterial cannula was fixed to the PA with a purse string of Mersutures® 1 (Ethicon). A cold retrograde flush with 1 L Perfadex® (XVIVO Perfusion) was performed. Lungs were placed in a sterile isolation bag with 500 ml Perfadex® and stored at 4°C. The mean duration of warm ischemia between euthanasia and cold storage was 82.8 ± 9.3 min. The mean duration of cold ischemia between cold storage and cross circulation initiation was 103.9 ± 15.6 min.

Perfusing Pig Conditioning

Pigs ($n = 10$) were anesthetized as described above. Septotryl® (0.08 ml/kg) (Vetoquinol) was injected i.m. prior to catheterization. A femoral arterial line (Arrow International, Cleveland, Ohio, USA) was placed percutaneously under ultrasound guiding for haemodynamic monitoring (Supplemental 2). A 25,000 U heparin bolus was administered. The superior vena cava was cannulated with a 20 F double lumen canula (Avalon Elite, Maquet Cardiopulmonary, Rastatt, Germany) using the Seldinger percutaneous technique¹⁴. The correct positioning of the wire and cannula in the inferior vena cava were checked by ultrasound (Supplemental 1). Pig was positioned on the left lateral decubitus with a heating blanket to avoid hypothermia. Physiological parameters, including heart rate, electrocardiogram, blood pressure, oxygen saturation, end-tidal CO₂, temperature, and respiratory rate were continuously monitored. About 30 min before the initiation of cross-circulation, 25 mg carboxyfluorescein succinimidyl ester (CFSE) (Sigma-Aldrich, Saint-Louis, MS, USA) was administered i.v. in the perfusing pig, diluted in 4 ml DMSO + 40 µl heparin. Sedation was maintained for 10 h by continuous administration of 2–4 mg/kg/h propofol

(Proposure®, Axience, Pantin, France) + 0.6% isoflurane and analgesia was obtained by administration of 0.2 mg/kg nalbuphine i.v. every 3 hours.

Cross-circulation.

We basically followed the procedure described in ^{12,14}. The circuit was filled with 1.5 L ringer lactate (Vetivex®, Dechra Northwich, UK). As shown in Fig. 1, the circuit consists in a main console (SCPC centrifugal pump console, LivaNova, London, UK), a disposable pump (Revolution centrifugal pump, LivaNova), a hard-shell reservoir (LivaNova) and three 8-inch tubing (Smart coated tubing, LivaNova). A EOS® oxygenator (LivaNova) was used for heating the circuit. Gas connection was occluded. The PA and pulmonary vein (PV) pressures, the PA flow, and the temperature data were continuously monitored. The perfusing pig was maintained on a continuous heparin infusion (100 U/Kg/h). The activated clotting time was measured using a IStat® kit (Abbott, Chicago, IL, USA) and the heparin drip was adjusted to maintain a target clotting time value of 150–200 sec. Donor lungs were placed in dorsal position on an XVIVO® chambers (XVIVO Perfusion) and the trachea was cannulated with a 7.5 mm diameter cuffed endotracheal tube (Mallinckrodt, Staines-upon-Thames, UK). The tubing was spliced to connect the perfusing pig to the dedicated circuit, marking the start of cross-circulation. Initial flow rates were set to 5% of the estimated cardiac output and were gradually increased to 10% with an initial PA target pressure below 15 mm Hg and a PV pressure of 3–5 mm Hg. The ventilation (Elisée 350 Resmed San Diego, USA) was initiated within the first 10 min with the following initial settings: volume control mode, 10/min respiratory rate, 6 mL/kg tidal volume, 5 cm H₂O positive end-expiratory pressure, and 21% FiO₂. Atelectatic lung regions were recruited by increasing the tidal volume and the positive end-expiratory pressure and by performing inspiratory hold maneuvers (up to 25 cm H₂O). PV was dependent on the hydrostatic pressure difference between the lungs and reservoir. Reservoir blood level was dependent on the hydrostatic pressure difference between the reservoir and the swine recipient. These two values were controlled by adjusting the height difference between the lung, the reservoir and the swine host ¹³.

