Enhanced anti-metastatic bioactivity of an IGF-Trap re-engineered to improve physicochemical properties

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Running Title: Distinct pharmacodynamic properties of bio-engineered IGF-TRAPs

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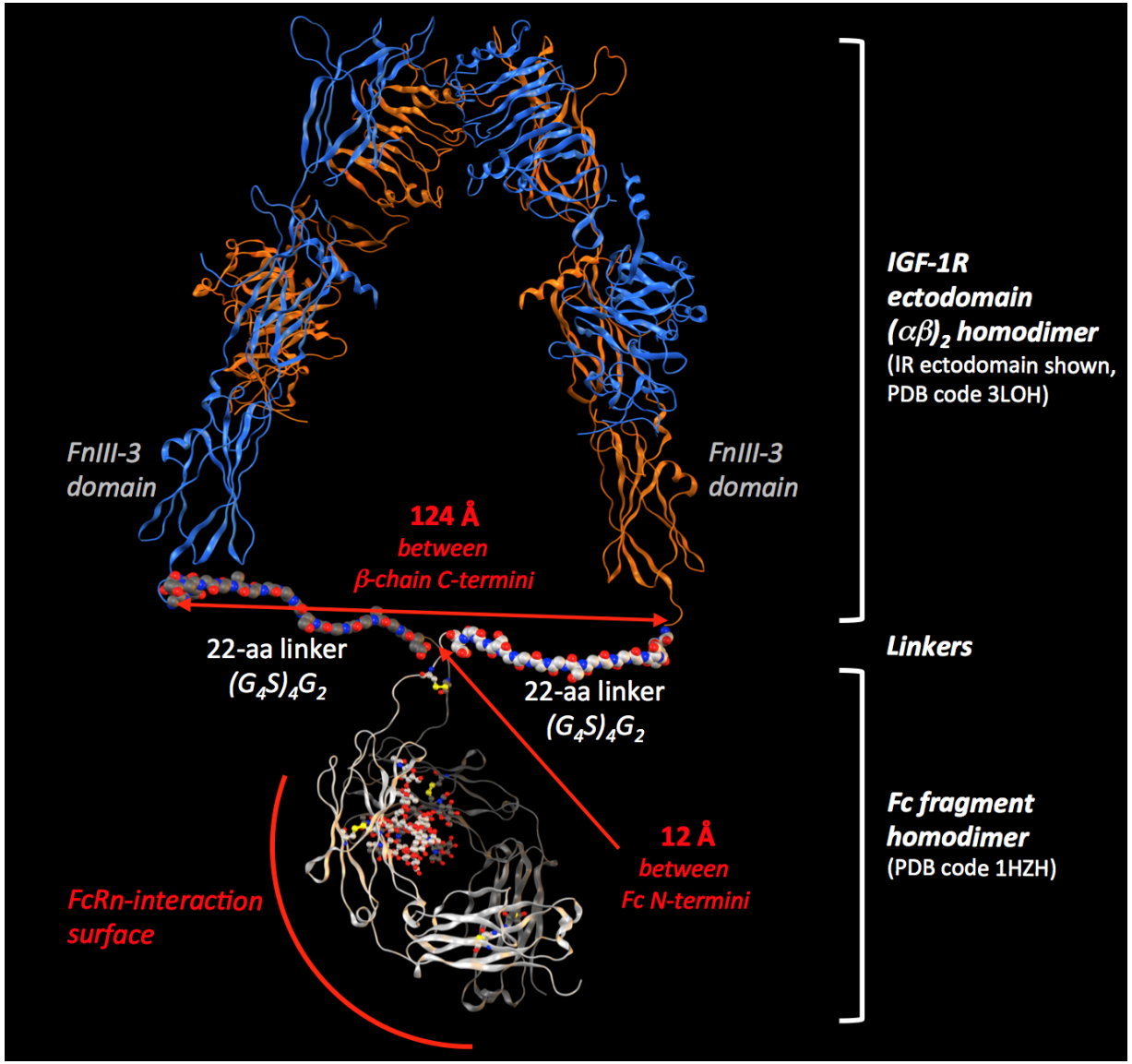
Supplementary Figure legends

