Objective:

1. The main objective of the research work to develop stability indicating RP-HPLC method for determination of impurities present in the Olmesartan Medoximil, and Hydrochlorothiazide Tablet dosage form.

2. To develop stability indicating RP-HPLC method for determination of impurities present in the Emtricitabine and Tenofovir Disoproxil Fumarate Tablet dosage form.

3. To develop a single RP-HPLC method for estimation of impurities and degradation products of drug from the combination of drug product.

4. The developed RP-HPLC method for estimation of impurities was validated as per current ICH guideline.