

# **SNS COLLEGE OF PHARMACY AND HEALTH SCIENCES**

Affiliated To The Tamil Nadu Dr. MGR Medical University, Chennai  
Approved by Pharmacy Council of India, New Delhi. Coimbatore -641035

**COURSE NAME : PHARMACOLOGY(ER20-21 T)**

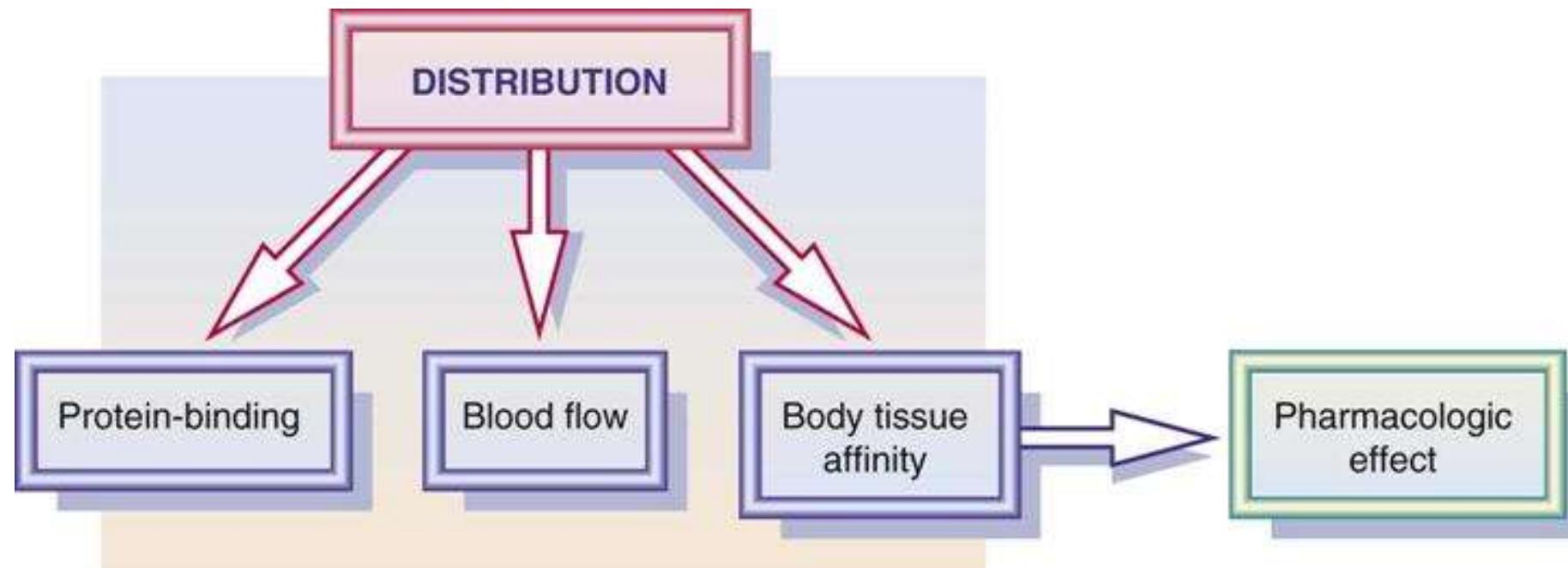
**YEAR : DPHARM-II YEAR**

**TOPIC 4 : PHARMACOKINETICS (DISTRIBUTION)**

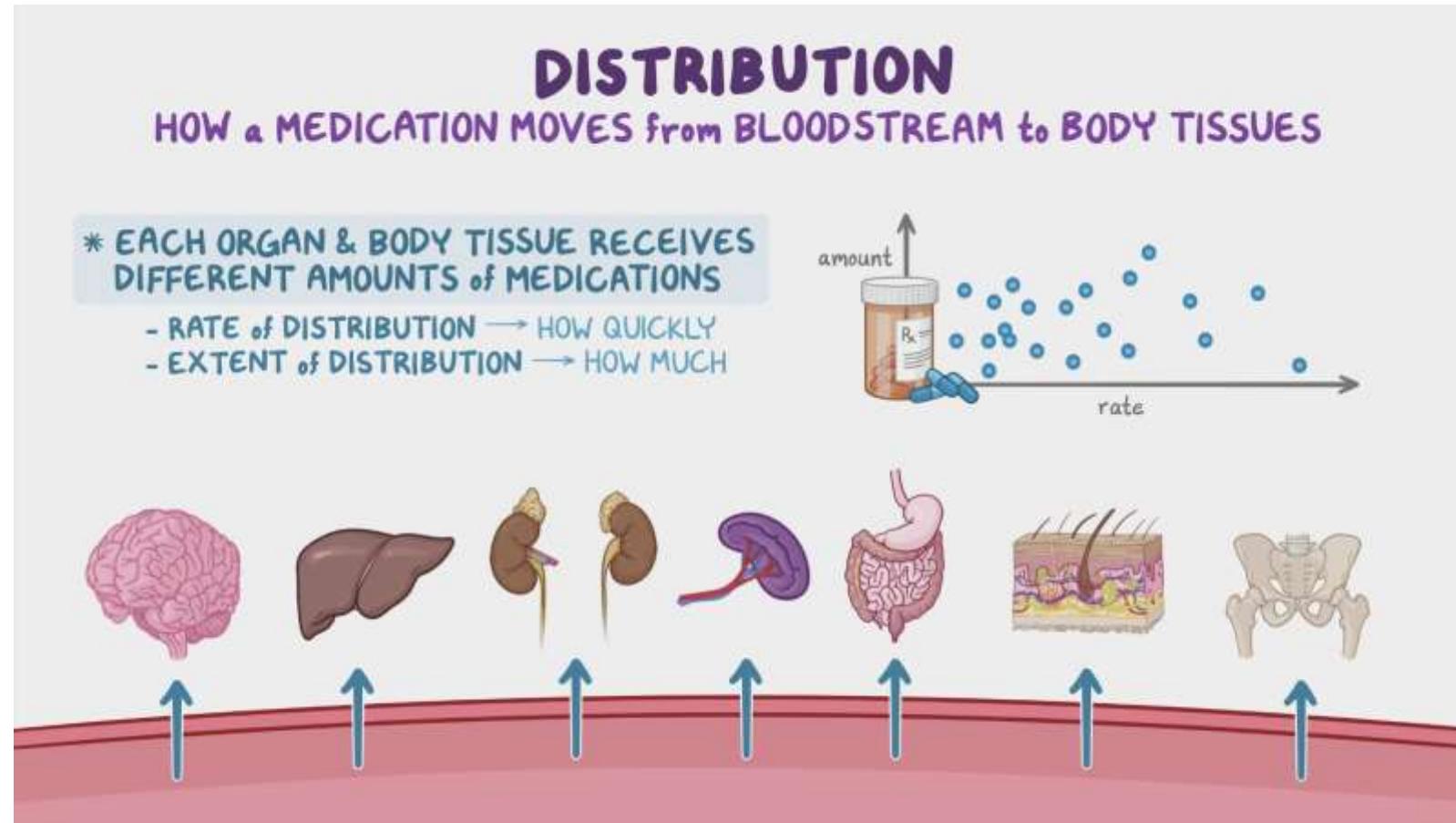
# DESIGN THINKING IN DRUG DISTRIBUTION

- **Empathize:** Deeply understand the user or patient's challenges, needs, and experiences related to drug distribution in the body.
- **Define:** Reframe the problem based on insights from the empathize phase and establish clear context.
- **Ideate:** Brainstorm and explore a wide range of ideas and potential solutions, including innovative formulations or carriers that enhance selective distribution.
- **Prototype:** Simulate and build drug delivery prototypes to improve tissue-specific distribution and patient outcomes.