Extracorporeal haemodynamic and lung function monitoring.

Blood samples were collected from the main PA and PV cannula every 1 h, and hemo-gas analysis was performed using a Istat® kit. Static compliance (Tidal Volume/(plate pressure – positive end expiratory pressure)), Δ PCO₂ (arterial PCO₂ – venous PCO₂) and Δ PO₂/FIO₂ ((venous PO₂ – arterial PO₂) / FIO₂) were calculated every 1 h. The transpulmonary pressure (PA - PV) and the pulmonary vascular resistance were calculated ((PA pressure – left atrial pressure) x 80 / flow rate).

Blood counts and biochemical monitoring.

Blood samples were collected by venipuncture of the auricular vein or directly from the extracorporeal circuit after cross-circulation. Plasma was analyzed immediately for biochemical profiling. Blood count and biochemical profiling were performed on a MS4.5 analyzer and a M-Scan II analyzer (Melet Schloesing Laboratoires, Cergy-Pontoise, France).

BAL and lung biopsies.

BAL was performed before cross-circulation in the subsegmental bronchi of the azygos lobe (0 h) and after 10 h cross-circulation in the cranial and caudal lobes. Lung biopsies for cell dissociation (about 2 g) were sampled in the cranial and caudal lobes using a surgical stapler (Endo GIA™ universal stapling system, Medtronic, Minneapolis, USA) and were immediately immersed in cold hypothermic preservation media (HypoThermosol® FRS, Stemcell Technologies Inc, Vancouver, Canada). For immuno-histo-fluorescence, biopsies (about 5 mm³) were snap-frozen in a matrix gel (Sakura, Paris, France). For histology, biopsies were fixed in cold phosphate-buffered 4% paraformaldehyde for 24 h and subsequently paraffin-embedded.

Lung cell extractions.

Tissues (2 g) were minced and incubated for 45 minutes at 37°C on a rotary shaker in RPMI 1640 supplemented with 100 IU/ml penicillin, 100 µg/ml streptomycin, 2 mM L-glutamine and 10% inactivated fetal calf serum (FCS) (all from Invitrogen, Paisley, UK), containing 3 mg/ml collagenase D, 0.25 mg/ml Dnase I (Sigma-Aldrich) and 0.7 mg/ml dispase II (Gibco®, ThermoFisher Scientific, St Aubin, France). The minced preparation was crushed and filtered on a nylon mesh (1 mm diameter) and filtered through successive cell strainers (500 µm, 100 µm, 40 µm). Red blood cells were lysed with erythrocytes lysis buffer (10 mM NaHCO₃, 155 mM NH₄Cl, and 10 mM EDTA). After a wash in PBS, 10⁸ cells were used for cell surface staining. The rest of the cells was frozen in FCS + 10% DMSO in a Mister Frosty freezing container (Nalgene, Rochester, NY, USA) and kept in liquid N₂.

Cell surface staining and flow cytometry.

Cell surface staining was performed in RPMI + 10 mM Hepes supplemented with 5% horse serum and 5% swine serum (Gibco, Life Technologies Europe, Bleiswijk, Netherlands). Primary and secondary Abs and their working dilutions are listed in Supplemental 5. Matched isotype controls for mouse IgG1, IgG2b, and IgG2a were used at the same concentration as the corresponding mAbs of interest, using the fluorescence minus one method³⁴. In cases of third step labelling with a directly conjugated mAb of the same IgG1 isotype as the primary Ab (anti-CD163-RPE and anti-CD3-RPE), an excess of mouse IgG1 (50 µg/ml) was used in an additional saturation step. Dead cells were excluded by DAPI staining (Sigma-Aldrich). Samples were acquired on BD LSR Fortessa™ Cell Analyzer (BD-Biosciences). Acquired data were analyzed using FlowJo software (version 10.7.1; Tree Star, Ashland, OR, USA).

Histology.

