

Formulation Additives

Study of Different Formulation Additives

- Formulation additives (also called excipients) are substances added to a drug product to help in its preparation, stability, effectiveness, and patient acceptability.
- They do not have therapeutic action but are essential for proper drug delivery.

Main categories of formulation additives

Formulation Additives



1. Diluents (Fillers)

- Used to increase the bulk of the formulation when the drug dose is very small
- Make tablets or capsules of practical size

Examples:

- Lactose
- Microcrystalline cellulose
- Dicalcium phosphate

Functions

- Improve compressibility
- Provide uniformity in tablet size and weight

2. Binders (Adhesives)

- Help powder particles stick together to form granules
- Provide mechanical strength to tablets

Examples

- Starch paste
- PVP (polyvinylpyrrolidone)
- Gelatin

Functions

- Improve tablet hardness
- Prevent tablet breakage

3. Disintegrants

- Help tablets break apart after ingestion
- Ensure drug is released quickly for absorption

Examples

- Starch
- Sodium starch glycolate
- Croscarmellose sodium

Mechanism

- Swell when in contact with water
- Create pressure that breaks tablet

4. Lubricants and Glidants

Lubricants

- Reduce friction between tablet and machine parts
- Prevent sticking during compression
- **Examples**
 - Magnesium stearate
 - Stearic acid

Glidants

- Improve flow of powder
- **Examples**
 - Talc
 - Colloidal silica

5. Coating Agents

- Applied to tablet surface

Functions

- Protect drug from environment (moisture, light)
- Mask unpleasant taste
- Control drug release (enteric coating)

6. Preservatives

- Preventing microbial growth

Examples

- Parabens
- Benzalkonium chloride

7. Coloring and Flavoring Agents

- Improve appearance and taste
- Increase patient compliance

8. Surfactants

- Improve wetting and solubility

Examples

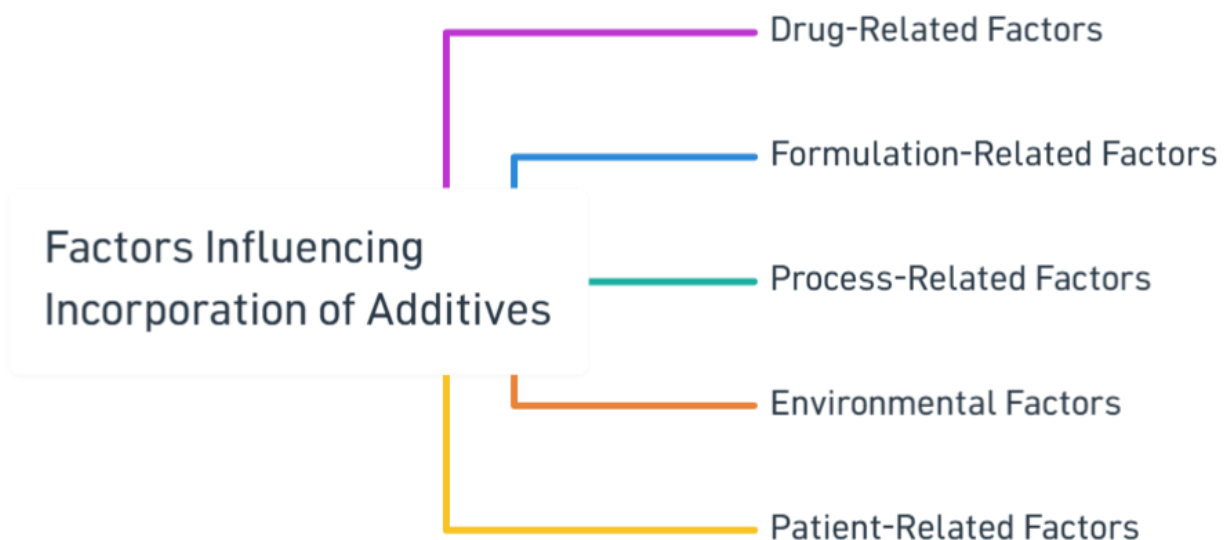
- Sodium lauryl sulfate

Important Points

- Excipients must be safe and non-toxic
- Must be compatible with drug
- Should not affect drug action

Factors Influencing Incorporation of Additives

Not all excipients can be added freely their selection and amount depend on several factors related to the drug and the formulation process.



1. Drug-Related Factors

1. Drug Dose

- Low dose → more diluent needed
- High dose → less excipient space

2. Drug Solubility

- Poorly soluble drugs need solubilizers or disintegrants
- Highly soluble drugs need controlled release agents

3. Drug Stability

- Sensitive drugs require stabilizers
- Avoid excipients that cause degradation

4. Drug Compatibility

- Must not react chemically with excipients

2. Formulation-Related Factors

1. Type of Dosage Form

- Tablet, capsule, liquid, injection
- Each requires different excipients

2. Route of Administration

- Oral → flavoring agents
- Injectable → must be sterile and pyrogen-free

3. Release Profile

- Immediate release
- Sustained/controlled release

3. Process-Related Factors

1. Manufacturing Method

- Direct compression
- Wet granulation
- Dry granulation

2. Compression Force

- Affects binder and lubricant selection

3. Mixing Time

- Overmixing may affect lubricant performance

4. Environmental Factors

- Temperature
- Humidity
- Light

5. Patient-Related Factors

- Age (children need sweeteners)
- Allergies (avoid certain excipients like lactose)

Outcome

- Proper excipient selection ensures stability, effectiveness, and manufacturability

Role of Formulation Development and Processing

- Formulation development is the process of designing drug products that is safe, effective, stable, and easy to manufacture.

Stages of Formulation Development

1. Preformulation Studies

- Study drug properties
- Helps in selecting excipients

2. Prototype Formulation

- Initial trial formulation
- Tested for basic properties

3. Optimization

- Adjusting composition for best results
- Improving stability, dissolution, and manufacturability

4. Scale-Up

- Moving from lab scale to industrial scale
- Ensures reproducibility

Role of formulation development

1. Selection of excipients

- Choose suitable additives based on drug properties
- Ensure compatibility

2. Design of dosage form

- Tablet, capsule, suspension, injection
- Depends on drug and therapeutic need

3. Optimization of drug release

- Immediate or controlled release



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