

## Pharmacokinetics III summary

| Topic                            | Details  |
|----------------------------------|--|
| Pharmacokinetics                 | Absorption, Distribution, Metabolism, Excretion  |
| Biotransformation                | Unionized drugs to ionized, water-soluble metabolites for easier excretion   |
| Organs Involved                  | Mainly liver, but also lungs, kidneys, intestines  |
| Drug Metabolism<br>Outcomes      | <ol style="list-style-type: none"> <li>1. Active drug → Inactive metabolite</li> <li>2. Active drug → Active metabolite (codeine → morphine)</li> <li>3. Prodrug → Active drug (enalapril → enalaprilat)</li> <li>4. Toxic metabolite formation</li> </ol> |
| Phases of Metabolism             | <p>Phase I: Oxidation, reduction, hydrolysis (CYP450)</p> <p>Phase II: Conjugation (glucuronic acid, sulfate, etc.)</p>  |
| Factors Affecting<br>Metabolism  | Age, Sex, Liver disease, Genetic variations (slow/fast acetylators), Enzyme induction/inhibition   |
| Enzyme Induction<br>& Inhibition | <p>Inducing drugs:</p> <p>Phenobarbitone ,phenytoin ,nicotine , rifampicin</p> <p>Inhibiting drugs:</p> <p>Erythromycin, clarithromycin ,cimetidine ,contraceptives</p>  |

Done by: Dr. Farah Breik