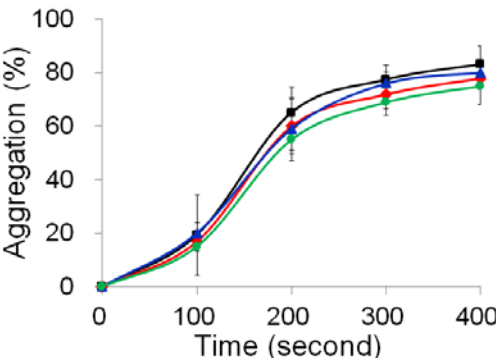
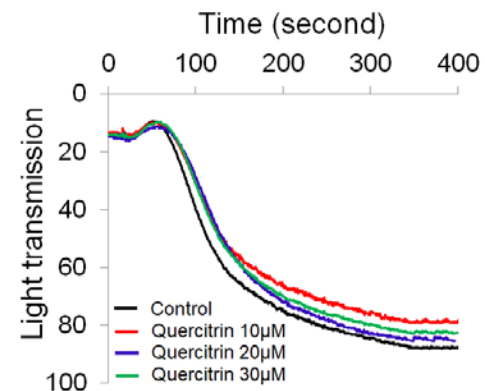
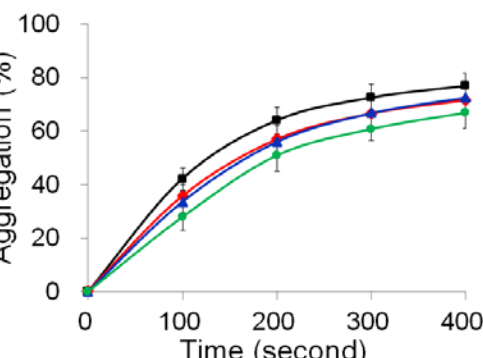
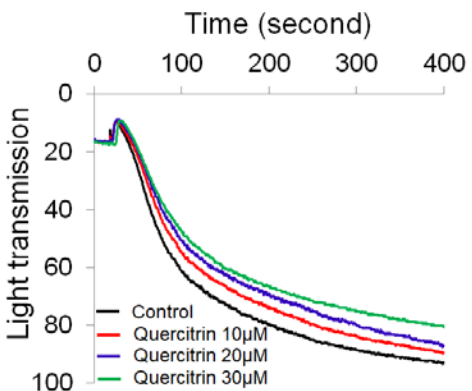


Figure S1

A) CRP 0.5 $\mu\text{g/ml}$



B) U46619 10 μM



B) Collagen 3 $\mu\text{g/ml}$

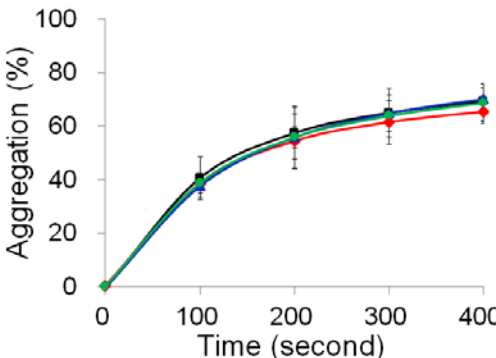
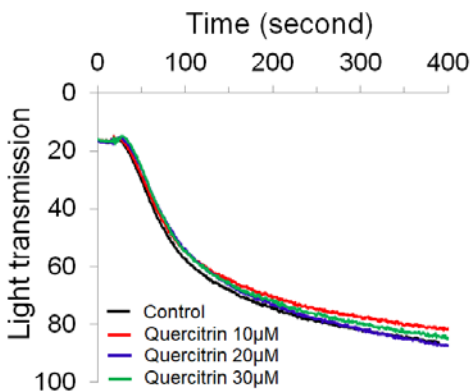
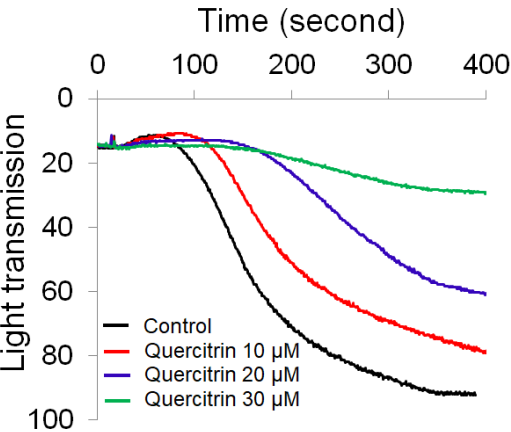


