Ans 1) BA & BE

BA/BE Studies are needed by regulations to guarantee remedial proportionality b/w a pharmaceutically comparable test item and a reference item.

:- BA/BE studies are finished early and late clinical trial definitions, formulations utilized as a part of clinical trial and steadiness Studies.

:- Everybody has more heaped on their plate than any time in recent remembrance, and numerous consultant discover themselves always reorganizing their work ~~exp~~ exercises.

:- BA (Bioavailability) is defined as the rate and extent to the which the active ingredient or active moiety is absorbed from a drug product and becomes available at the site of action.

:- BE (Bioequivalence) is defined as the absence of a significant difference in the rate and extent to which the active ingredient or active moiety in pharmaceutical equivalents or pharmaceutical alternatives becomes available at the site of drug action when administered at the same molar dose under similar conditions in an appropriately designed study.