Formalin-fixed paraffin-embedded lung tissues at 0, 6, 10 h were sectioned every 50 µm for generating six 5 µm tissue slices per sample and stained with hematoxylin-eosin-saffron (HES). The slides were imaged with a slide scanner (Pannoramic SCAN II, v3.0.2, 3DHitech, Medipixel Ltd, Budapest, Hungary) and analyzed by an external pathologist and a veterinarian in a blinded fashion. Five randomly-selected high

power fields ($7 \times 10^4 \mu\text{m}^2$ area) from 6 slides per sample were observed and scored by quantification of airway and alveolar polymorphonuclear cells and interstitial edema according to a reference scoring ¹¹.

Immunohistofluorescence.

Cryosections (10 μm) of lung parenchyma frozen biopsies were obtained using a

cryostat (Leica CM3050S, Nanterre, France). Sections were fixed in methanol/acetone (1:1) at -20°C for 20 min and stained with sheep IgG anti-FITC IgG in order to amplify the CFSE signal, followed by donkey anti-sheep IgG-A594 and with matched sheep IgG controls (Supplemental 5). Sections were stained with DAPI and mounted in SlowFade medium (Invitrogen). The sections were scanned at a x 20 magnification with the Panoramic SCAN II, v3.0.2.

Statistics

Data were analyzed with the GraphPad Prism 7.0 software. After subjecting the data to a normality test, a paired parametric two-tailed t-test was used to compare values between different time points. When the data did not present a normal distribution (Supplemental 2, vital parameters), a non-parametric Wilcoxon signed rank test was used.

Study Approval

The animal experiments were conducted in accordance with the EU guidelines and the French regulations (DIRECTIVE 2010/63/EU, 2010; Code rural, 2018; Décret n°2013 – 118, 2013). The experiments were approved by the COMETHEA ethic committee under the APAFIS number authorization 25174-2020011414322379 and were authorized by the French “ministère de l’enseignement supérieur et de la recherche”. The authors complied with the ARRIVE guidelines.

Abbreviations

Lung transplantation (LT), alveolar macrophages (AMs), monocytic cells (MoCs), dendritic cells (DCs), polymorphonuclear neutrophils (PMNs), primary graft dysfunction (PGD), carboxyfluorescein succinimidyl ester (CFSE), pulmonary artery (PA), pulmonary vein (PV), swine (sw), human (hu), broncho-alveolar lavage (BAL), isotype control (ISC), single cell RNA-seq (scRNA-seq), hematoxylin-eosin-saffron (HES).

Declarations

Data availability statement

The data generated during this study are included in this article or in the supplemental file. In case of needed complementary information or raw data, they are available from the corresponding author.

Competing interest

Authors have no financial competing interest.

Author contribution

G.M.: PhD training, participated to the design of the study, performed the pig surgery, conducted experiments, analyzed data, performed statistical analyses, prepared several figures.

P.F.: participated to the pig surgery, conducted most cell/molecular/histological analysis experiments, analyzed data, performed statistical analyzes, prepared the majority of figures.

E.J.: participated to pig surgery, collection of material, and cell analysis experiments

H.M.: participated to pig surgery, collection of material, cell analysis experiments, generated one of the figures.

G.C.: performed biochemical measurements and cell isolations

U.C.: performed histo-immunofluorescence experiments and cell isolations

B.F.: performed biochemical measurements

R.J.: acquired histo-immunofluorescence images

R.C.: managed the surgery platform, participated to the ethic application

G.V.: support for surgery, organization of material

D.W.J.: performed the pig surgery and participated to the design of the study

L.G.M.: participated to the design of the study, supervised the long anesthesia, and participated to data analyses, edited the manuscript

M.A.: participated to the design of the study, to the data analyses, edited the manuscript

R.A.: participated to the design of the study, to the data analyses, edited the manuscript

S.-C.I.: participated to the design of the study regarding the cellular aspects, initiated and supervised the immunological/histological/immuno-histo-fluorescence analyses, analyzed the data, prepared several figures, participated to the ethic application, wrote the manuscript

S.E.: conceptualized the study, raised funding, participated to the pig surgery, analyzed data.

