

Quality control of Crude Drugs

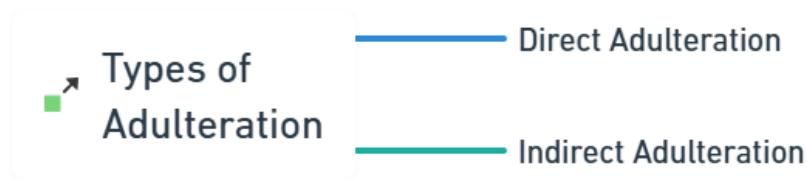
Adulteration of Drugs of Natural Origin

- **Adulteration** refers to the intentional or unintentional addition, substitution, or removal of substances in a drug, compromising its quality and safety.

Types of Adulteration

- Adulteration refers to the contamination or addition of inferior substances to food, beverages, or other products, making them unsafe for consumption.

It can be broadly classified into two types:



1. Direct Adulteration

- Direct adulteration occurs when harmful substances are deliberately added to food products to increase quantity, enhance appearance, or reduce costs.
- These substances are often toxic and pose serious health risks.

Examples of Direct Adulteration:

- **Addition of non-edible colors** – e.g., Metanil yellow in turmeric
- **Mixing toxic chemicals** – e.g., Formalin in milk and fish
- **Use of non-food-grade additives** – e.g., Washing powder in ice cream
- **Blending with inferior substances** – e.g., Starch in milk or ghee

2. Indirect Adulteration

- Indirect adulteration occurs unintentionally due to environmental contamination, improper handling, or poor storage conditions.
- It usually happens during production, processing, transportation, or packaging.

Examples of Indirect Adulteration:

- **Pesticide residues in fruits and vegetables** – e.g., DDT, lead, mercury
- **Heavy metal contamination in water** – e.g., Arsenic, cadmium
- **Oil rancidity due to improper storage** – leading to toxic effects
- **Rodent or insect contamination** – e.g., Presence of insect parts in grains

Causes of Adulteration

- **Economic Gain:** Lower production costs by using cheaper materials.

- **Lack of Regulation:** Inadequate enforcement of quality standards.
- **Misidentification:** Errors in plant or species identification leading to the use of incorrect materials.
- **Environmental Factors:** Contamination from soil, water, or air pollutants.

Examples of Adulteration

A. Plant-Based Drugs:

- **Turmeric (*Curcuma longa*):** Adulterated with metanil yellow (a synthetic dye) to enhance color.
- **Ginger (*Zingiber officinale*):** Adulterated with wheat flour or other fillers to increase weight.

B. Animal-Based Drugs:

- **Honey:** Adulterated with sugar syrups or corn syrup.
- **Insulin:** Originally extracted from animal pancreases, potential contamination if not properly purified.

Detection and Prevention of Adulteration

- **Strict Regulatory Frameworks:** Implementing and enforcing standards like the United States Pharmacopeia (USP) or the European Pharmacopoeia.
- **Supplier Verification:** Ensuring that raw materials are sourced from reputable suppliers.
- **Good Manufacturing Practices (GMP):** Adhering to GMP to minimize contamination and ensure consistency.
- **Advanced Analytical Techniques:** Utilizing sophisticated methods for detection (discussed in evaluation methods below).

Organoleptic (Sensory) Evaluation of Drugs

Definition

- Involves the use of the **sense organs** (sight, smell, taste, touch) to inspect the crude drug.

Key Properties Assessed

1. **Appearance** (color, shape, size): e.g., bright yellowish-orange for turmeric rhizome.
2. **Odor:** e.g., characteristic pungent-spicy smell of clove.
3. **Taste:** e.g., bitter taste of quinine bark.
4. **Texture:** e.g., brittle leaves of senna.

Significance

- **Preliminary screening** for authenticity and obvious adulteration.
- **Quick and cost-effective** check.

Limitations

- Subjective and requires **experience**.
- Cannot detect subtle adulterants.

Microscopic Evaluation of Drugs

Definition

- **Microscopic examination** of crude drugs (whole or powdered) to study **histological** or **cellular** features.

Procedure

1. Preparation of **thin sections** (e.g., transverse or longitudinal) of the plant part.
2. **Staining or clearing** the sample (e.g., with chloral hydrate, phloroglucinol + HCl).
3. Observation of **distinctive tissue structures** (e.g., trichomes, stomata, vascular bundles, calcium oxalate crystals).

Key Properties Assessed

- **Diagnostic cell structures** (types of fibers, vessels, glands).
- **Characteristic inclusions** (starch grains, crystals, pigments).

Significance

- **Definitive identification** of plant drugs.
- Detection of **foreign matter** (e.g., other plant tissues, insects).

Limitations

- Requires **expertise** in plant anatomy.
- May not reveal chemically similar adulterants if they share similar microscopic features.

Physical Evaluation of Drugs

Definition

- Measurement of **physical parameters** and constants that help gauge purity and identity.

Common Physical Tests

1. **Foreign Matter Content:** Visual or sieve-based inspection.
2. **Moisture Content / Loss on Drying:** Prevents microbial growth, ensures stability.
3. **Ash Values**
 - **Total Ash:** Inorganic residue after incineration.
 - **Acid-Insoluble Ash:** Detects silica and dirt.
 - **Water-Soluble Ash.**
4. **Extractive Values:** Water-soluble, alcohol-soluble extractives indicating content of soluble constituents.
5. **Volatile Oil Content:** For aromatic drugs (clove, cinnamon).
6. **Melting/Boiling Point:** Helps identify purified natural compounds (e.g., camphor).
7. **Specific Gravity / Optical Rotation:** Characteristic for certain oils and solutions.

Significance

- **Quantitative** measure of purity (limits set in pharmacopoeias).
- **Quality control** benchmark across batches.

Limitations

- Does not definitively confirm **chemical identity** of active constituents.
- Some tests can vary based on environmental or processing factors.

Chemical Evaluation of Drugs

Definition

- **Qualitative** and **quantitative** chemical analyses to identify and measure **active constituents** or marker compounds.

Approaches

1. Simple Chemical Tests:

- Color reactions (e.g., Mayer's reagent for alkaloids, Ferric chloride test for phenolics).

2. Chromatographic Techniques:

- **TLC / HPTLC:** Quick, cost-effective separation and identification.
- **HPLC / GC:** More precise, can quantify individual components.

3. Spectral Methods:

- **UV-Vis, IR, NMR, Mass Spectrometry** for structural and purity analysis.

4. Assay Methods:



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