UNIT-II ICD



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DISEASE

What is Disease?

- "A disease is a condition that deteriorates the normal functioning of the cells, tissues, and organs." Diseases are often thought of as medical conditions that are characterized by their signs and symptoms.
- The disease can also be defined as: "Any dangerous divergence from a functional or normal state of an entity."
- When a person is infected with a disease, he exhibits a few symptoms and signs that range from normal to severe depending upon the medical condition. Hence, in order to identify different diseases, the normalcy of an entity needs to be studied and understood as a clear demarcation between disease and disease-free is not always apparent.

Types of Diseases

Infectious Diseases

- Diseases that **spread from one person to another** are called **communicable diseases.**
- They are usually caused by microorganisms called pathogens (fungi, rickettsia, bacteria, viruses, protozoans, and worms).
- When an infected person discharges bodily fluids, pathogens may exit the host and infect a new person (sneezing, coughing etc). Examples include Cholera, chickenpox, malaria etc.

Non-infectious Diseases

- These diseases are caused by pathogens, but other factors such as age, nutritional deficiency, gender of an individual, and lifestyle also influence the disease.
- Examples include <u>hypertension</u>, diabetes, and cancer.
- They do not spread to others and they restrain within a person who has comes in contact with them.
- Alzheimer's disease, asthma, cataract and heart diseases are other non-infectious diseases.

Classification of Diseases

Classification of Diseases		
Туре	Explanation	Example
Anatomic Classification	This type refers to the affected organ or tissue	Heart disease
Topographical Classification	Further classified into types such as vascular disease, chest disease, gastrointestinal disease, and abdominal diseases. These are then handled by specializations in medicine that follow these topographical classifications	An ENT specialist (Ear- Nose-Throat) A Gastroenterology specialist etc.
Physiological Classification	This type includes diseases that affect a process or a function (such as metabolism,	Diabetes
Pathological Classification	This type considers the nature of the disease. For instance, cancer is associated with uncontrolled cell growth, and there are variations or types in the disease.	Neoplastic diseases (uncontrolled cell growth that is characteristic of cancer) Inflammatory diseases (autoimmunity)
Epidemiological Classification	This classification refers to the rate of occurrence, distribution and the control of the disease in a population.	Epidemic diseases such as the plague and Influenza pandemic of 1918– 1919

International Classification of Diseases (ICD)

- ICD is an international tool for the "standard diagnostic tool for management, epidemiology, and clinical purposes."
- ICD's official name is the international statistical classification of disease and health problems.
- ICD is maintained by WHO globally. WHO is coordinating and directing authority for health within the United Nations system.
- The ICD is properly designed as a health care classification system and it provides a very unique system of diagnostic code for classifying the disease which includes classification of a wide variety of signs, symptoms, social circumstances complaints an external cause of injury.

- This ICD system is specially designed to map health conditions for the corresponding generic categories together with a specific variation.
- The ICD is revised periodically and is currently in its 10th revision.
- The ICD-11 is the final draft which is expected to be submitted to the WHO's World Health Assembly (WHA) and it has the endorsement in 2015. This version was released on 18 June 2018.

- ICD (International Statistical Classification of Diseases and Related Health Problems), shortly known as International Classification of Diseases is the base/foundation for the identification of health trends and statistics.
- ICD is an international uniform standard for reporting diseases and health conditions.
- It is the diagnostic classification standard for all clinical and research purposes.
- ICD maps the human health condition from birth to death. No matter, however we die of, it will be recorded/coded under ICD.

Application

- Morbidity and mortality statistics
- Reimbursement (compensation/settlement) of systems
- Automated decision support in health care.
- The ICD is a major project to statistically classify all health disorders, and provide diagnostic assistance.

Historical synopsis

- In 1860, Florence nightingale →made first model of systemic collection of hospital data.
- In 1893, French physician, Jacques Bertillon → introduced Bertillon
 classification of cause of death
- In 1898, American public health association → recommended revision of ICD system every 10 years.
- The revision followed minor changes until 6th version of ICD \rightarrow morbidity and mortality condition and section on mental disorders
- WHO →responsibility for preparing and publishing the ICD revisions every 10 years.