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Figures

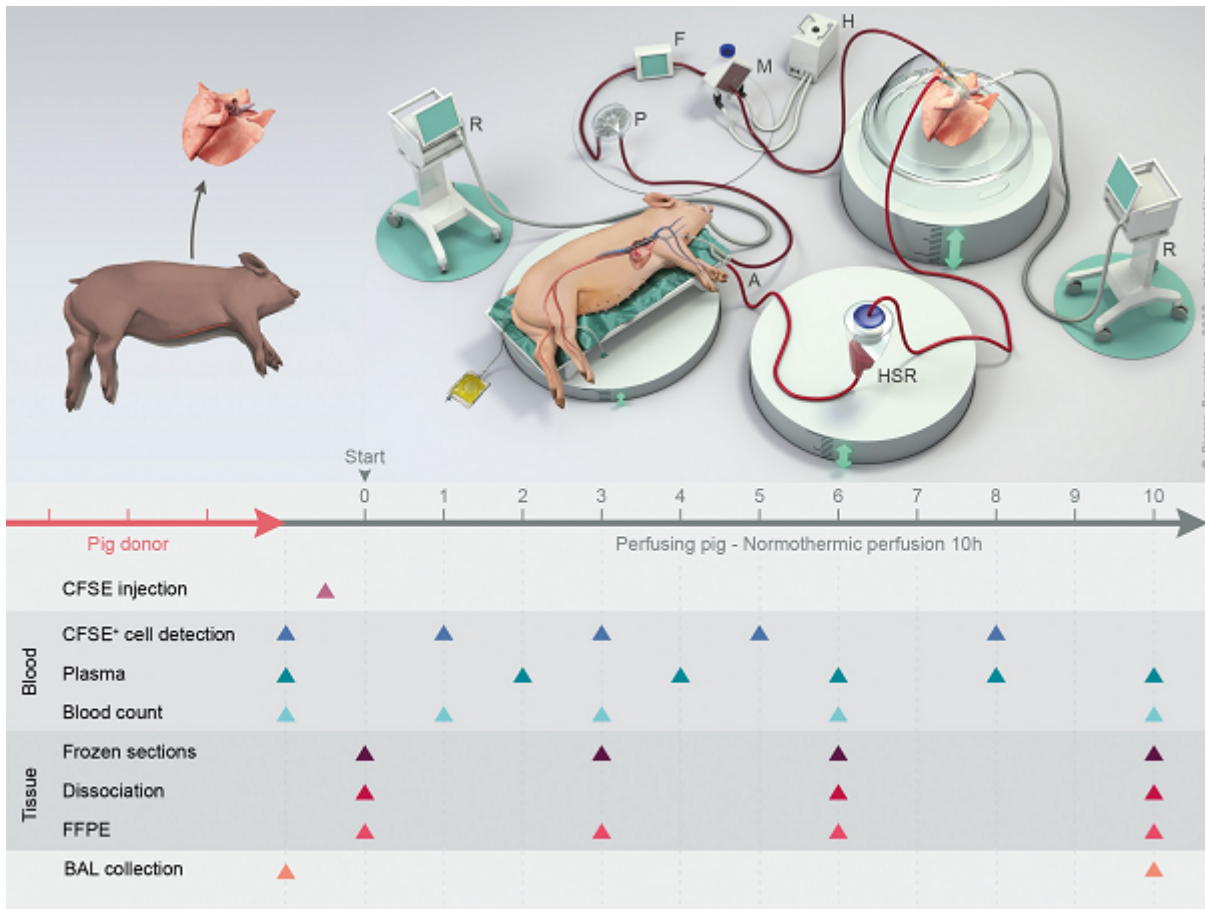


Figure 1

Figure 1

Experimental overview of the cross-circulation model for cell mapping. **Top.** Setup of the cross-circulation technique. A double lumen canula, HSR hard-shell reservoir, R ventilator, H heater, M membrane, F flow-meter, P pump. **Bottom.** Experimental time-line of injection procedures and samples collections. CFSE (25 mg) was injected 30 min before cross-circulation. Blood, broncho-alveolar lavage (BAL) and lung tissue biopsies were collected at different time points. Blood was used for the detection of CFSE⁺ cells in the perfusing pig, for biochemical profiling in plasma, and for blood cell counts. Lung tissue was used for producing frozen sections (location of CFSE⁺ cells in the tissue), for enzymatic dissociation (CFSE⁺ cells composition in the tissue), and for formalin-fixed paraffin-embedded section generation (histological analyses). BAL was used for CFSE⁺ cell detection.

