PHARMACEUTICAL TECHNOLOGY

1. WEEK

What is Pharmaceutical Technology?

"All of the technologies that involve the development and use of drugs"

- Preparation, production, control of drug forms
- Properties of active substance release, interactions with human body
- Production, control of biotechnological, radiopharmaceutical, cosmetic and cosmeceutical productions

Pharmaceutical Technology,

is a science that encompasses all the processes for turning an active pharmaceutical ingredient into a medicine that can be used safely and effectively by patients. Fenni İspençiyari

Ottoman

Materia Medica

Rome

Galenic Pharmacy

Europe

Pharmaceutics

Anglo-Saxon Scandinavia

Pharmaceutics definition covers the issues of

- General pharmaceutical technology,
- Clinical pharmacology ,
- Biopharmaceutics and pharmacokinetics,
- Pharmacy applications
- Cosmetics

Industrial Pharmaceutical Technology

involves the areas given below:

- drug product preparation,
- methods of application of scientific basesto industry,
- the scientific basis of the instruments and machines used in the operations,
- pharmaceutical engineering and
- Effects of fabrication on drugs

Codex and Pharmacopoeia

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is a manuscript, official and antique book containing a list of chemical and medical items

Pharmacopoeia

They are official books containing the pharmaceutical active substances and the necessary properties for the preparation of medicinal forms prepared therefore for the protection of life and for therapeutic purposes.

The therapeutically effective amount of the active ingredients, the dosage forms prepared therefrom, the excipients used are given in pharmacopoeias.

They contain;

- Physical, chemical and physicochemical properties,
- Control and Identification reactions,
- Quantity assignments,
- Storage conditions,
- **■**Some formulations
- They are prepared and printed by the authorities assigned by each country. This ensures a legal standard.

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Formulary:

These are the active substances and dosage forms which are not important enough to enter the pharmacopoeia.

National Formulary (N.F.)
British National Formulary (B.N.F.)

Drug or pharmaceutical product?

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the medicine or other substance which has a physiological effect when ingested or otherwise introduced into the body.

Under the «Health Topics» title, WHO (World Health Organization) defines drugs under two different subtitles;

- essential medicines,
- pharmaceutical products,

which the latter is commonly called as drug or medicine.

- protecting the living from disease,
- Used with the aim of diagnosing and treating the diseases,
- Contain one (or more) active ingredient,
- Designed to be easily received by the patient,
- Prepared in the form of a formula (auxiliary substances) which will be effected according to the desired purpose and duration.

Pharmaceutical products:

More commonly known as medicines or drugs – are a fundamental component of both modern and traditional medicine. It is essential that such products are safe, effective, and of good quality, and are prescribed and used rationally.

Pharmaceutical products can be categorized as:

Magistral:

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This is the common name of medicines prepared by a pharmacist according to a prescription written by a doctor, veterinary doctor or a dentist.

Officinal:

Pharmacist usually prepares this kind of medicines according to the formulations given in codex or pharmacopoeia. They are prepared as stock formulations.

Pharmaceutical preparations:

These are the pharmaceutical products prepared in a factory after licenced by the health authority of the country TURKISH MEDICINES AND MEDICAL DEVICES AGENCY (TMMDA; TİTCK) is the authority in our country. These medicines can be over the counter (OTC) or can be given with a prescription.

structure of the pharmaceutical product

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- They can be classified human and veterinary according to the organism applied to
- They contain Active substance
 - Auxiliary substances / vehicles

Active substance (drug): is an organic / inorganic substance that provides the therapeutic and diagnostic purpose of the drug.

Excipient (vehicle): is the inert substance used in the formulation of the active substance and used according to the dosage form to be formulated.

According to their structure they can be classified as

- Natural
- Semisynthetic
- Synthetic

They can also be classified as,

>Simple:

Pharmaceutical product contains only one drug

>Composed:

Pharmaceutical product contains more than one drug

Excipient, vehicle

Excipient term is used for semisolids while vehicle term is generally used for solid dosage forms.

when the active ingredient is formulated with a suitable excipient as a pharmaceutical product:

- It can be easily taken by the patient,
- Dose is precisely adjusted,
- It is well absorbed,
- It stays long-lasting

can be done according to the

- structure of drug
- application site of product
- organs they are applied
- formulation and preparation techniques
- amount of active ingredient they

contain

prescription

application area of product

- ■Internal (oral)
- External

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parenteral



Oral use:

Syrups, capsules, tablets, efervescent powders, granules, suspensions etc..

External use:

Eye/ear preparations, creams, semisolids, suppositories, lotions, etc...

Parenteral use:

Enjectable preparations, solutions packaged in ampules or vials, serum solutions, dialysis solutions...

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- organs they are applied
 - Ophtalmic
 - Nasal
 - Otic
 - Rectal vaginal
 - Transdermal
- amount of active ingredient they contain
 - Adult dose
 - Pediatric dose (for children)





Formulation and preparation techniques

Solutions (syrups, elixirs etc)



Disperse systems

(colloidal preparations, suspensions, emulsions etc

- Semisolid dosage forms
 (ointments, creams, suppositories, ovules etc.)
- Solid dosage forms
 (tablets, granules, powders, capsules etc.)