General Principles of Disease Classification

- A statistical classification of diseases must be confined to a limited number of mutually exclusive categories. The categories have to be chosen to facilitate the statistical study of disease phenomena.
- A specific disease entity that is of particular public health importance or that occurs frequently should have its own category. Every disease or morbid condition must have a well-defined place in the list of categories.
- A statistical classification of diseases should retain the ability both to identify specific disease entities and to allow statistical presentation of data for broader groups, to enable useful and understandable information to be obtained.

Idea and Need for Classification in ICD:

Classification or Sorting of disease is needed due to the following facts:

- Determines the natural cause of the disease mechanism in a human body.
- Determines the **exact problem and solution of the disease**.
- Determines the normal stages and the critical stage of the disease
- Formulation of health statistics. ICD is the bedrock for health statistics.
- Concludes the **risk factors of the illness**.
- Calculates the **prevalence and incidence rate**, which ultimately helps to develop mitigating measures against the disease.
- Helps to categorize vulnerable disease.
- Determines the health cost and resources required to combat disease.
- Critical analysis of the disease based on the severity.

ICD YEARS

- ICD 1-1900
- ICD 2-1910
- ICD 3-1921
- ICD4-1930
- ICD5-1939
- ICD6-1949
- ICD7-1958
- ICD8A-1968
- ICD10-1999
- ICD11-2019

Volumes of ICD-10

Volume 1: Tabular list

 An alphanumeric listing of diseases and disease groups, along with inclusion and exclusion notes, some coding tabulation lists for mortality and morbidity, definition and regulation.

Volume 2: Instruction manual

- It provides
 - Instructions on how to use volume 1 and 3.
 - Guidelines for certification and rules for mortality coding
 - Guidelines for recording and coding for morbidity coding
 - Statistical presentation

Volume 3: Alphabetical index

• Is the comprehensive Alphabetical index of the diseases and conditions found in the tabular list Certain Infectious and parasitic diseases.[A00- B99]

CHAPTERS OF ICD-10

- Neoplasms [C00-D48]
- Diseases of the blood and blood forming organs and certain disorders involving the immune mechanism.
- Endocrine nutritional and metabolic diseases [E00-E90]
- Mental and behavioral disorder [F00-F99]
- Disorders of nervous system [G00-G99]
- Disease of eye [H00-H99]
- Disease of the ear and mastoid process [H60-H95]

The main innovations of ICD 10th Revisions are :

1. Relevant information for ambulatory(adapted)

and managed care encounter

- 2. Expanded injury codes
- 3. New combination codes for diagnosis /

symptoms to reduce the number of codes

needed to describe a problem fully.

ICD-10CM

- The ICD-10 Clinical Modification (ICD-10-CM) is a modification of the ICD-10, authorized by the World Health Organization, used as a source for diagnosis codes in the United States of America.
- It replaces the earlier ICD-9-CM
- The world health organization is currently revising the ICD towards ICD 11.
- The development is taking place on an internet based workspace called "icat " platform.
- The draft of the ICD -11 system → submitted to who's World Health
 Assembly (WHA)→ by 2017 and a final version → in 2018.

- Officially endorsed by all who members during the 72nd world health assembly on 25 may 2019.
- The ICD-11 officially came into effect on 1 JANUARY 2022
- Currently, **no country has officially switched** to the ICD-11 yet
- The ICD-11 is a large taxonomy consisting of about 85,000 entities, also called classes or nodes.
- An entity can be anything that is relevant to health care. It usually represents a disease or a pathogen.

Features :

- About 300 specialists of 55 countries, organized in 30 main working groups have provided their input to make ICD-11 scientifically up-to-date, and structural problems that became apparent using ICD-10 have been solved.
- ICD-11 is much easier to use than ICD-10.
- The systematic reliance on the use of code combinations and extension codes makes ICD finally clinically relevant.
- Primary care, cancer coding, traditional medicine and a section for assessment of functioning are now included.

Principles of classification of ICD

- The ICD is a variable-axis classification. The structure of ICD structure was developed of that proposed by William Farr during early days of international discussions. According to him, for all practical, epidemiological purposes, statistical data on diseases should be grouped in the following way:
 - epidemic diseases
 - constitutional or general diseases
 - local diseases arranged by site
 - developmental diseases
 - injuries

How are Diseases Classified in ICD?

The diseases are categorized:

On the basis of	Disease
Body system or region	Abdominal disease
	Gastrointestinal disease
Anatomic	Lung disease
	Liver disease
Function	Metabolic disease
Pathological/ nature of the disease process	Inflammatory disease
	Tumors formation
Etiologic/ cause	Bacterial infection
Epidemiology	Epidemic disease
Statistical	High prevalence and incidence rate disease.