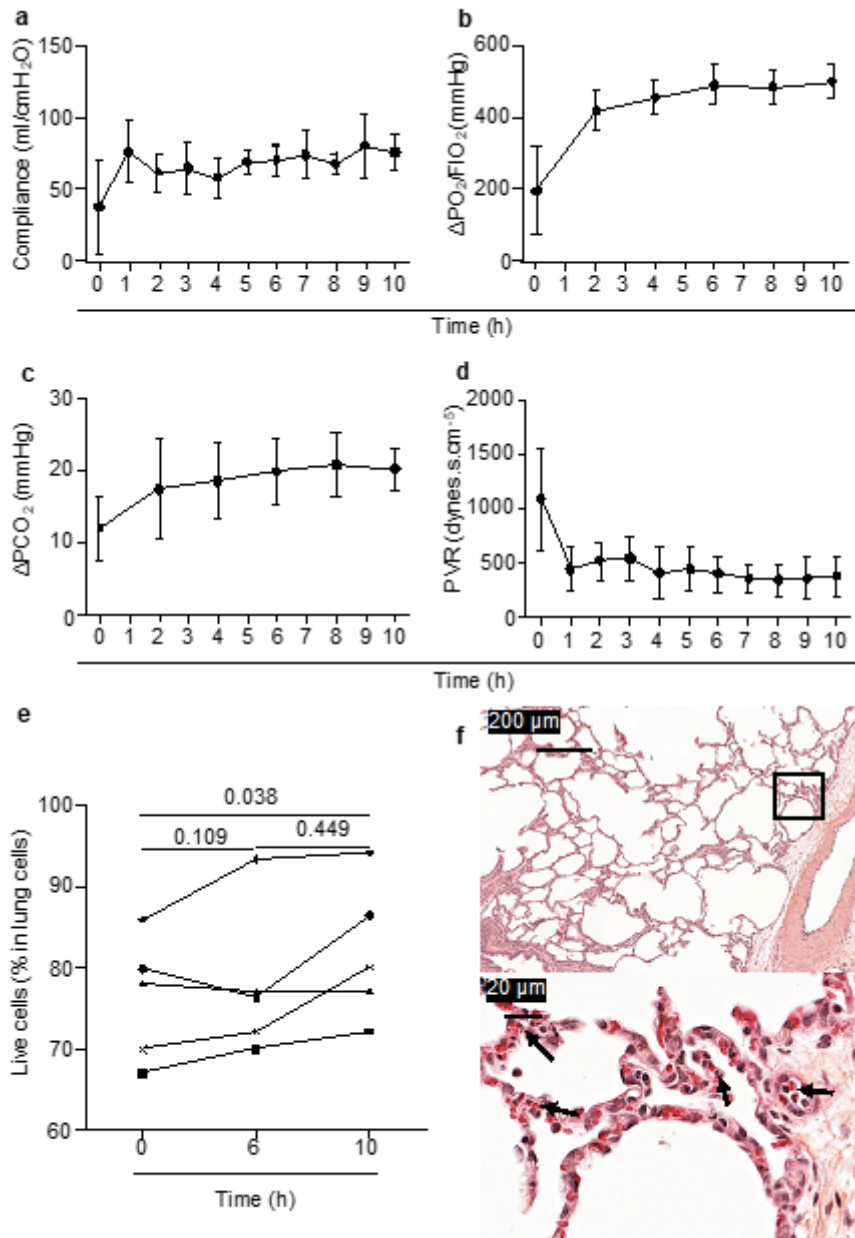


Figure 2

Figure 2

Extracorporeal lung stability and performance throughout 10 h of cross-circulation. In a, b, c, d, all values represent means \pm standard deviations (n = 5) **a.** Static compliance. **b.** $\Delta PO_2/FIO_2$: (venous PO_2 - arterial PO_2)/ FIO_2 . **c.** ΔPCO_2 = arterial PCO_2 - venous PCO_2 . **d.** Pulmonary Vascular Resistance (PVR): ((pulmonary arterial pressure - left atrium pressure) x 80) / flow rate. **e.** Live cells in total lung cells evaluated with exclusion of DAPI staining from enzymatically dissociated lung fragments collected at 0, 6, 10 h. Each pig is labelled with a unique symbol throughout the paper. After passing a normal distribution test, the values were compared between different timings with a paired t-test and the p-values are reported. **f.** Conventional histopathological assessment of the pig lung after 10 h of cross-circulation (haematoxylin and eosin staining). A higher magnification of the squared field on top is shown on the

bottom. See the weak infiltration with PMNs around blood vessels (black arrows, < 25 PMNs per high power field, 30 fields/sample).