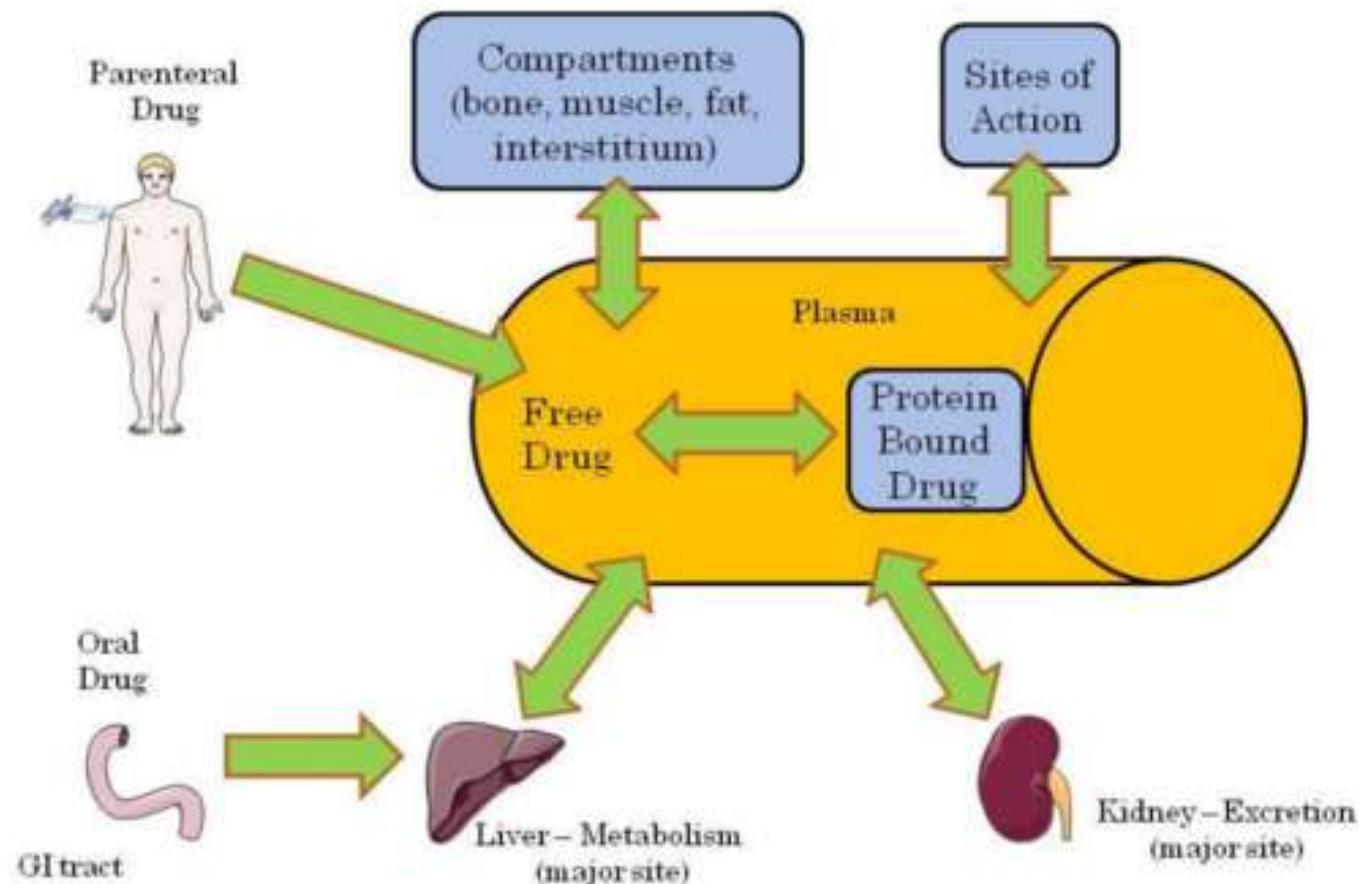
# MINDMAP



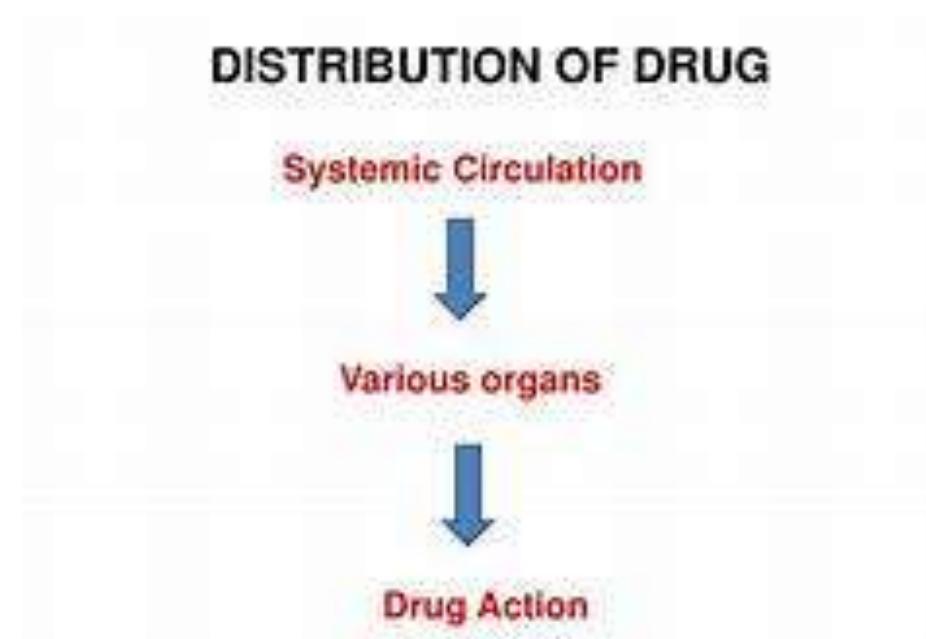
# DISTRIBUTION



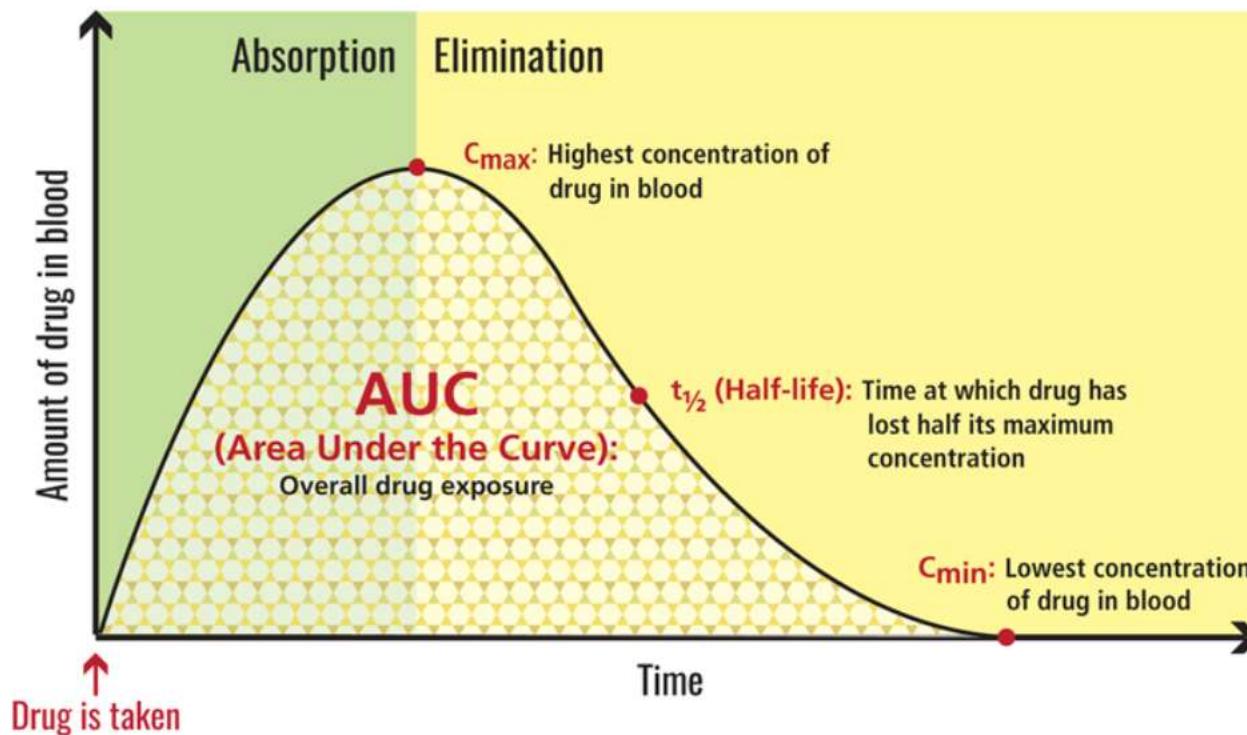