Figure S1. Role of quercitrin in human platelet aggregation induced by numerous agonists. Washed human platelets were preincubated with various concentrations of quercitrin (10, 20, and 30 μ M) for 5 minutes at 37°C and then stimulated with 0.5 μ g/ml CRP (A), 10 μ M U46619 (B), and 3 μ g/ml Collagen (C). The representative aggregation traces were obtained from three independent experiments. Data represent the mean \pm SD (n = 3).

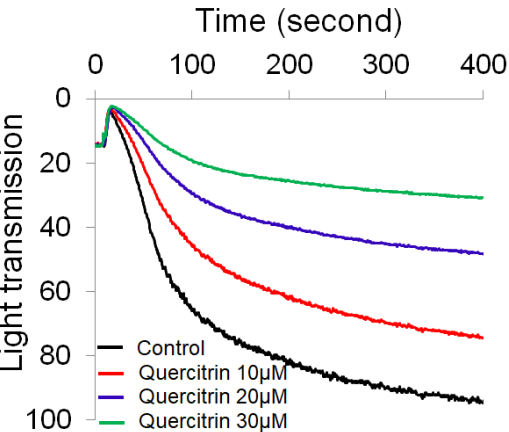
Figure S2

Mouse platelets

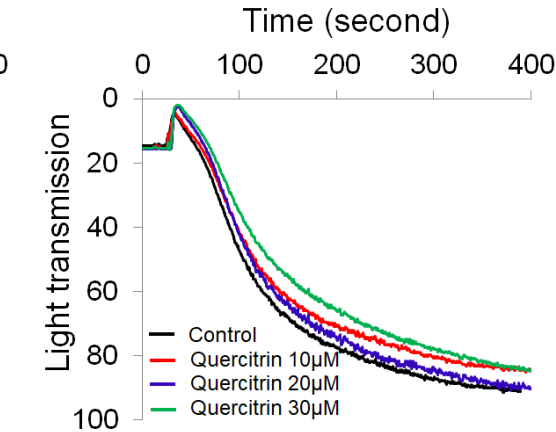
A) CRP 0.1 $\mu\text{g/ml}$



B) U46619 3 μM



C) Thrombin 0.025 U/ml



D) ADP 2.5 μM

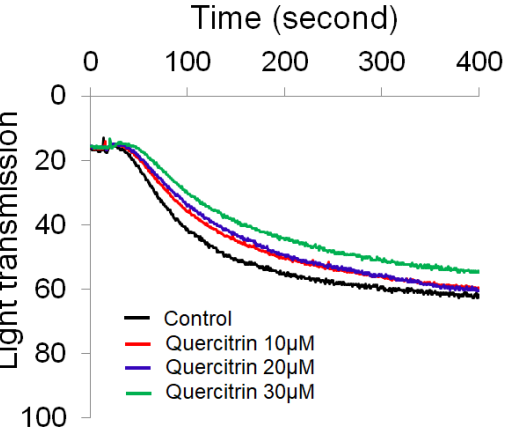


Figure S2. Role of quercitrin in mouse platelet aggregation induced by numerous agonists. Washed mouse platelets were preincubated with various concentrations of quercitrin (10, 20, and 30 μ M) for 5 minutes at 37°C and then stimulated with 0.1 μ g/ml CRP (A), 3 μ M U46619 (B), 0.025 μ g/ml Collagen (C), and 2.5 μ M ADP (D). The representative aggregation traces were obtained from three independent experiments. Data represent the mean \pm SD (n = 3).

