

1/6

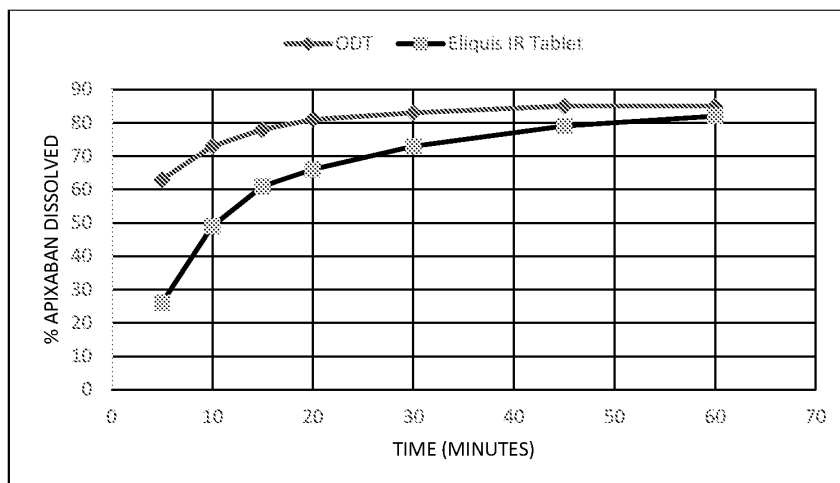


Figure 1: Comparative dissolution profile of apixaban (5 mg) orally disintegrating tablet and film-coated immediate release tablets ELIQUIS® of at pH 6.8

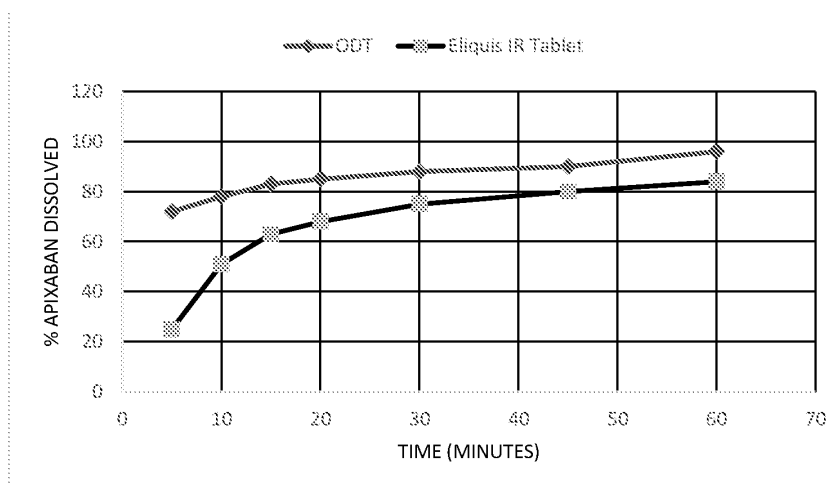


Figure 2: Comparative dissolution profile of apixaban (5 mg) orally disintegrating tablet and film-coated immediate release tablets ELIQUIS® of at pH 4.5

2/6

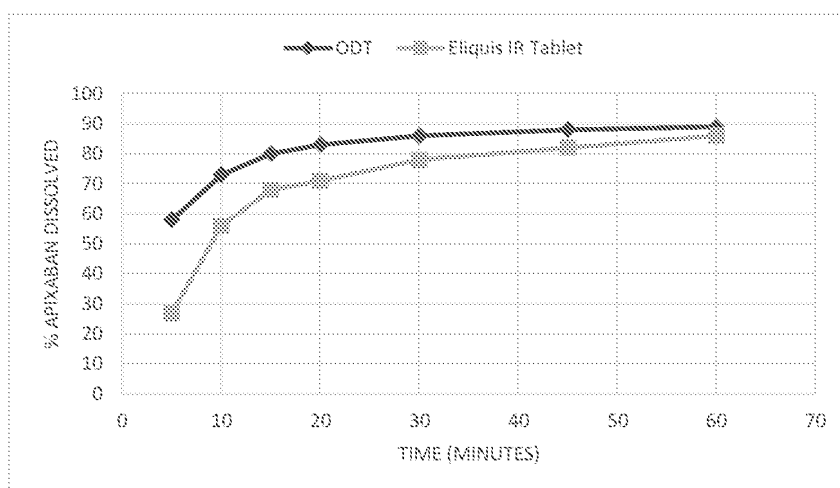


Figure 3: Comparative dissolution profile of apixaban (5 mg) orally disintegrating tablet and film-coated immediate release tablets ELIQUIS® of at pH 1.2