## IMPORTANCE OF DRUG DISTRIBUTION

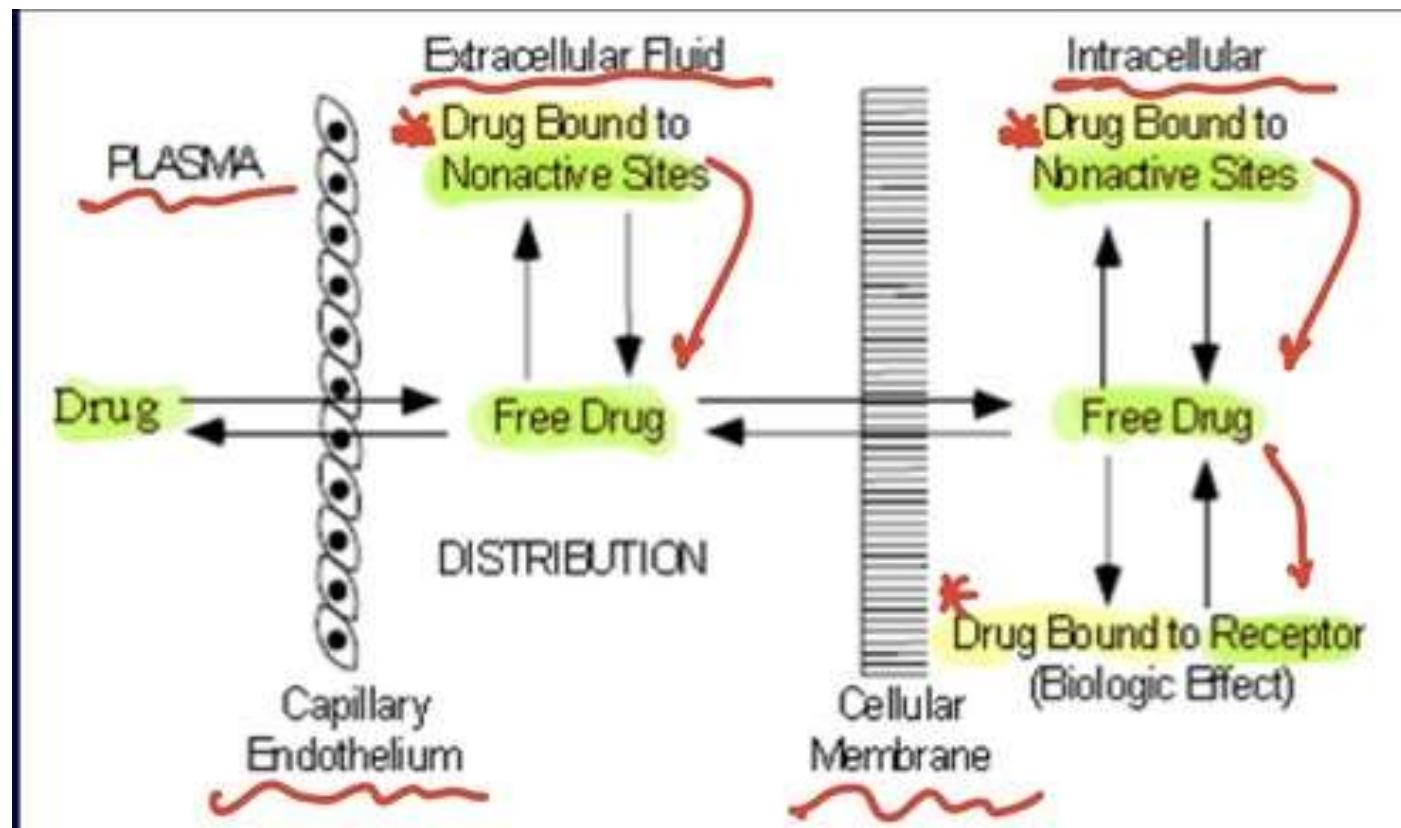


# PROCESS OF DRUG DISTRIBUTION



# TERMINOLOGY USED IN DRUG DISTRIBUTION





# VOLUME OF