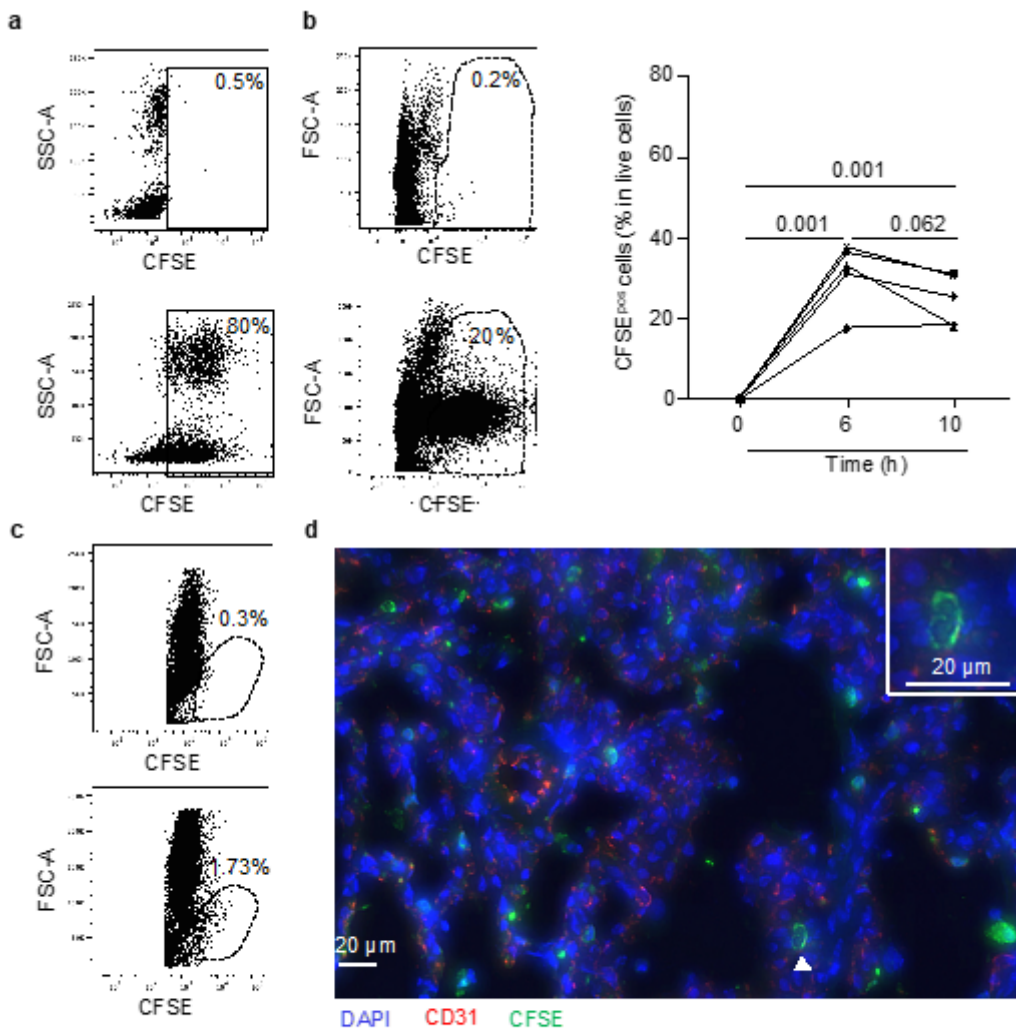


Figure 3

Figure 3

Detection of CFSE⁺ cells in the perfusing pig blood and in the lung upon cross-circulation. **a.** Blood was collected 1 h before and 1 h after the cross-circulation initiation (top and bottom respectively), and CFSE was injected 30 min. before cross-circulation initiation. Whole blood cells were analyzed by flow cytometry after lysis of erythrocytes and the % CFSE⁺ cells is indicated. **b.** Lung biopsies were collected at 0 h, 6 h and 10 h of cross-circulation and a single cell suspension was generated by enzymatic treatment. Left, the FACS profiles at 0 h (top) and 10 h (bottom) of one representative experiment are shown and the % CFSE⁺ among DAPI^{neg} live cells is indicated. Right, the proportion of CFSE⁺ cells among live lung cells is shown (5 pigs/group). Each pig is labelled with a unique symbol that is conserved throughout the paper. After passing a normal distribution test, the values were compared between different timings with