3/6

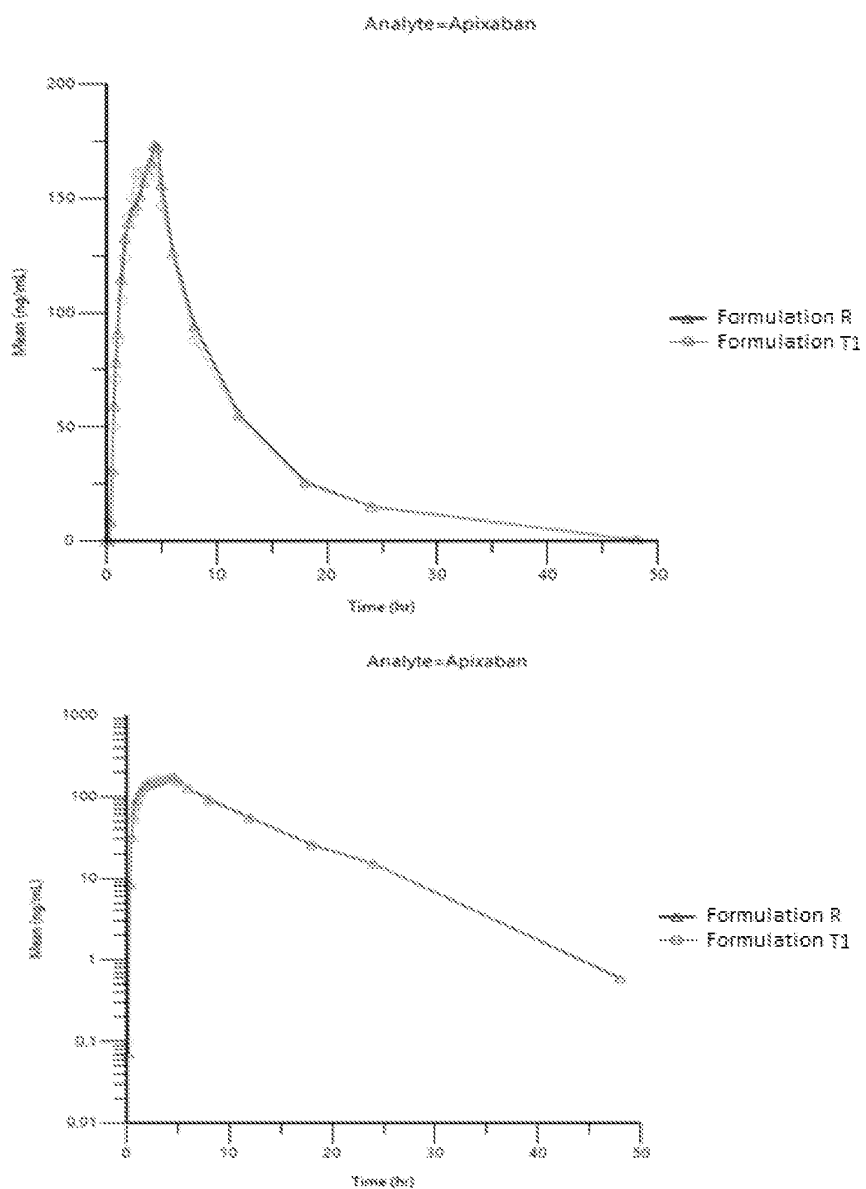


Figure 4: Linear and semi-logarithmic plots of mean plasma concentration (MPC) versus time (t) for Apixaban orally disintegrating tablet (T1: Without water) compared to ELIQUIS®