DISTRIBUTION

## Volume of distribution (Vd)

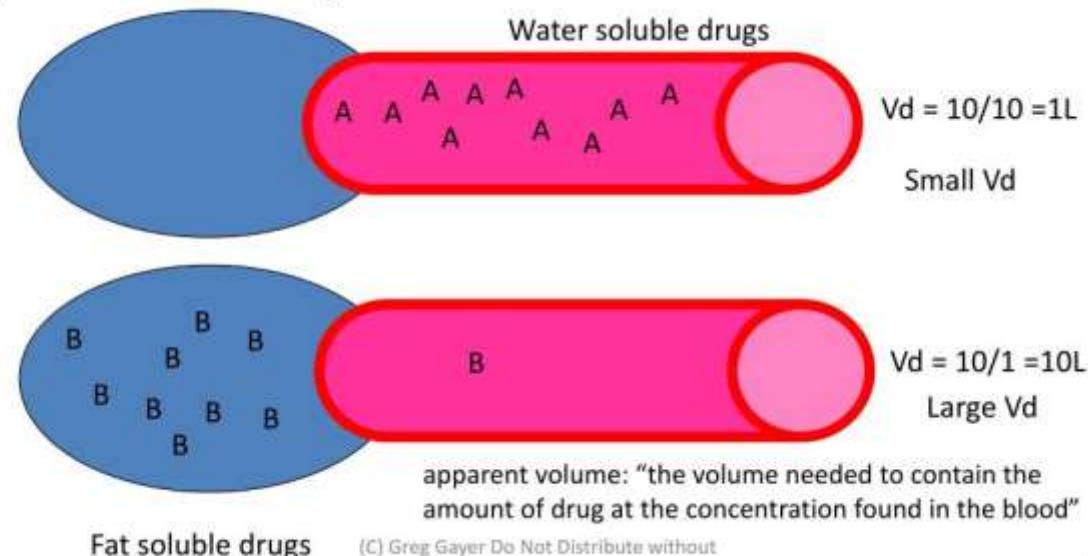
$$V_d = \frac{\text{Amount of drug in body}}{C_{\text{plasma}}}$$

(Units=volume)