a paired t-test and the p-values are reported. **c.** BAL was collected at 0 h from the azygos lobe (top) and after 10 h of cross-circulation from the rest of the lung (bottom), stained with DAPI and analyzed by flow cytometry. The live CFSE⁺ cell population is shown. **d.** Cryosections of lung biopsies were fixed with acetone:methanol, stained with APC-conjugated anti-swine CD31 (red) and sheep anti-FITC IgG followed by A594-conjugated anti-sheep IgG in order to amplify the CFSE signal for microscopy (green). Nuclei were counterstained with DAPI (blue). The sections were scanned at a x 20 magnification with the Panoramic SCAN II. The white arrow points to a CFSE⁺ monocyte expanded at a higher magnification.

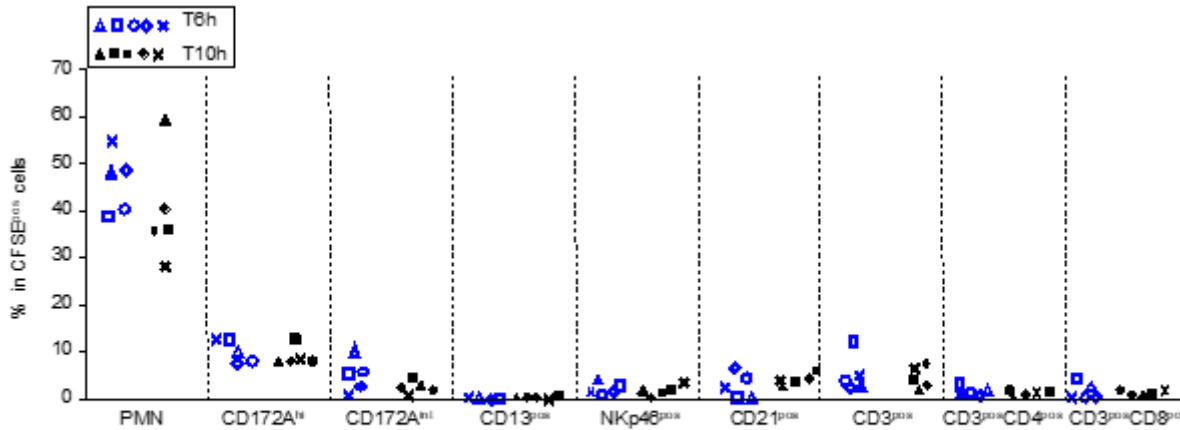
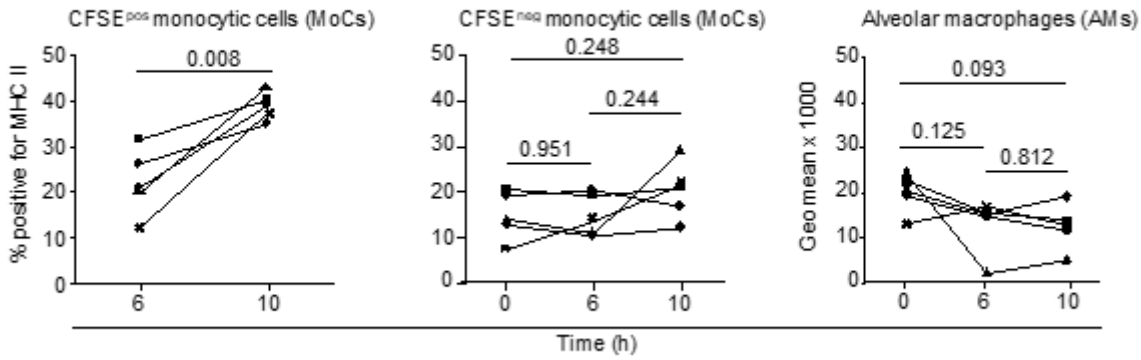


