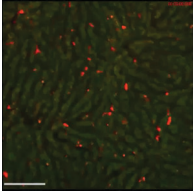
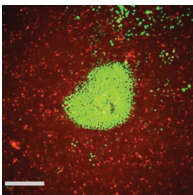


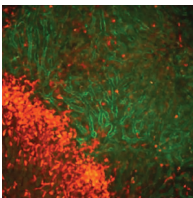
SUPPLEMENTAL MATERIAL

Dal-Secco, <http://www.jem.org/cgi/content/full/jem.20141539/DC1>

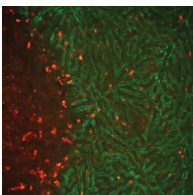
Video 1. Patrolling of the liver vasculature by CCR2⁺ monocytes under basal conditions. Under control conditions, a population of RFP⁺ cells can be observed patrolling the liver vasculature in *Ccr2^{RFP/+}* mice. Autofluorescent hepatocytes (green); RFP⁺ monocytes (red); scale bar, 100 μ m. 4D imaging confirms these cells are patrolling within the liver vasculature. Endothelium (blue); RFP⁺ monocytes (red).



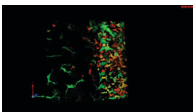
Video 2. Early monocyte response to focal sterile injury in the liver is characterized by a slow accumulation of monocytes around the lesion site. Time-lapsed SD-IVM imaging of a *Ccr2^{RFP/+}* mouse during the first 4 h after focal sterile injury in the liver shows a slow recruitment of patrolling CCR2⁺ monocytes to the injury site. Focal necrotic injury labelled with Sytox (green); RFP⁺ monocytes (red).



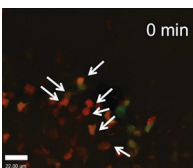
Video 3. Meandering behavior of CCR2⁺ monocytes around the site of a focal necrotic injury in the liver. SD-IVM of a *Ccr2^{RFP/+}* mouse shows that CCR2⁺ cells can be observed crawling in-and-out of a monocytic ring that forms around a focal sterile necrotic lesion 24 h after injury. Additionally, CCR2⁺ monocytes can be observed meandering around within the ring surrounding the injury. Vasculature labelled with FITC-conjugated anti-CD31 (green); RFP⁺ monocytes (red). Injury site is located in the lower left corner of the image.



Video 4. CCR2-deficient monocytes fail to accumulate around and within the site of focal necrotic injury in the liver. SD-IVM of a *Ccr2^{RFP/RFP}* mouse demonstrates that RFP⁺ cells fail to recruit to the site of injury and form a ring surrounding the lesion. RFP⁺ cells can be observed patrolling the liver vasculature distal from the injury site; however, these cells are not seen to be recruited to the lesion. Vasculature labelled with FITC-conjugated anti-CD31 (green); RFP⁺ monocytes (red). Injury site is located on the left side of the image.



Video 5. A spectrum of monocytes surround a focal sterile injury of the liver. 3D reconstruction from sequential z stacks obtained by SD-IVM of *Ccr2^{RFP/+}/Cx3cr1^{GFP/+}* mouse. Imaging of a focal sterile lesion 48 h after injury demonstrates the accumulation of a spectrum of RFP/GFP-expressing monocytes (red, orange, yellow, or green). Image compiled and rendered as 3D opacity models using the Volocity software package. Bar, 100 μ m.



Video 6. In situ conversion of monocytes recruited to a site of injury. A 2 mm x 1mm biopsy punch containing the sterile injury site was harvested from a *Ccr2^{RFP/+}/Cx3cr1^{GFP/+}* mouse 24 h after injury, maintained at 37°C and 5% CO₂, and imaged. Images of the tissue biopsy were taken every 20 min and the resulting images were compiled into a time-lapsed video demonstrating the shift from red to green in individual cells.