

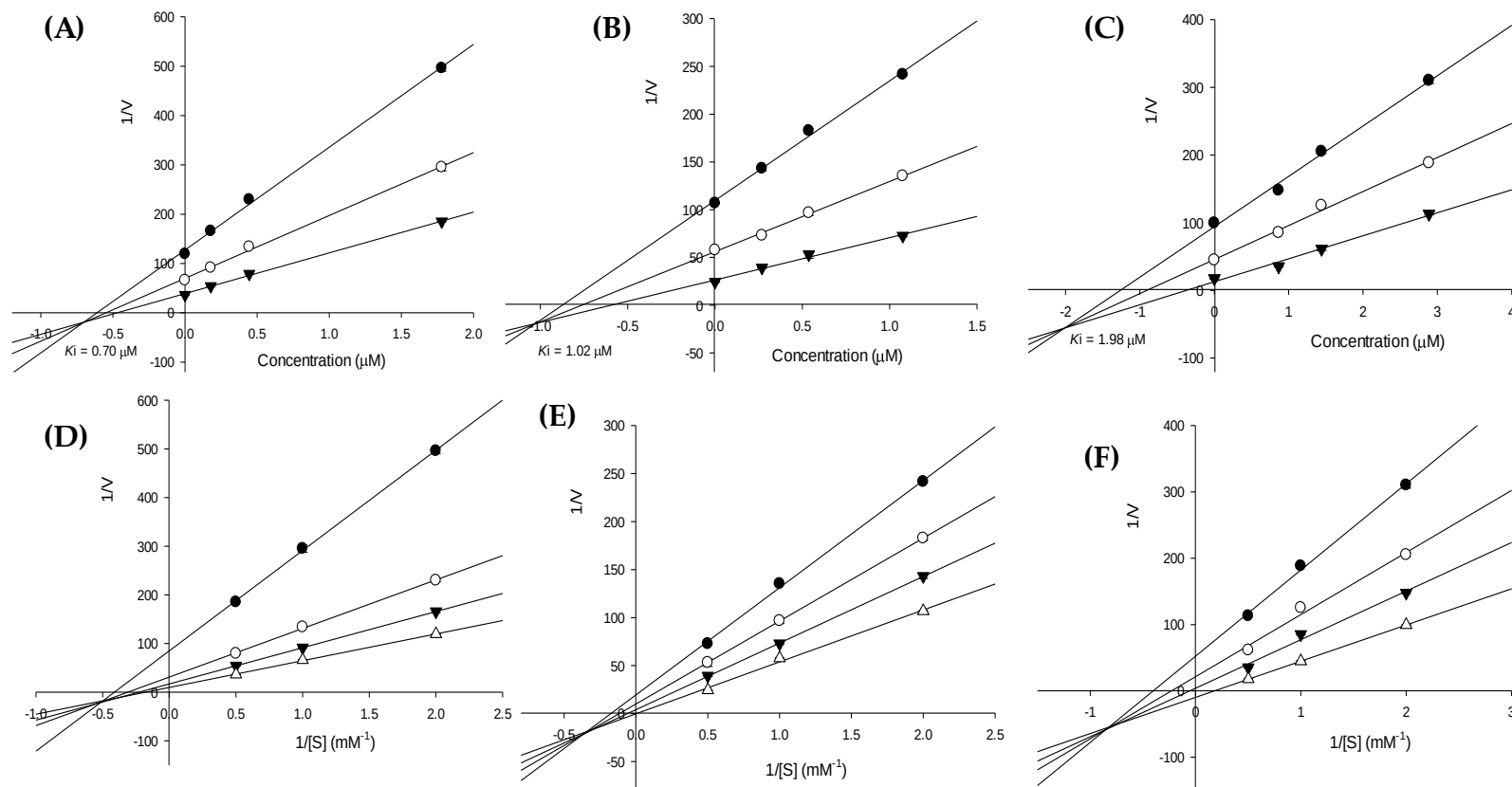


Figure S1. Enzyme kinetic plots of PTP1B inhibition by 1-3. (A-C) Dixon plots of PTP1B inhibition. 1 (A), 2 (B) and 3 (C) were tested in the presence of different substrate concentrations: 0.5 mM (●), 1.0 mM (○) and 2.0 mM (▼). (D-F) Lineweaver-Burk plots of PTP1B inhibition by 1-3. PTP1B inhibition was analyzed in the presence of different concentrations of test samples as follows: 1.78 μM (●), 0.44 μM (○), 0.18 μM (▼), 0 μM (△) for 1; 1.07 μM (●), 0.54 μM (○), 0.27 μM (▼), 0 μM (△) for 2; 2.89 μM (●), 1.44 μM (○), 0.87 μM (▼), 0 μM (△) for 3.

