

Figure S6. Related to Fig. 7. Influence of serum on the binding affinities of selected αllbβ3 inhibitors. (A) Binding affinity of αllbβ3 WT expressed on transiently transfected Expi293F integrin α5 and αV knock-out cells for FITC-Echistatin determined by binding of 5 nM Alexa647-MBC319.4 Fab in L15 medium supplemented with varying concentrations of serum. Binding curves were fitted to Eq.3 for the binding affinity of FITC-Echistatin at different serum concentrations. Errors were standard fitting errors from nonlinear least square fits. (B-C) Competitive binding to measure the affinity of αllbβ3 inhibitors. αllbβ3 inhibitors at the indicated concentrations competed binding of FITC-Echistatin at the concentration shown on the y axis to αllbβ3 WT expressed on transiently transfected Expi293 α5 &αV KO cells. Mean fluorescence intensity (MFI) was determined by flow cytometry without washing.