Used to calculate Loading dose and  $t_{1/2}$

$$\text{Loading dose} = \frac{C_p \times V_d}{F}$$

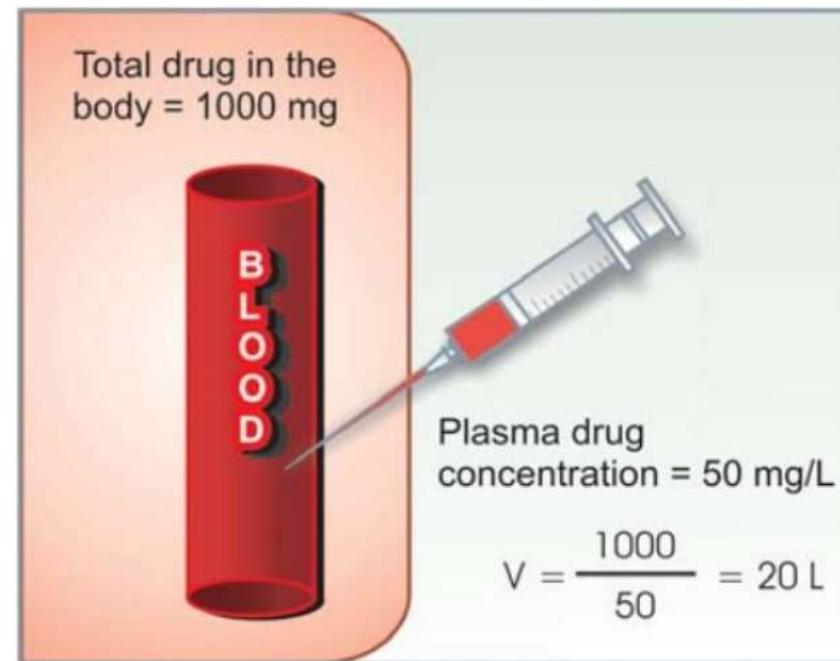
$$t_{1/2} = \frac{C_p \times V_d}{F}$$



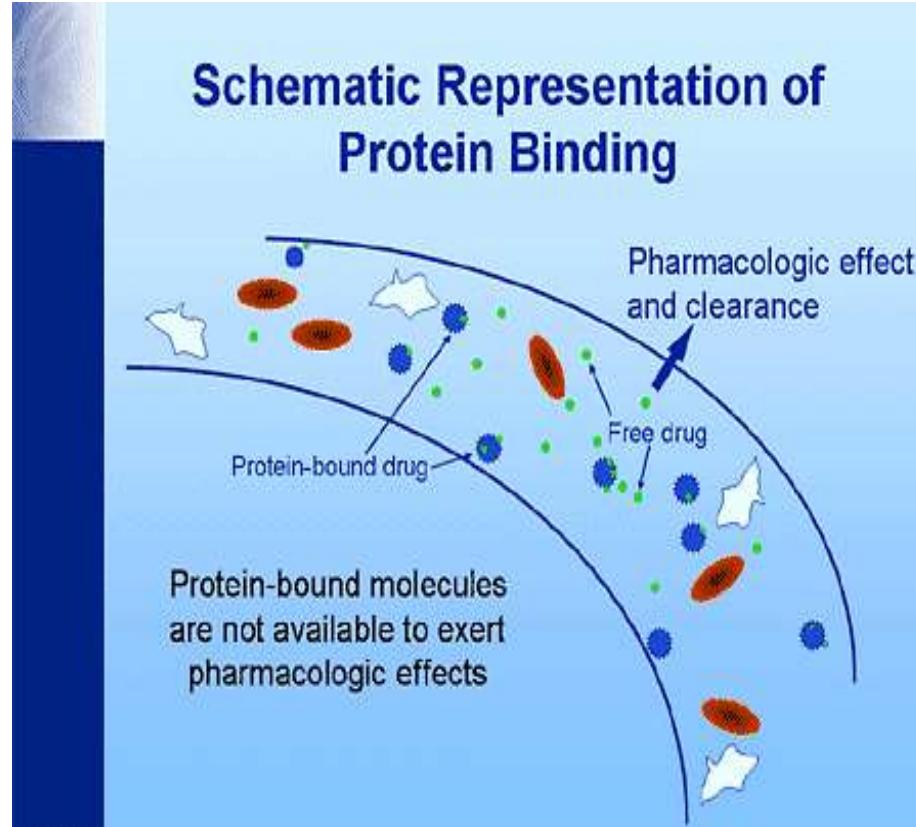
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## APPARENT VOLUME OF DISTRIBUTION