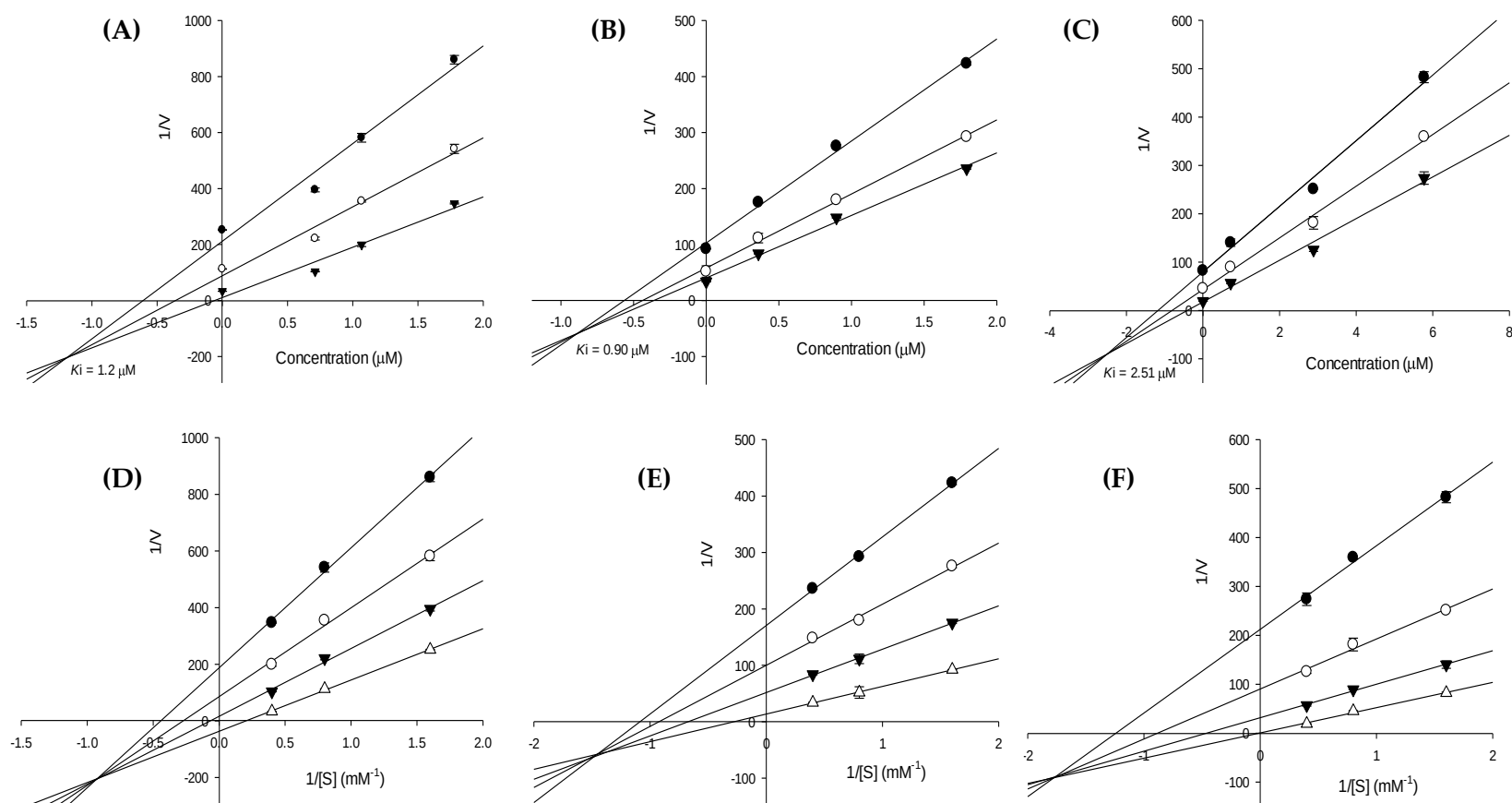


Figure S2. Enzyme kinetic plots of α -glucosidase inhibition by 1-3. (A-C) Dixon plots of α -glucosidase inhibition. 1 (A), 2 (B) and 3 (C) were tested in the presence of different substrate concentrations: 0.4 mM ($\bullet$), 0.8 mM (O) and 1.6 mM ($\blacktriangledown$). (D-F) Lineweaver-Burk plots of α -glucosidase inhibition by 1-3. α -Glucosidase inhibition was analyzed in the presence of different concentrations of test samples as follows: 1.78 μM ($\bullet$), 1.07 μM (O), 0.71 μM ($\blacktriangledown$), 0 μM ($\triangle$) for 1; 1.79 μM ($\bullet$), 0.9 μM (O), 0.36 μM ($\blacktriangledown$), 0 μM ($\triangle$) for 2; 5.77 μM ($\bullet$), 2.88 μM (O), 0.72 μM ($\blacktriangledown$), 0 μM ($\triangle$) for 3.