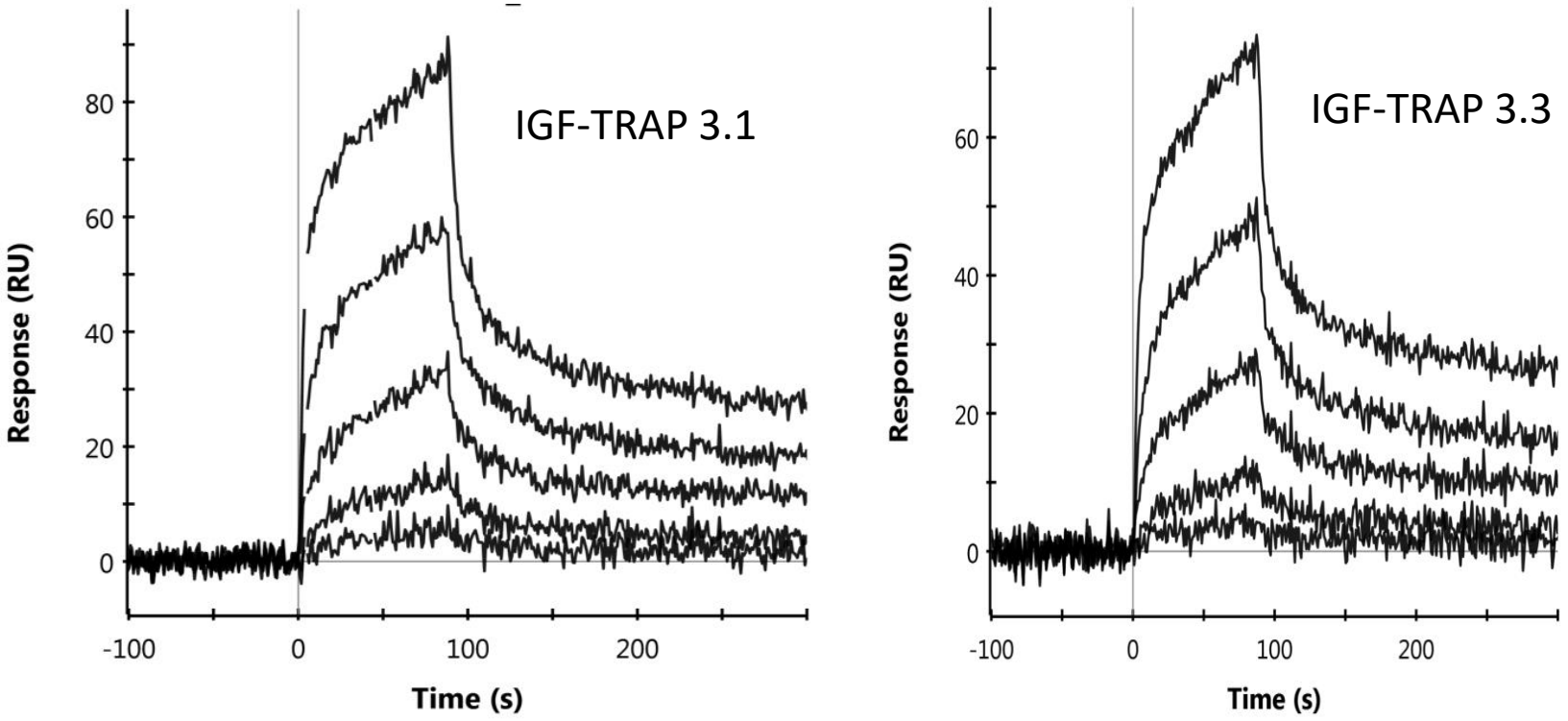
Figure S1. A proposed 3D model of the IGF-TRAP. The extracellular domain of IGF-1R and the fused Fc domain of human IgG_1 are based on the known crystal structures of the ectodomain of hIR and the Fc domain of a $hIgG_1$, respectively.

Figure S2. IGF-TRAPs 3.1 and 3.3 bind to the FcRn with similar affinities. Shown are SPR sensorgrams of IGF-TRAP 3.1 and IGF-TRAP 3.3 flowing over 80 RUs of C-terminally biotinylated FcRn on a neutravidin sensor surface. The apparent K_D's based on an equilibrium fit gave similar results for each of the two IGF-TRAPs.