4/6

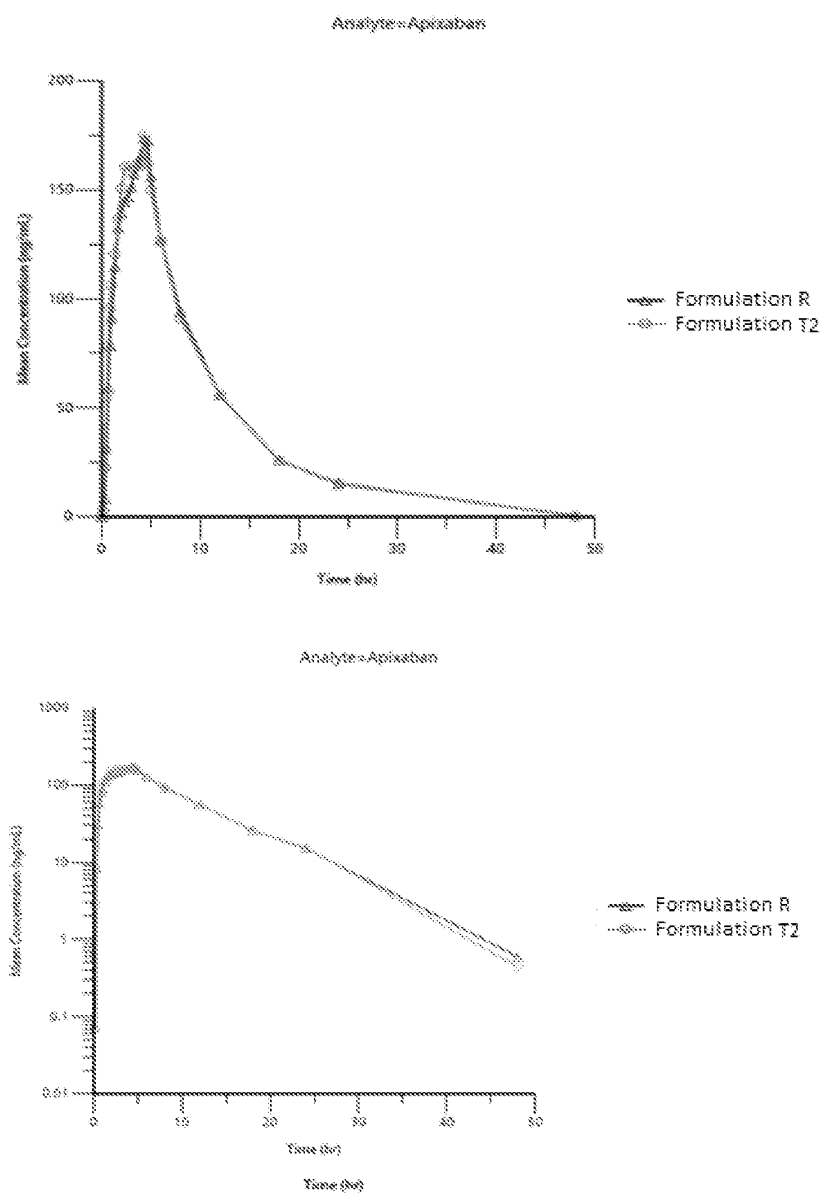


Figure 5: Linear and semi-logarithmic plots of mean plasma concentration (MPC) versus time (t) for Apixaban orally disintegrating tablet (T2: With water) compared to ELIQUIS[®]

