

Supplementary Material

Loss of zinc transporters ZIP1 and ZIP3 augments platelet reactivity in response to thrombin and accelerates thrombus formation *in vivo*

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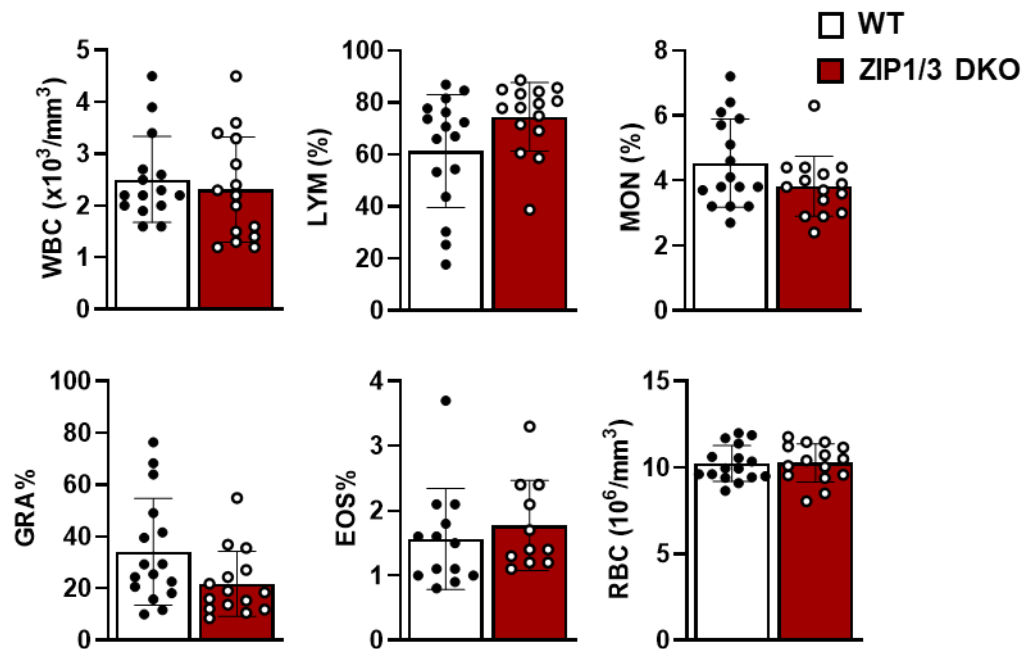
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1 Supplementary Data

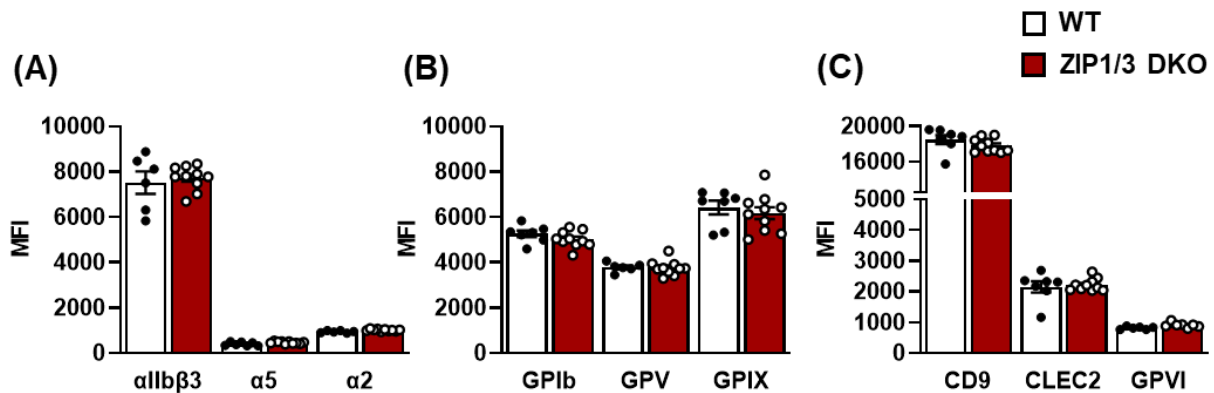
No supplementary data

2 Supplementary Figures and Tables

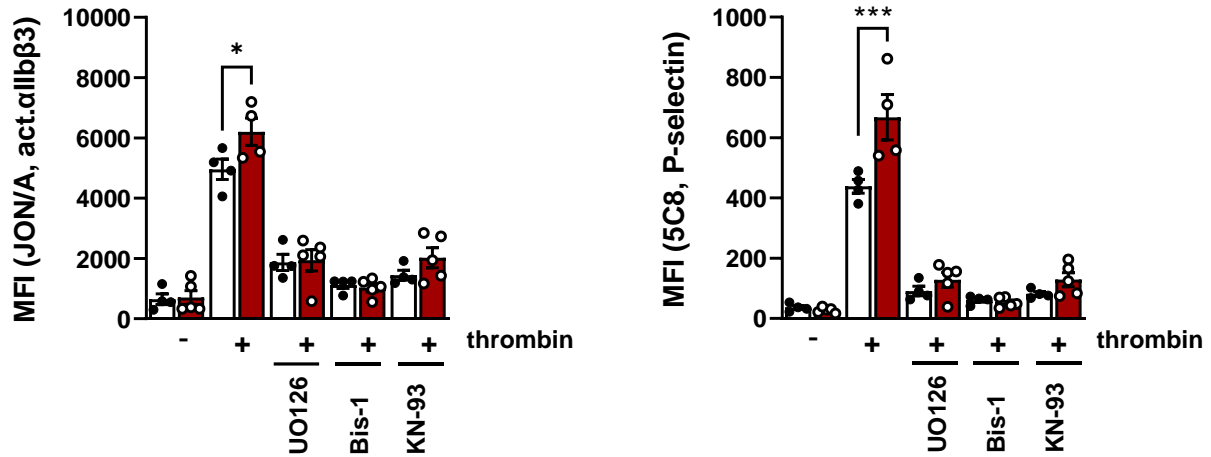
2.1 Supplementary Figures



Supplementary Figure 1. ZIP1/3 deficiency has no effect on white and red blood cell count. Analyses of blood parameters were performed from EDTA blood using a ScilVET hemacytometer. n=15-16 (Students t-test).



Supplementary Figure 2. ZIP1/3 deficiency does not affect integrin or platelet glycoprotein cell surface levels. Flow cytometric analysis of the cell surface expression of the indicated integrins (A), the GPIb-V-IX complex (B) and CD9, CLEC2, GPVI (C) using directly fluorescently-labelled specific antibodies in saturating amounts. n=6-10 (Students t-test).



Supplementary Figure 3: Influence of PKC, ERK1/2 and CaMKII inhibitors on thrombin-stimulated integrin activation and degranulation. Platelets from WT and ZIP1/3 DKO mice were pre-incubated for 1h at RT with inhibitors against PKC (bisindolylmaleimide-1 (Bis-1); 10 μ M), MEK1/2 (UO126; 5 μ M) or CamKII (KN-93; 10 μ M), activated with a threshold dose of thrombin and stained with saturating amounts of PE-coupled JON/A detecting activated $\alpha_{IIb}\beta_3$ integrin (left) and Dylight649-coupled anti-P-selectin (right); n=4-5 (two-way ANOVA + Bonferroni; *P<0.05, ***P<0.001).