



**Biopharmaceutics &
Pharmacokinetics**
8th SEM

**Drug
Distribution**

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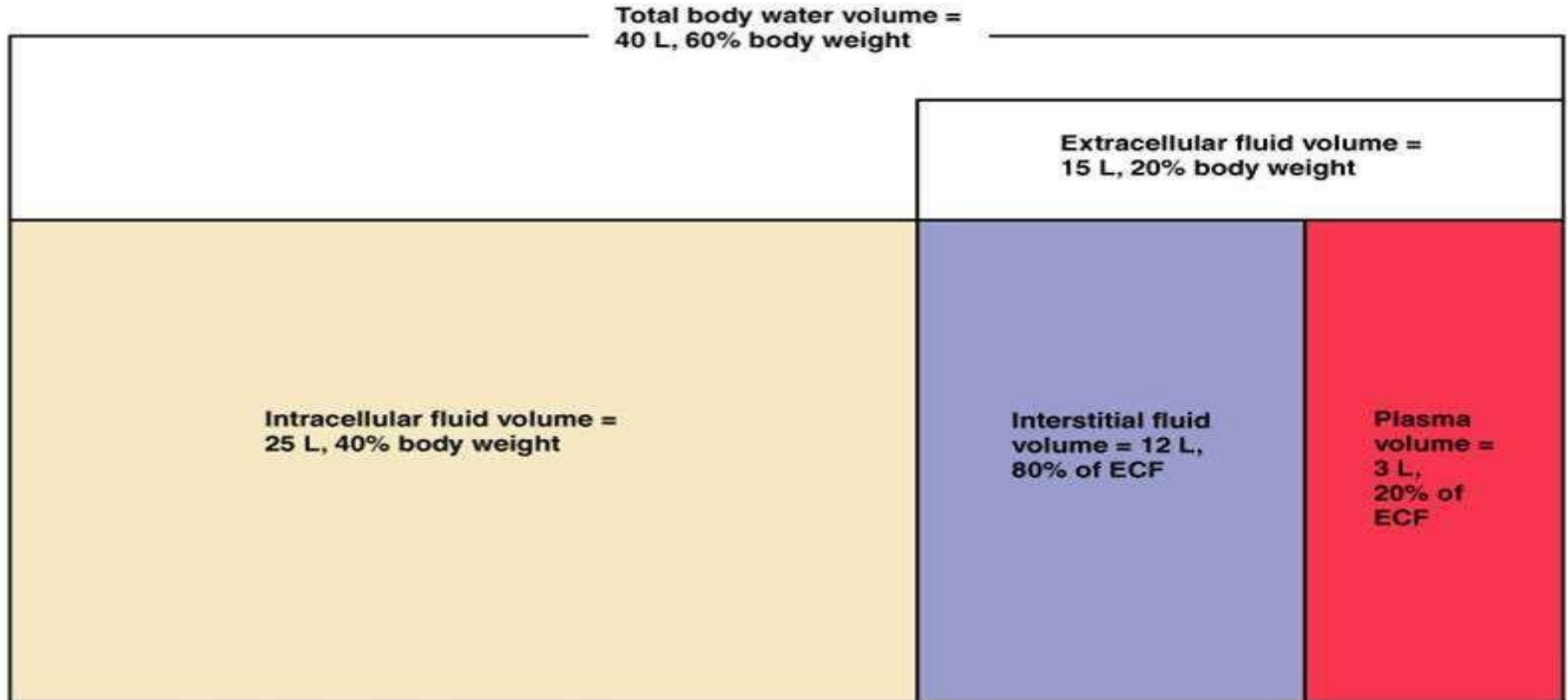
INTRODUCTION

- Once a drug enters into a blood stream the drug is subjected to a number of processes called as **DISPOSITION PROCESSES** that tend to lower the plasma concentration of drug.
- **DISTRIBUTION** is a process which involves **reversible transfer of a drug** between compartments.
- **ELIMINATION** is a process which involves **irreversible loss of drug** from the body.

DEFINITION OF DISTRIBUTION

- ✓ **DISTRIBUTION** is reversible transfer of a drug between the **ONE COMPARTMENT** (Blood) to **ANOTHER** (Extravascular fluids and tissues).
- ✓ Distribution is a passive process for which the driving force is **CONCENTRATION GRADIENT** between the blood and the extravascular tissues.

Drugs distribution in different compartments of body.



Transcellular fluid(2.5%)– CSF, intraocular, peritoneal, pleural, synovial, digestive secretion, fetus is also special type. Drugs will be in free or bound form.

