



SNS COLLEGE OF PHARMACY AND HEALTH SCIENCES Coimbatore -641035

COURSE NAME: Pharmaceutical Inorganic Chemistry (BP 104 T)

I YEAR / I SEM

TOPIC 1 : Origins of Pharmacopoeia

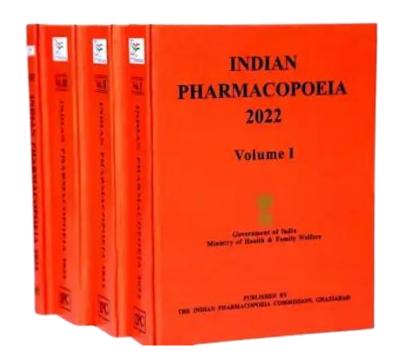






Definition & Purpose

- A pharmacopoeia is an official, authoritative compendium published by a recognized authority, typically a government or professional body, that establishes standards for the identity, quality, purity, strength, and packaging of pharmaceutical substances and medicinal products.
- ➤ It serves as a legal standard for ensuring the consistency and safety of drugs, guiding manufacturers, pharmacists, and regulatory bodies in maintaining high quality medicines.
- The primary purpose is to protect public health by minimizing variations in drug composition, preventing adulteration, and specifying analytical methods for quality control.



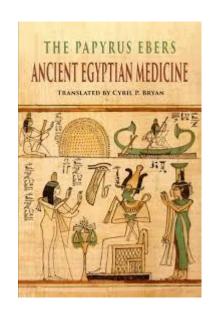




Ancient Egypt (Ebers Papyrus, 1550 BC)

- ➤ Ebers Papyrus (1550 BC)
- ➤ 20m scroll, 877 remedies for diseases
- ➤ Ingredients: Honey, aloe; used fresh for purity
- > Example: Pomegranate for worms







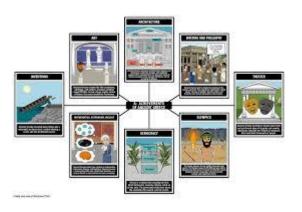




Contributions from Ancient Greece

- ➤ **Hippocrates (460–370 BC):** Father of Medicine; rational approaches; >200 drugs in Corpus Hippocraticum; emphasized purity
- > Theophrastus (371–287 BC): Enquiry into Plants; classified >500 medicinal plants; prevented impure substitutions
- ➤ Dioscorides (60–78 AD): De Materia Medica; 600+ plants, 100 animals, 80 minerals; guidelines for collection/storage; sensory adulteration tests (e.g., opium from unripe poppies)





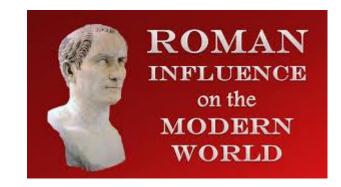


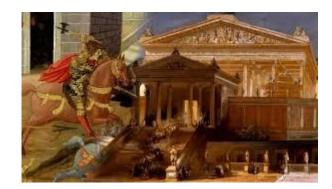


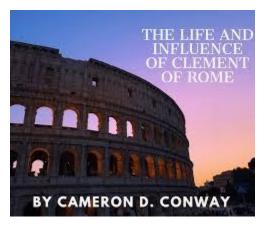
Roman Influences

- Galen (129-216 AD): Developed galenicals
- Over 400 remedies with precise compounding
- Introduced percolation, expression; warned of mold
- ➤ Theriac: 64-ingredient antidote, standardized proportions











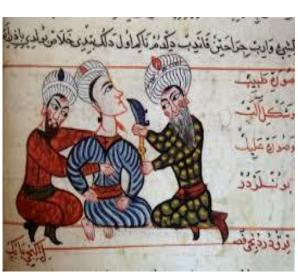


Medieval Developments in the Arab World

- ➤ Rhazes (865-925 AD): Kitab al-Mansuri with purity via distillation
- Avicenna (980-1037 AD): Canon with 760 drugs, taste/color impurity tests
- > Introduced alcohol distillation, warned of mercury toxicity











Renaissance and Early Modern Pharmacopoeias in Europe

- > Renaissance: Shift to printed pharmacopoeias
- ➤ Nuovo Receptario (1498, Florence): 168 standardized recipes
- ➤ London Pharmacopoeia (1618): 1,028 simples, 932 compounds
- Regional inconsistencies noted (e.g., Edinburgh 1699, Dublin 1807)







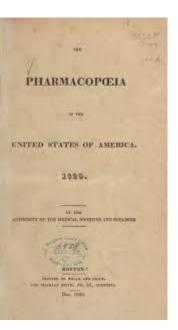


Establishment of the United States Pharmacopeia (1820)

- > Founded January 1, 1820, by 11 physicians led by Lyman Spalding
- > First edition (Dec 1820): 217 drugs, bilingual (English/Latin)
- > Set standards for identity, purity (e.g., ignition tests), strength
- > Reduced adulteration risks, e.g., calomel purity with lime water test





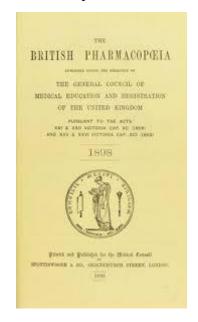




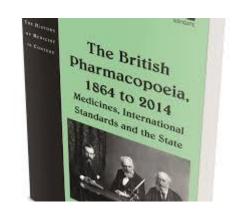


Formation of the British Pharmacopoeia (1864)

- First published in 1864 by General Medical Council
- ➤ Unified London, Edinburgh, Dublin standards
- ➤ 311 preparations with purity tests for arsenic
- ➤ Required ≥31.8% hydrochloric acid consistency







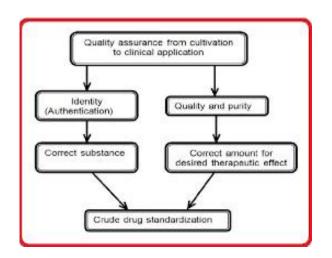




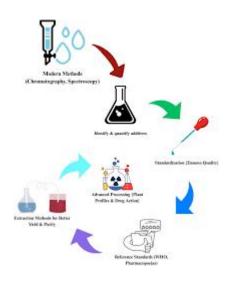


Role in Standardizing Drug Purity

- Pivotal role via mandatory tests and impurity limits
- > Evolved from sensory tests (e.g., taste for saffron) to chemical assays (USP 1820)
- ➤ BP 1864 added quantitative limits (e.g., <0.005%)











History of Pharmacopoeia:

Figure 1. Timeline indicating the year when many of the pharmacopoeias were created.

