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Routes of Drug Administration

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- A. Enteral
- **B.** Parenteral
- C. Others

- 1. Oral route (PO):
- The drug should be swallowed.
- Most commonly used route.
- Safest, most convenient, and most economical
- Duodenum is the major site of absorption, but other parts of GIT may be involved.

Disadvantages:

- a. The patient must be cooperative (compliant).
- b. Absorption is variable because of several factors affecting the rate and extent of absorption:
- Vomiting
- Failure of disintegration and dissolution

- First-pass effect
- Drug may be destroyed by gastric acid or intestinal flora.
- Food may delay absorption.
- Alteration in intestinal motility may affect absorption.
- Absorption may be affected by splanchnic blood flow.

- 2. Sublingual route (SL):
- Drug is placed under the tongue.
- Avoids first-pass effect.
- Used when a rapid onset is required such as in angina pectoris.
- Not commonly used.

- 3. Rectal route (PR):
- Avoids first-pass effect partially (~ 50%). Why?
- Useful in unconscious or vomiting patients.
- Absorption is often irregular, incomplete and unpredictable.
- Can be used for a local effect.

- Used for drugs poorly absorbed from, or unstable in the GIT.
- Used for rapid effect.
- Aseptic technique is required.
- 1. Intravenous route (IV):
- Bolus vs infusion.
- Rapid onset of action.
- Only aqueous solutions may be
 injected IV.

- Oily vehicles or those that precipitate blood constituents should not be given IV.
- No first-pass hepatic metabolism, the drug goes first to the right side of the heart, the lung, the left side of the heart, then to the systemic circulation.

Disadvantages:

- 1. Produce high initial concentration of the drug that might be toxic.
- 2. Once injected, the drug is there...??

- 2. Intramuscular route (IM):
- The drug is injected within muscle fibers of deltoid, gluteus maximus or vastus lateralis.
- Absorption of drug depends on blood supply (slower for g.m.
- Absorption is reduced in circulatory failure or shock.

- To be injected IM, the drug must be non-irritating to tissues.
- Can utilize:
- a. Aqueous solutions for fast absorption and rapid action.
- b. Depot preperations and suspensions for slow or sustained absorption (oily vehicles or ethylene glycol).
- Can accommodate large volumes.

- 3. Subcutaneous injections (SC, or SQ):
- The drug is injected under the skin.
- Absorption is affected by blood flow.
- Drug should be non-irritating to tissues.
- Absorption is slow and sustained.
- Accommodate smaller volumes than IM.

 Solid pellets can be implanted under the skin to produce effects over weeksmonths.

- 1. Inhalational or pulmonary route:
- Used for gaseous or volatile drugs, such as general anesthetics.
- Can also be used for solids that can be put in an aerosol, such as drugs for bronchial asthma.
- Drugs are absorbed across pulmonary epithelium and mucous membranes of respiratory tract.

- Absorption is rapid.
- Avoids first-pass effect.
- The lung acts as a route of elimination also.

- 2. Topical application:
- For a local effect on:

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a. mucous membranes: conjunctiva, nose, mouth, nasopharynx, oropharynx, vagina, rectum, colon, urethra, and urinary bladder.

b. skin: highly lipid-soluble drugs can be absorbed systemically.

Systemic absorption also occurs from abraded, burned and inflamed skin.

- 3. Transdermal route (TD):
- The drug is applied to the skin for systemic effect, such as in angina.
- For a sustained effect.
- Avoids first-pass metabolism.

ROUTE	BIOAVAILABILITY	ADVANTAGES	DISADVANTAGES
Parenteral Routes	the computer line (because	ne a. a.ur phenoispe hu	string mentation by Children
Intravenous bolus (IV)	Complete (100%) systemic drug absorption. Rate of bioavailability con- sidered instantaneous.	Drug is given for immedi- ate effect.	Increased chance for adverse reaction. Possible anaphylaxis.
Intravenous infusion (IV inf)	Complete (100%) systemic drug absorption. Rate of drug absorption controlled by infusion rate.	 Plasma drug levels more precisely controlled. May inject large fluid volumes. May use drugs with poor lipid solubility and/or irritating drugs. 	Requires skill in insertion of infusion set. Tissue damage at site of injection (infiltration, necrosis, or sterile abscess).
Intramuscular injection (IM)	Rapid from aqueous solution. Slow absorption from non- aqueous (oil) solutions.	Easier to inject than intra- venous injection. Larger volumes may be used compared to sub- cutaneous solutions.	Irritating drugs may be very painful. Different rates of absorp- tion depending on mus- cle group injected and blood flow.
Subcutaneous injection (SC)	Prompt from aqueous solu- tion. Slow absorption from repository formulations.	Generally, used for insulin injection.	Rate of drug absorption depends on blood flow and injection volume.

Buccal or sublingual (SL) Rapid absorption from lipid- No "first-pass" effects. Some drugs may be swalsoluble drugs. lowed.

Oral (PO)

Absorption may vary. Generally, slower absorption rate compared to IV bolus or IM injection. Safest and easiest route of drug administration. May use immediate-release and modified-release drug products.

Not for most drugs or drugs with high doses. Some drugs may have erratic absorption, be unstable in the gastointestinal tract, or be metabolized by liver prior to systemic absorption. Absorption may be erratic. Suppository may migrate to different position. Some patient discomfort.

Rectal (PR)

Absorption may vary from
suppository.Useful when patient can-
not swallow medication.More reliable absorption
from enema (solution).Used for local and systemic
effects.

Transdermal

Slow absorption, rate may
vary.Transdermal delivery system
(patch) is easy to use.Increased absorption with
occlusive dressing.Used for lipid-soluble drugs
with low dose and low

Some irritation by patch or drug. Permeability of skin variable with condition, anatomic site, age, and gender. Type of cream or ointment base affects drug release and absorption. Particle size of drug determines anatomic placement in respiratory tract. May stimulate cough reflex. Some drug may be swallowed.

Inhalation and intranasal Rapid absorption.May be used for local orTotal dose absorbed is variable.systemic effects.

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