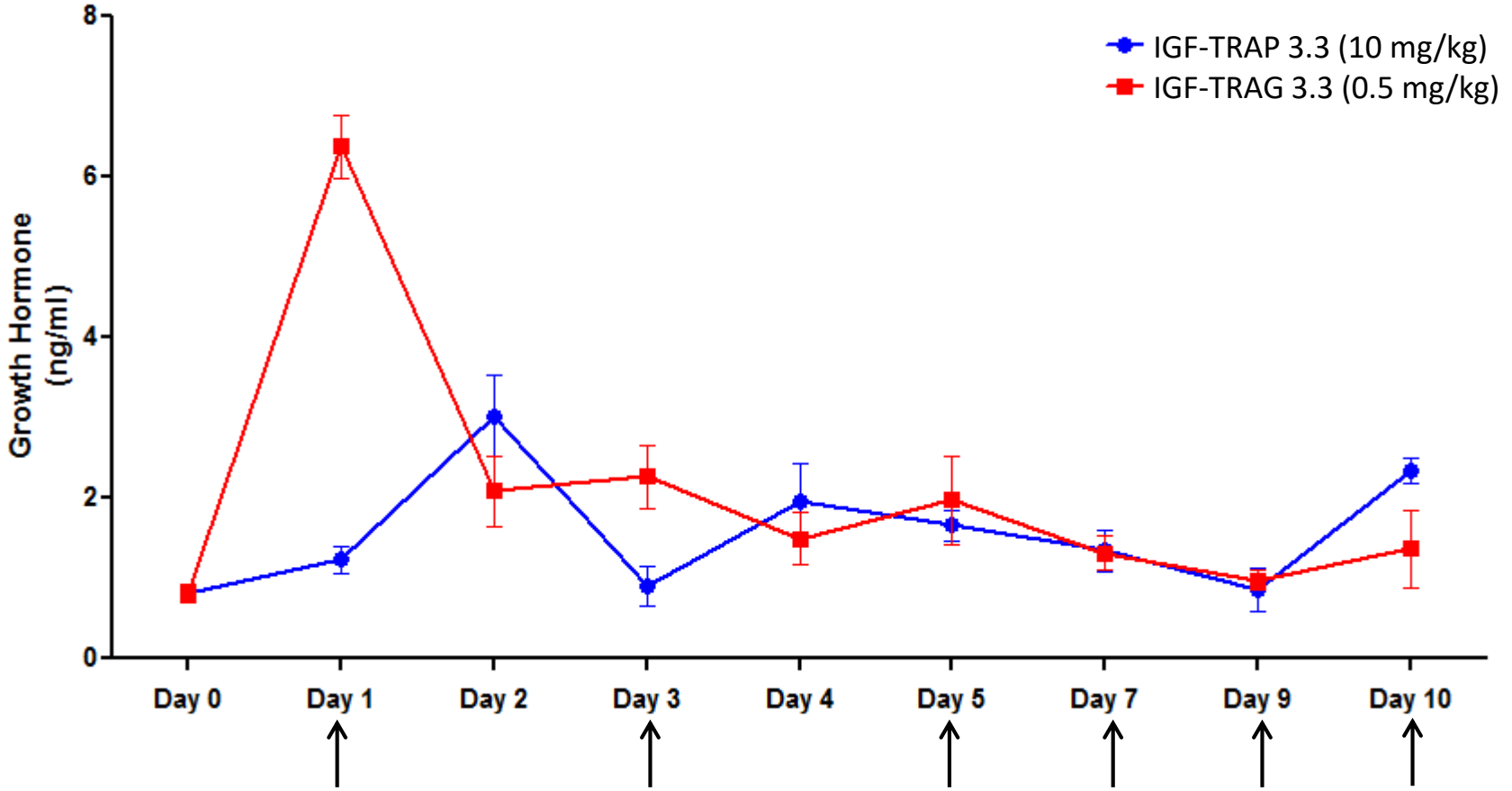
Figure S3. IGF-TRAP 3.3 injections alter circulating growth hormone (GH) levels. Blood was collected daily from mice injected with 0.5 or 10 mg/kg IGF-TRAP 3.3 on alternate days, as indicated. Shown are mean circulating GH levels (±SD) based on 3 animals per time point as measured by ELISA.



Supplementary Fig S1



Supplementary Fig S2



Supplementary Fig S3