ADVANTAGES

- Fast aggregation of health data
- To know the real time information about current health situation
- Wider analysis of community health
- Global health assessment

LIMITATIONS

- Discussion is limited to classified diseases
- Common term description of diseases
- Covers most common diseases
- Country specific modifications of certain diseases

Daily defined doses (DDD)

The basic **definition** of the defined daily dose (DDD) is:

- The DDD is the assumed average maintenance dose per day for a drug used for its main indication in adults.
- The DDD is a unit of measurement and does not necessarily reflect the recommended or Prescribed Daily Dose (PDD).

- Therapeutic doses for individual patients and patient groups will often differ from the DDD as they will be based on individual characteristics (such as age, weight, ethnic differences, type and severity of disease) and pharmacokinetic considerations.
- Only one DDD is assigned per ATC code and route of administration (e.g. oral formulation). The DDD is nearly always a compromise based on a review of available information including doses used in various countries when this information is available.

- The DDD is sometimes a "dose" that is rarely if ever prescribed, because it might be an average of two or more commonly used doses.
- Drug utilization data presented in DDDs only give a rough estimate of consumption and not an exact picture of actual use.
- DDDs provide a fixed unit of measurement independent of price, currencies, package size and strength enabling the researcher to assess trends in drug consumption and to perform comparisons between population groups.

Objectives of DDD: By applying DDD it is possible to

- Examine changes in drug utilization over time.
- Make international comparisons.
- Evaluate the effect of an intervention on drug use.
- Document the relative therapy intensity with various groups of drugs.
- Follow the changes in the use of a class of drugs.
- Evaluate regulatory effects and effects of interventions on prescribing patterns.

General principles for DDD assignment

- DDDs are only assigned to drugs with an ATC code and a DDD will normally not be assigned for a substance before a product is approved and marketed in at least one country.
- The basic principle is to assign only one DDD per route of administration within an ATC code.
- DDDs for single substances are normally based on monotherapy.
- Exceptions to this rule are given in the guidelines of the relevant ATC groups.

- For substances indicated for rare disorders with highly individual dosing schedules, the Working Group could decide not to assign a DDD.
- DDDs are not established for topical products, sera, vaccines, antineoplastic agents, allergen extracts, general and local anesthetics and contrast media.
- When a new DDD is assigned, various sources are used to get the best overview of the actual or expected use of a substance. The assigned DDD is based on the following principles:

- The average adult dose recommended for the main indication as reflected by the ATC code. When the recommended dose refers to body weight, an adult is considered to be a person of 70 kg. It should be emphasized that even special pharmaceutical forms mainly intended for children (e.g. mixtures, suppositories) are assigned the DDD used for adults. Exceptions are made for some products only used by children, e.g. growth hormones and fluoride tablets.
- For some groups of medicinal products specific principles for DDD assignment are established (e.g. the DDDs for the selective serotonin agonists in the treatment of migraine are based on the approved initial dose). These principles are given in the guidelines for the relevant ATC groups.

- The recommended maintenance dose (long term therapeutic dose) is usually preferred when establishing the DDD. The initial dose may differ from the maintenance dose but this is not reflected in the DDD.
- If the approved dose recommendation provides limited information about maintenance dose, the DDD will usually be the average of the maintenance dose range. Examples of interpretation of approved dose titration recommendations:
 - "Titrate up to a high dose if it is tolerated": the high dose would normally be chosen as the DDD.
 - "Consider to increase the dose only if efficacy is not satisfactory with initial dose": the DDD would normally be based on the initial dose.

- The treatment dose is generally used. If, however, prophylaxis is the main indication, this dose is used, e.g. for fluoride tablets (A01AA01) and some anti malaria.
- A DDD is usually established according to the declared content (strength) of the product. Various salts of a substance are usually not given different DDDs.
- Different stereoisomeric forms are normally assigned separate DDDs and ATC codes. The DDDs for stereoisomeric forms are described in the respective ATC groups.
- **Prodrugs**, have **not been given a separate ATC code**.