Figure 4

Figure 4

Recruitment of immune cell subsets upon 6 and 10 h of cross-circulation. Contribution of immune cell subsets in the CFSE⁺ cells. Within live CFSE⁺ gated cells, the percent of PMNs, SSC-A^{lo}CD172A^{hi} cells (MoCs), SSC-A^{lo}CD172^{int} cells, CD13⁺ (cDC1 dendritic cells), NKp46⁺ (NK-cells), CD21⁺ (B-cells), CD3⁺ T-cells, CD3⁺CD4⁺ T-cells, CD3⁺CD8⁺ T-cells are reported (4 pigs per group). Each pig is labelled with a unique symbol throughout the paper. The gating strategy is presented in Supplement Fig. 6.

a. MHC II



b. CD80/86

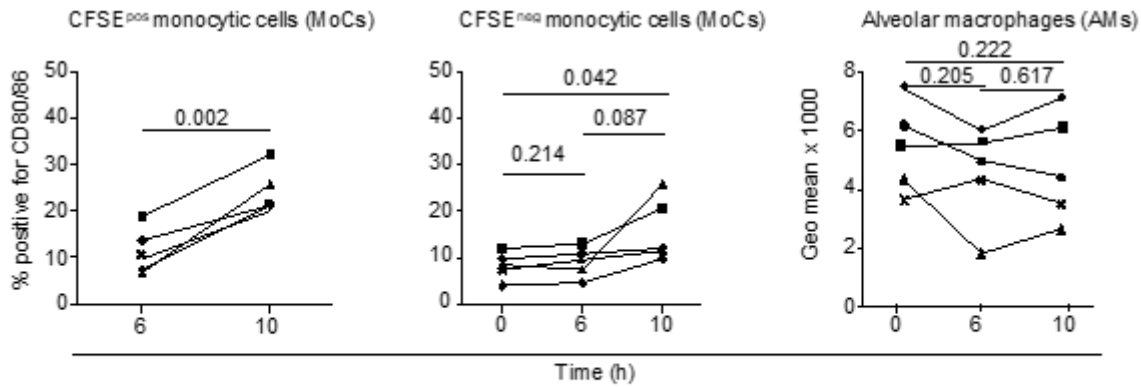


Figure 5

Figure 5

Modulation of MHC class II and CD80/86 expression on lung monocytes and macrophages upon cross-circulation a. From left to right, percent of MHC class II-positive cells within live CFSE⁺SSC-A^{lo}CD172A^{hi} cells (CFSE⁺ monocytic cells, MoCs), within live CFSE⁻SSC-A^{lo}CD172A^{hi} cells (CFSE⁻ MoCs) and expression of MHC class II (geometric mean intensity) on live alveolar macrophages (AMs, which are all positive for MHC class II) analyzed as presented in Supplement Fig. 7 and 8. **b.** Same as in a for analysis of CD80/86 expression. In a and b, each pig is labelled with a unique symbol throughout the paper, 5 pigs per group were analyzed. After passing a normal distribution test, the values were compared between different timings with a paired t-test and the p-values are reported.

Supplementary Files

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