

# Drug absorption

Absorption is the transfer of the drug into the bloodstream. In IV this process is complete thus the whole dose reaches the systemic circulation. However in other routes of administration the rate and efficiency is dependent on many factors and usually only results in partial absorption.

## Transport in GIT

Due to their chemical properties, drugs are absorbed either by passive diffusion or active transport.

1. **Passive diffusion:** Passive diffusion is facilitated by the concentration gradient across the membrane. The drug moves from high concentration to low concentration, it does not require an extra carrier or extra energy. Water-soluble agents use aqueous pores in the membrane while lipid soluble diffuse directly through the membrane.
2. **Active transport:** Active transport requires specific carriers and ATP. The specific carrier protein carries the drugs that closely resemble the structure of the naturally occurring metabolites specific for the carrier. It is capable of transporting from low to high concentration compartments.
3. **Endocytosis:** Endocytosis means that the membrane engulfs the drug molecule and transports it into the cell by pinching off the drug-filled vesicle. E.g. Vitamin B<sub>12</sub> is transported this way.

## Factors affecting absorption

- **pH**, most drugs are either weak acids or weak bases. Drugs pass through membranes easier if uncharged. The effective concentration of each drug at the absorption site is determined by the relative concentrations of the charged and uncharged forms.
- **Blood Flow** to the intestine is much more extensive than to the stomach as well as larger surface area favours the intestine for absorption.
- **Movement** movement of the drug through the GIT has a large effect on its absorption. The faster it moves through thus decreasing its contact time at the absorption surface and therefore less is absorbed. This is the case in severe diarrhea. Also the presence of food dilutes the drug and slows gastric emptying and thus the drug is generally absorbed slower.

## Links

### Related articles

- Route of drug administration
- Bioavailability

### Bibliography

- FINKEL, CLARK, CUBEDDU, HARVEY, CHAMPE, et al. *Lippincott's Illustrated Reviews: Pharmacology*. 4. edition. 2009. ISBN 978-1605472003.