

Supplementary Information Titles

Journal: Nature Medicine

Article Title:	HETEROTYPIC INTERACTIONS ENABLED BY POLARIZED NEUTROPHIL MICRODOMAINS MEDIATE THROMBO-INFLAMMATORY INJURY
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Supplementary Item & Number (add rows as necessary)	Title or Caption
Supplementary Figure 1	Elevated expression of $\alpha M\beta 2$ on <i>Selp1g^{-/-}</i> leukocytes.
Supplementary Figure 2	Reduced nRBC capture in mice deficient in C3.
Supplementary Figure 3	Specificity of albumin-coated fluosphere capture <i>in vivo</i>.
Supplementary Figure 4	Fluosphere capture by adherent leukocytes correlates with the level of $\alpha M\beta 2$ expression.
Supplementary Figure 5	Inhibition of Src kinases protects from lung injury.
Supplementary Figure 6	<i>In vivo</i> detection of reactive oxygen species (ROS) in adherent leukocytes following anti-H2d infusion.
Supplementary Figure 7	N-acetyl-cysteine prevents ROS generation by adherent leukocytes and vascular permeability after anti-H2d infusion.
Supplementary Figure 8	Kinetics of leukocyte recruitment and RBC-leukocyte interactions in SCD mice.
Supplementary Figure 9	Frequency of adherent leukocyte subsets in venules of sickle transgenic mice.
Supplementary Figure 10	Leukocyte recruitment in venules of wild-type, <i>Selp^{-/-}</i> and <i>Sele^{-/-}</i> mice reconstituted with hematopoietic stem and progenitor cells from SCD mice.
Supplementary Figure 11	Effect of adhesion receptor deficiency and SCD mutation on $\alpha M\beta 2$ activity on adherent leukocytes.
Supplementary Table 1	Hemodynamic parameters in mice analyzed for RBC-leukocyte interactions
Supplementary Table 2	Hemodynamic parameters in mice analyzed for fluosphere capture.

Supplementary Methods	
Supplementary Video 1	nRBC captures are mediated by the leading edge of adherent PMNs. Adherent leukocytes in venules of C57BL/6 mice treated with TNF- α were imaged following the intravenous injection of PE-conjugated anti-L-selectin (red, 0.5 μ g) and FITC-conjugated anti-LFA-1 (clone M17/4; green, 1 μ g). L-selectin clusters identify the trailing edge of adherent leukocytes. Brightfield images of nRBC interactions with PMNs in inflamed venules were captured 180 min after cytokine administration.
Supplementary Video 2	Platelets interact mostly with leukocyte microdomains at the leading edge. Platelets were labeled by anti-CD41 (red, 1 μ g / mouse) and the trailing edge with anti-L-selectin (blue, 0.02 mg/Kg) in a TNF- α -stimulated mouse. Real time is shown in the left upper corner (h:min:s).
Supplementary Video 3	Regional activation of αMβ2 integrin at the leading edge of adherent leukocytes mediates fluosphere capture. TNF- α -treated C57BL/6 mice were injected with PE-conjugated anti-L-selectin (red; 0.5 μ g) to label the trailing edge of adherent leukocytes. Fluosphere interactions with adherent leukocytes in inflamed venules were imaged 180 min after cytokine treatment, immediately upon the injection of albumin-coated fluospheres (green) through a catheter placed in the left carotid artery.
Supplementary Video 4a and 4b	Platelet-WBC interactions are markedly induced by anti-H2d administration in Balb/c mice. Platelets were labeled by anti-CD41 (red, 1 μ g / mouse) and the trailing edge with anti-L-selectin (blue, 0.02 mg/Kg). (a) Sequence of images just before anti-H2d administration. (b) Sequence of images taken after anti-H2d injection. Note the increase in platelet captures by leukocytes and also the interactions of non-labeled RBCs with adherent leukocytes.