- The DDD is often identical for various dosage forms of the same drug. Different DDDs can be established when the bioavailability is substantially different for various routes of administration (e.g. oral and parenteral administration of morphine) or if the dosage forms are used for different indications.
- The DDDs are frequently used as indicators for antibacterial use in hospitals, and it has been decided that assigning different DDDs for oral and parenteral formulations could be important in some cases to improve the usefulness of the methodology in drug utilization monitoring and research.
- Parenteral products with different routes of administration (e.g. i.v. and i.m.) have the same DDD.

DDDs for combinations products:

- The DDDs assigned for combination products are based on the main principle of counting the combination as one daily dose, regardless of the number of active ingredients included in the combination.
- If a treatment schedule for a patient includes e.g. two single ingredient products, then the consumption will be measured by counting the DDDs of each single ingredient product separately.

• Example I:

Treatment with two products, each containing one active ingredient:

• Product A:

Tablets containing 20 mg of substance X (DDD = 20 mg)

• Product B:

Tablets containing 25 mg of substance Y (DDD = 25 mg)

- The dosing schedule 1 tablet of A plus 1 tablet of B daily will be calculated as a consumption of 2 DDDs.
- Example II:

Treatment with a combination product containing two active ingredients:

• Product C:

Tablets containing 20 mg of substance X and 12.5 mg of substance Y. The DDD of the combination products is assigned as 1 UD = 1 tablet.

• The dosing schedule 1 tablet of C daily will be calculated as 1 DDD (even though it will be equivalent to 1.5 DDD of the single active ingredients).

International Nonproprietary Name

• An international nonproprietary name (INN) is an official generic and non-

proprietary name given to a pharmaceutical drug or an active ingredient

- to make communication more precise by providing a unique standard name for

each active ingredient

- to avoid prescribing errors.
- The INN system has been coordinated by the World Health Organization (WHO) since 1953.
- Drugs are generally marketed by their brand names. Countries have their own systems of generic nomenclature for pharmaceutical substances. However owing to the various languages, there is a need to have a uniform standardized system of generic names that are accepted worldwide.

 The WHO administers an international generic nomenclature system called INNs also known as rINN, for recommended International Nonproprietary Name or p INN for proposed International Nonproprietary Name).

- A nonproprietary name is also known as a generic name.
- Initiated in 1950 by a World Health Assembly resolution
- Cumulative list of INN now stands at some 7000 names designated since that time .

AIMS OF INN SYSTEM

- To provide health professionals with a unique and universally available designated name
- To **identify** each pharmaceutical substance.
- For the clear identification, safe prescription and dispensing of medicines to patients.
- For **communication and exchange of information** among health professionals and scientists worldwide.
- Nonproprietary names are intended for use in Pharma, labeling, product information, advertising and other promotional material, drug regulation and scientific literature, and as a basis for product names, e.g. for generics.

SELECTION OF INN

• The names are selected by the World Health Organization on the advice of experts from the WHO Expert Advisory Panel.

STEPS:

- **a request/application** is made by the **manufacturer or inventor**; after a review of the request **a proposed INN** is selected and published for comments.
- after a time-period for objections has lapsed, the name will obtain the status of

 a recommended INN and will be published as such if no objection has been
 raised.
- Since the name is available in the public domain it may be used freely.
 However, it should not be registered as a trade-mark since this would prevent its use by other parties.

NAMING

- INN is designated for the active part of the molecule only to avoid the multiplication of entries in cases where several salts, esters, etc. are actually used. In such cases, the user of the INN has to create a modified INN (INNM).
- Ex: mepyramine maleate (a salt of mepyramine with maleic acid)

The primary principles for selection:

- Distinctive in sound and spelling; Not too long
- Not liable to confusion with other names in common use.
- Ex: -ac for anti-inflammatory agents, ibufenac derivatives -coxib for COX-2 inhibitors, a type of anti- inflammatory drugs (e.g. celecoxib)
- Do not select names for mixtures of substances.
- Do not select names for those substances that have a long history of use for medical purposes under well established names such as those of alkaloids (e.g. morphine, codeine), or trivial chemical names (e.g. acetic acid)

USE OF INNs IN INDIA

- In India, Section 13 (b) of the Trade Marks Act, 1999 prohibits the registration of names of chemical elements or INNs which have been declared by the World Health Organization (WHO) and notified by the Registrar of Trade Marks.
- Nevertheless, there are reported incidents of marketing of drugs under names that are similar to INNs. The WHO has brought the same to the attention of the Drug Controller General of India.
- The basic requirement of Section 13 (b) of the Trade Marks Act, 1999 is that INNs must be notified by the Registrar.