# PLASMA PROTEIN BINDING



## REDISTRIBUTION

### Redistribution of drug

Highly Lipid soluble drugs: Thiopentone sodium



Organs with High blood flow: Brain, Kidney, Liver

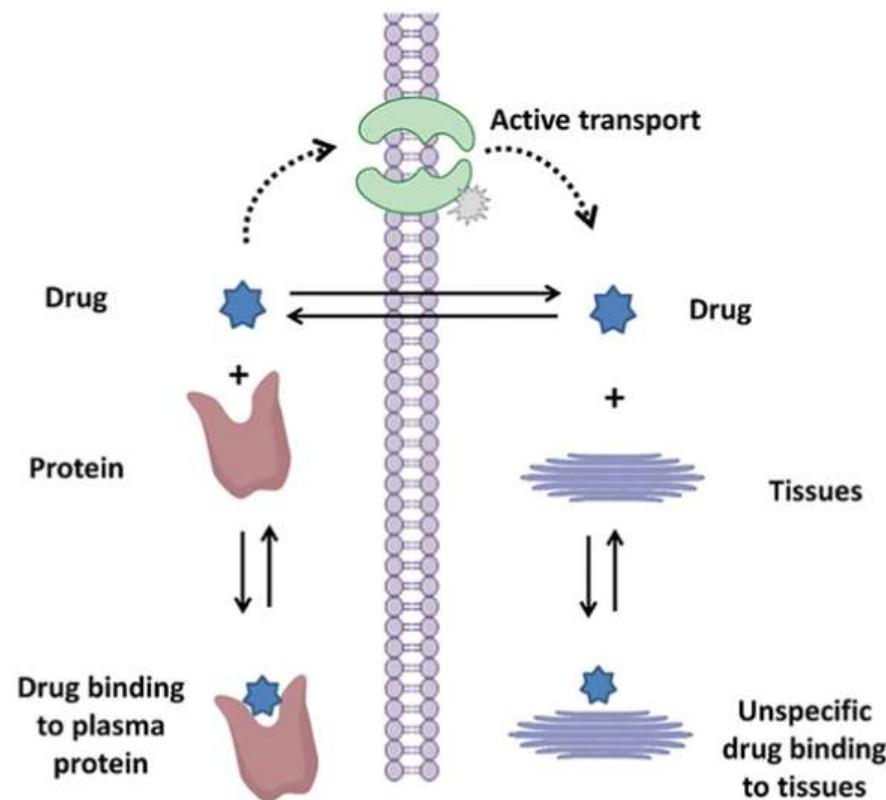


Less vascular & Bulky Tissue: Fat, Muscle

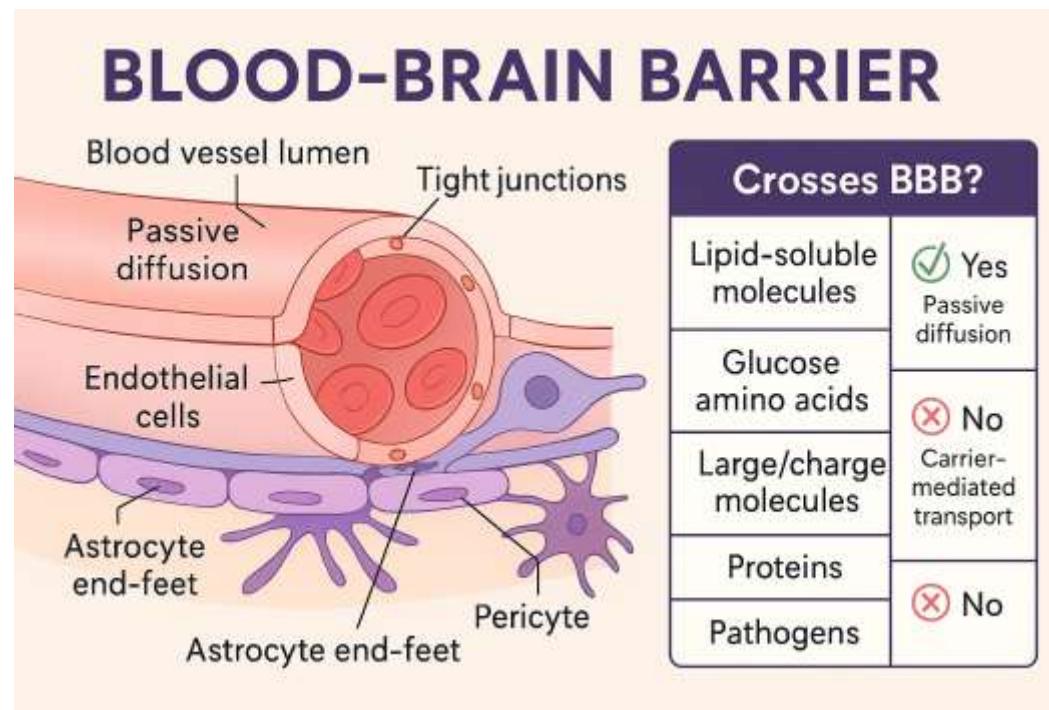


Termination of Drug Action

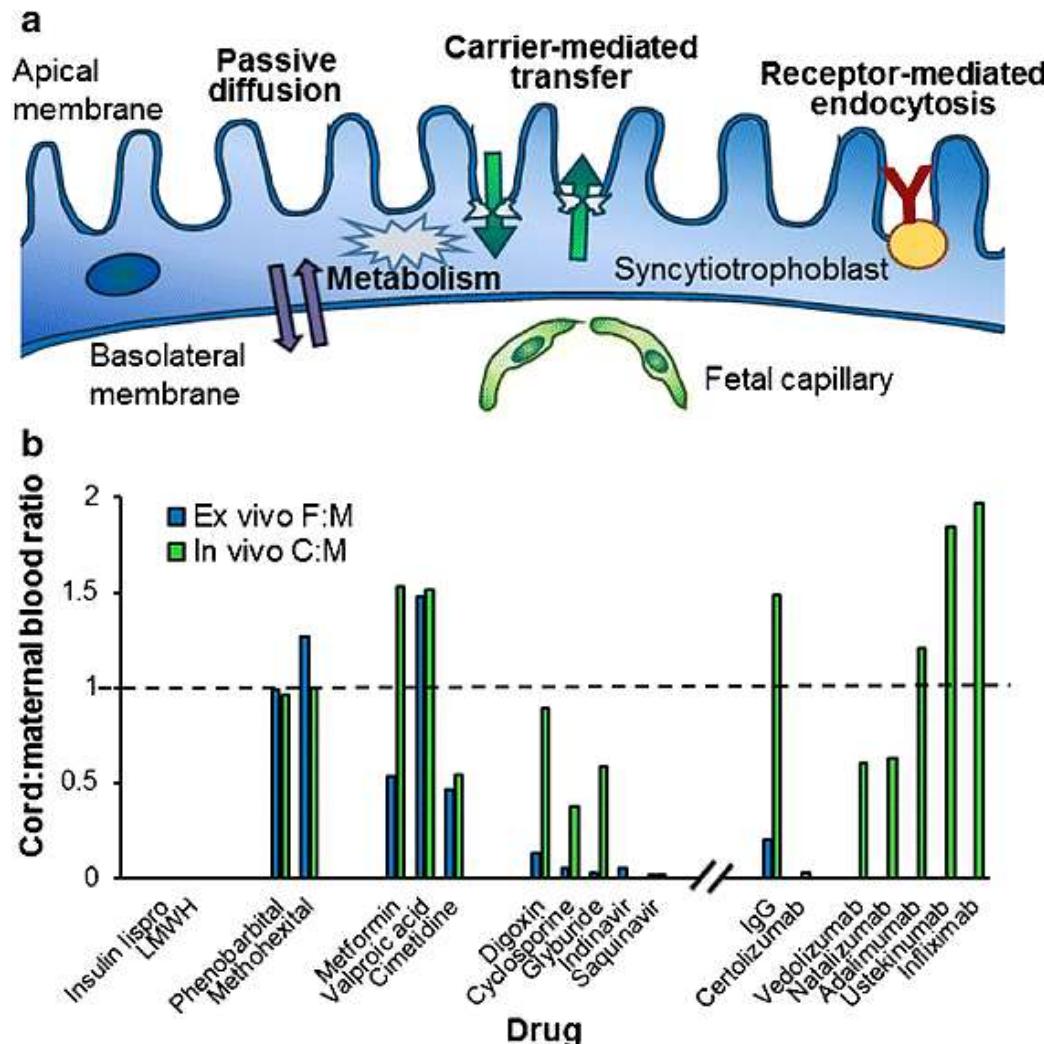
# TISSUE BINDING



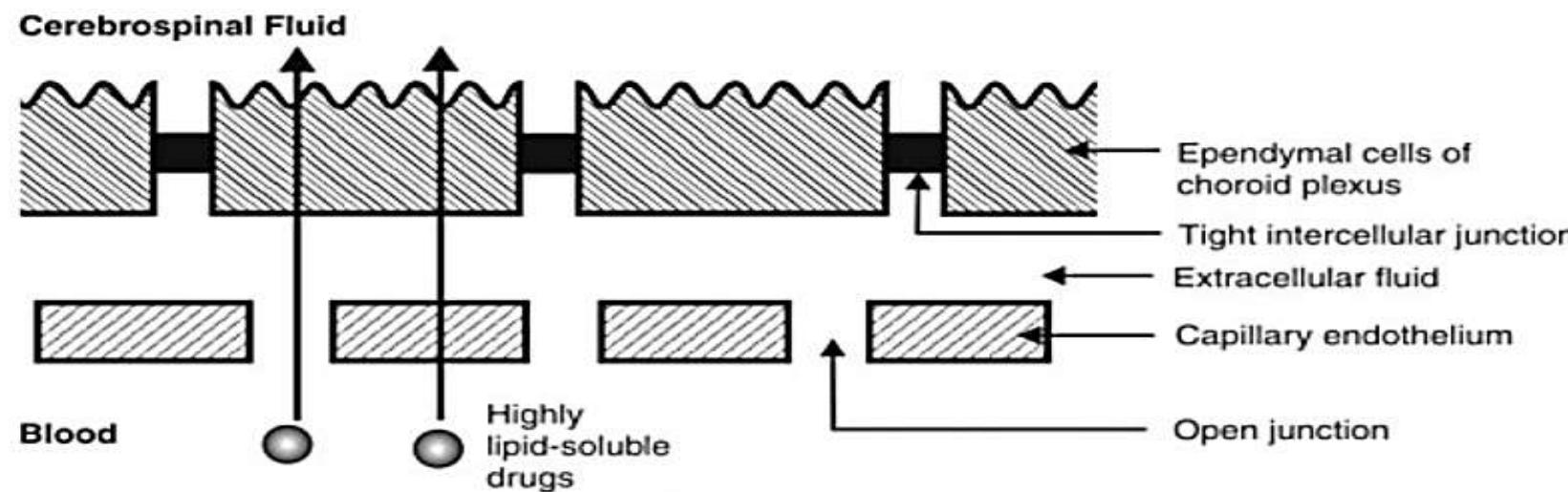
# BLOOD BRAIN BARRIER



# PLACENTAL BARRIER

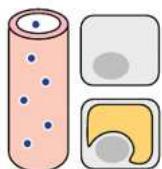