Figure S3

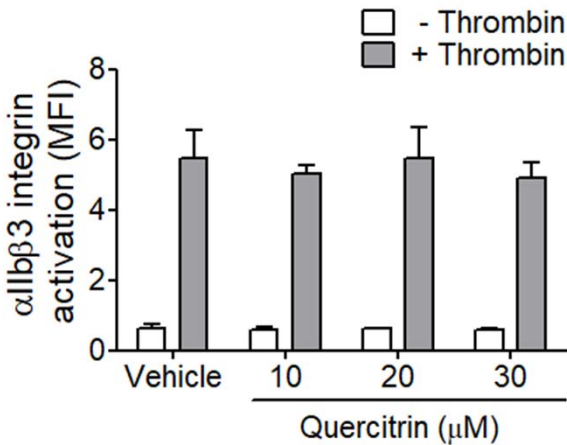
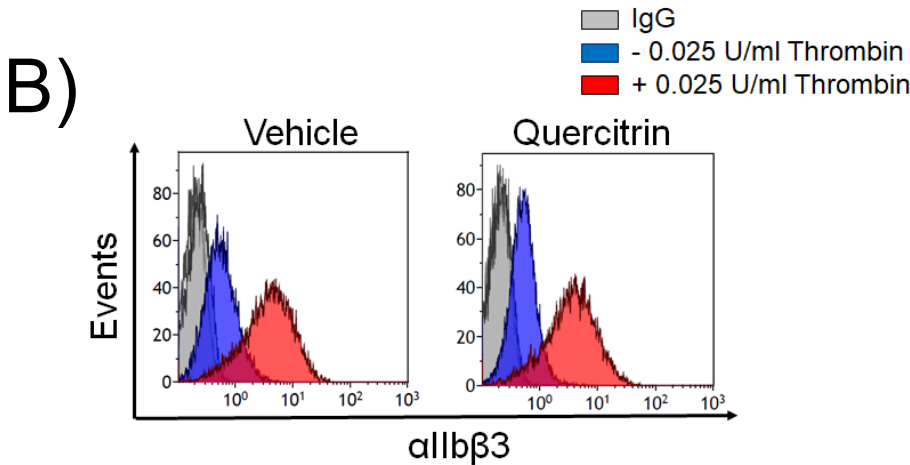
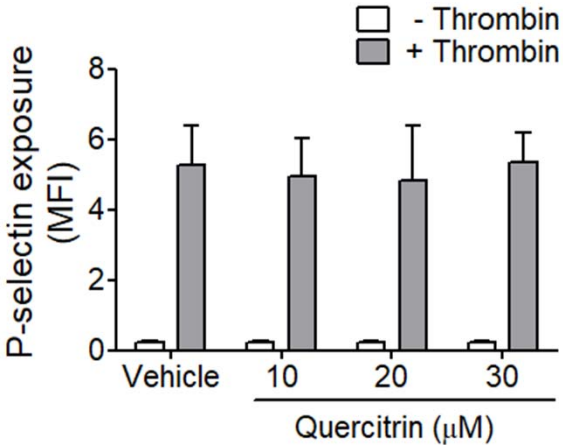
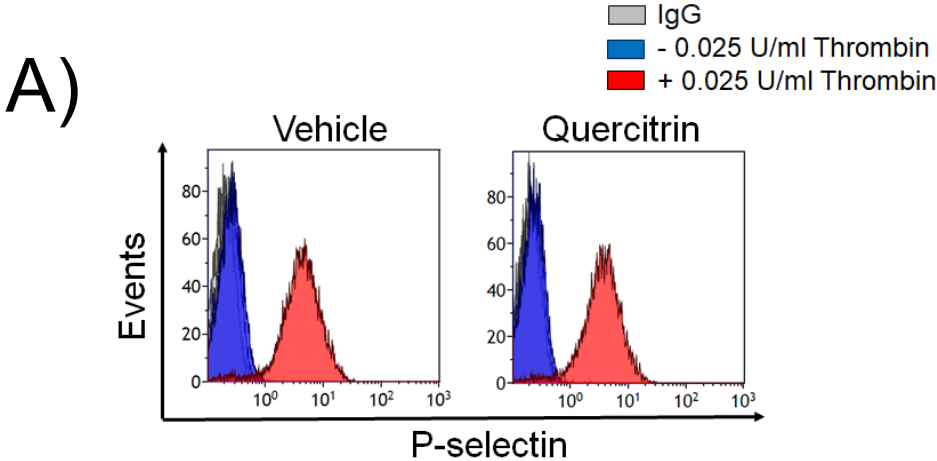