Body compartment	Types of drugs
Total body water	Small, water soluble alcohol and antipyrine.
Extracellular space	Large, water soluble mannitol
Intravascular space	Very large, strongly protein bound, heparin
Body fat	Highly lipid soluble DDT and thiopentone
Bones	Fluoride and lead

**Distribution is a Passive Process,
for which the
Driving Force is
the Conc. Gradient
between the Blood
and Extravascular
Tissues**

- **The Process occurs by the Diffusion of Free Drug until equilibrium is established**

SIGNIFICANCE

- ⦿ Pharmacological action of drug depends upon its concentration at the site of action.
- ⦿ Distribution plays a significant role in –
 - a) Onset of action.
 - b) Intensity of action.
 - c) Duration of action.

STEPS IN DRUG DISTRIBUTION

- ⦿ Permeation of free drug present in blood through capillary wall and entry into extracellular fluid (ECF)
- ⦿ Permeation of drugs present in ECF through the membrane of tissue cells and into intracellular fluid.
- ⦿ This is **RATE LIMITING STEPS** and depends upon two factors:-
 - a) Rate of Perfusion to the extracellular tissue.
 - b) Membrane permeability of the drug.

DISTRIBUTION PROCESS

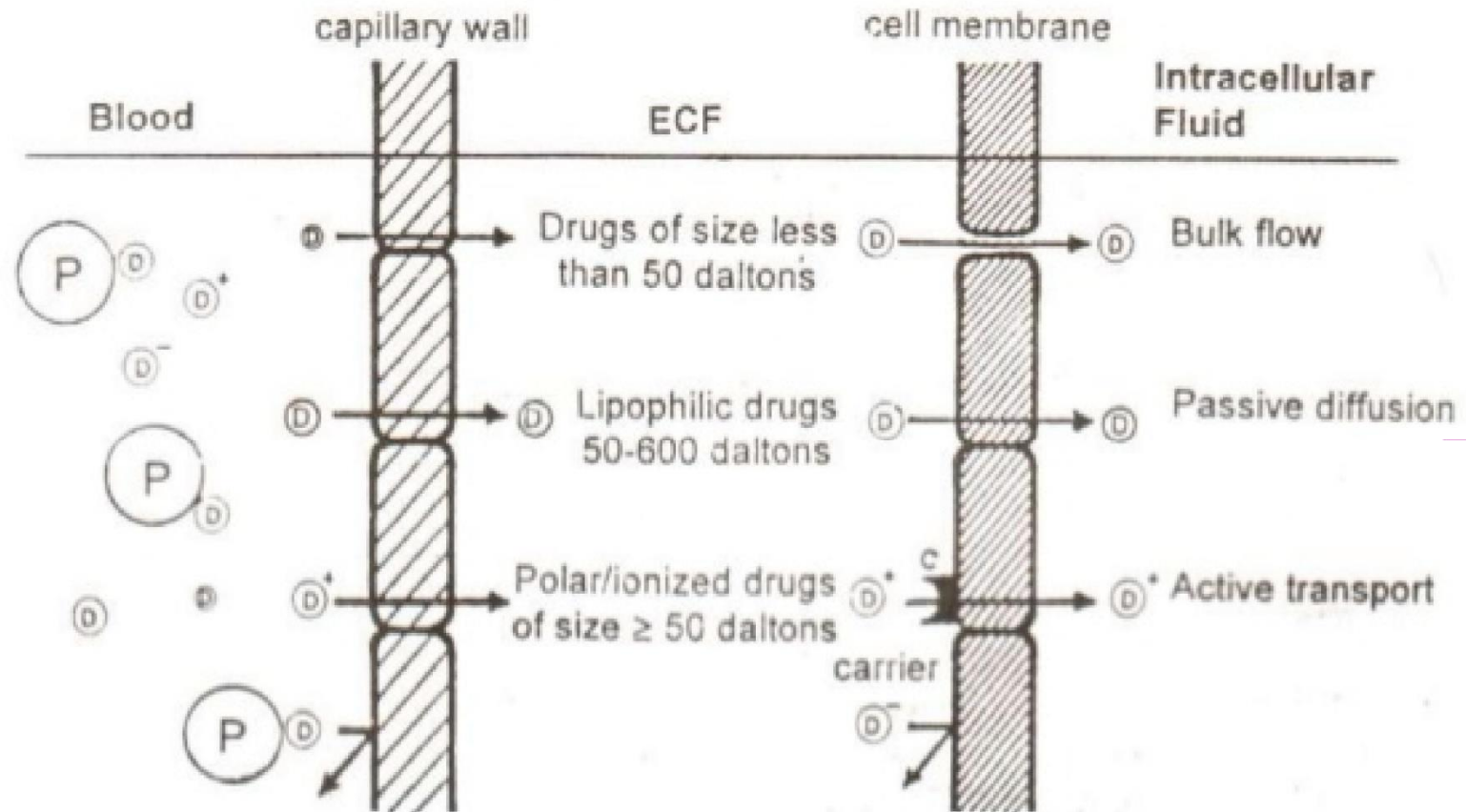


Fig. 3.3 Plasma membrane barrier and drug diffusion across it

Thank You