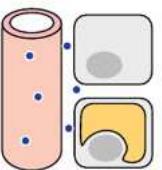
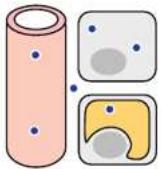


# CSF BARRIER

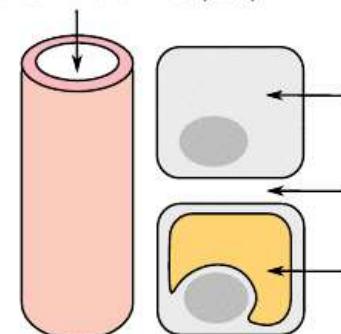


# CLASS ASSESSMENTS

How drugs distribute in the body?



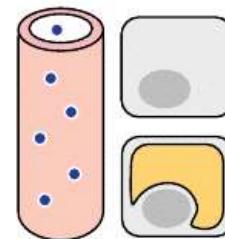
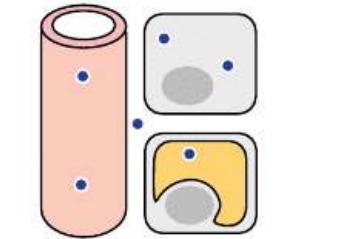
Intravascular volume (5%)



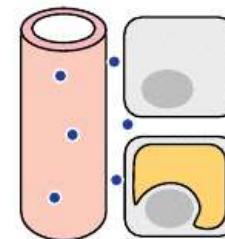
Intracellular volume (40%)

Interstitial volume (15%)

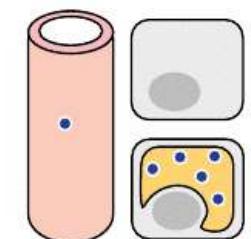
Fat (several %)



Drug confined to circulation  
(very large drug molecules)



Drug excluded by cell membranes  
(very polar drug molecules)



Drug enriched in fat  
(lipophilic drugs)

## SUMMARY

- Drug distribution is key to pharmacokinetics, influenced by solubility, binding, and barriers.
- Understanding it optimizes therapy, minimizes risks. Key concepts:  $V_d$ , protein binding, redistribution, special barriers.

## REFERENCE

- Rang & Dale's Pharmacology, 9th Edition, Elsevier, 2019.
- Goodman & Gilman's: The Pharmacological Basis of Therapeutics, 13th Edition, McGraw-Hill, 2018.
- Text book of Pharmacology by KD Tripathi.