5/6

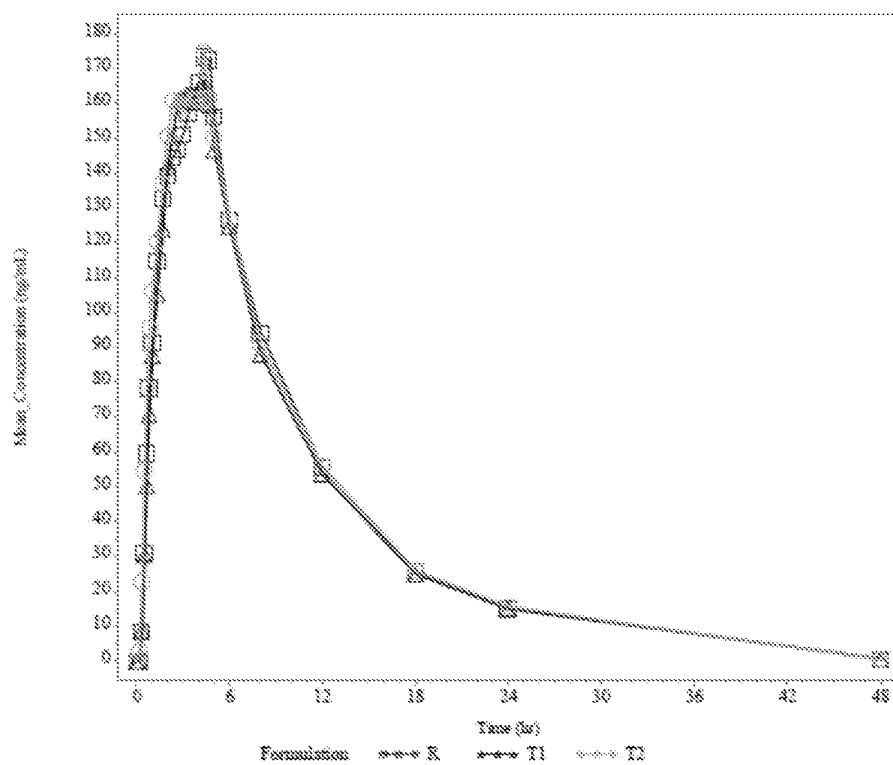


Figure 6: Comparative linear plot of mean plasma concentration (MPC) versus time (t) for Apixaban orally disintegrating tablet (T1: Without water; T2: With water & R: ELIQUIS®)

6/6

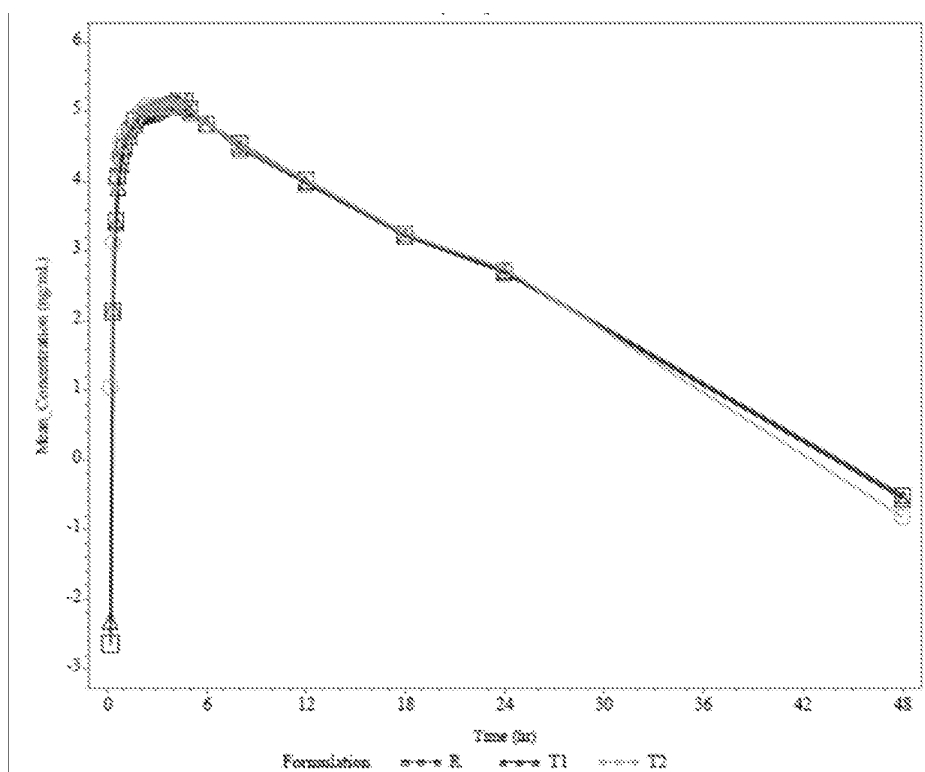


Figure 7: Comparative semi-logarithmic plot of mean plasma concentration (MPC) versus time (t) for Apixaban orally disintegrating tablet (T1: Without water; T2: With water & R: ELIQUIS®)