Figure S3. α IIb β 3 integrin activation and P-selectin exposure in thrombin-activated quercitrin treated platelets. Quercitrin treated platelets were activated with 0.025 U/ml thrombin, and α IIb β 3 integrin activation (A) and P-selectin exposure (B) was measured by flow cytometry. Binding of anti-activated α IIb β 3 (JON/A) and anti-P-selectin antibodies to platelets was calculated by the ratio of the geometric mean fluorescence intensity (MFI) value of antibodies to that of control IgG. Data represent mean \pm SD (n = 3).

Figure S4

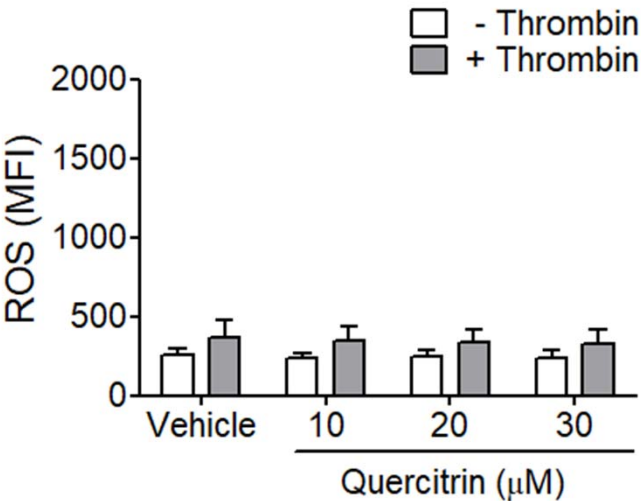
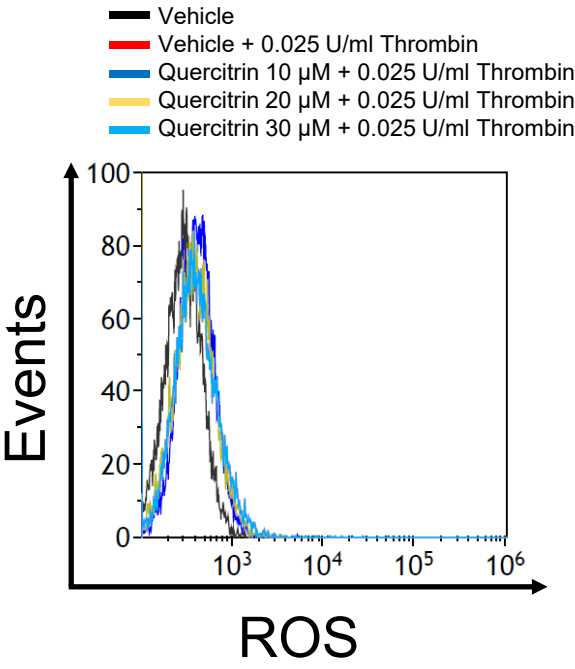


Figure S4. Intracellular H_2O_2 following thrombin stimulation in quercitrin treated platelets. Vehicle control (0.02% DMSO) and quercitrin (10, 20, and 30 μM) were incubated with H_2DCFDA and stimulated with 0.025 U/ml thrombin. DCF fluorescence was measured using flow cytometry and is quantified as mean \pm SD (n = 3).