

# Microencapsulation

• It applies a relatively thin coating to small particles of solids or droplets of liquids.

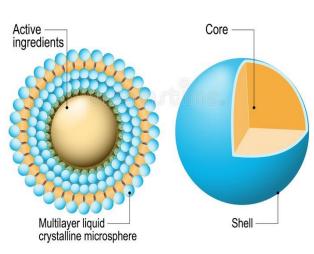
#### Advantages of microencapsulation:

- 1. The smallness of the particles, so, the active ingredient can be widely distributed throughout the GIT, thus improving drug absorption.
- 2. Taste-masking
- 3. Formulation of tablets or capsules containing incompatible ingredients.

#### Disadvantages or challenges:

- 1. No single microencapsulation process is useful for all materials.
- 2. Difficulties include **incomplete or discontinuous** coating and inadequate stability of sensitive drugs.

Microencapsulation



# **Core and Coating materials**

- The **core** material (the drug particles) can be liquid or solid in nature.
- The solid core material can be an active ingredient alone or as a mixture with diluents and other excipients.

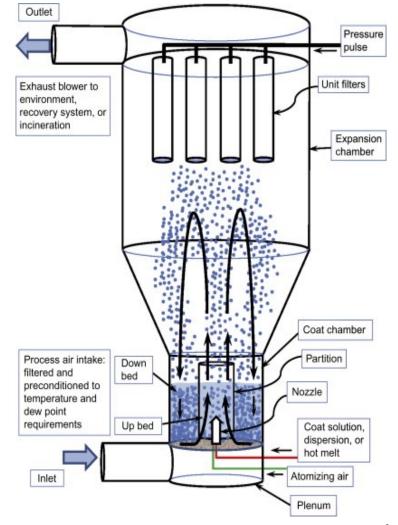
- The **coating material** should be:
- 1. Capable of **forming a film** that is cohesive with the core material.
- 2. Be chemically **compatible** and noncreative with the core material.

Examples of	Characteristic	Purpose of	Final Dosage
Core Material		•	Form
Core Material	Property	Encapsulation	FOITH
Acetaminophen	Slightly water	Taste masking	Tablet
	soluble solid		
Aspirin	Slightly water	Taste masking, sustained	Dry powder
!	soluble solid	release, reduced gastric	
		irritation.	
Islet of Langerhans	Viable cells	Replacement therapy for	Injection
		diabetic patients	
Isosorbide dinitrate	Water soluble	Sustained release	Capsule
	solid		
Menthol, methyl	Volatile solution	Reduction of volatility;	Lotion
Salicylate, camphor		sustained release	
mixture			
Vitamin A	Nonvolatile	Stabilization to prevent	Dry powder
palmitate	<mark>liquid</mark>	oxidation	

(NOT FOR SAVE).

- 1. Air suspension: (for solid core only)
- This method consists of suspending the solid material in the air and spraying the coating material with the air.

• However, this process generally is considered to be suitable only for the encapsulation of solid core materials.



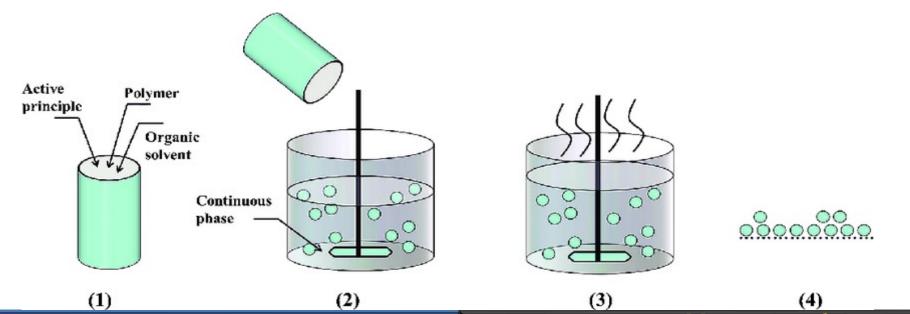
## 2. Pan Coating: (only for solids)

• In this method, the coating is applied as an atomized spray to the desired solid core material in the coating pan. To remove the coating solvent, warm air is used in a process similar to that of tablet coating.



# Solvent Evaporation (continue)

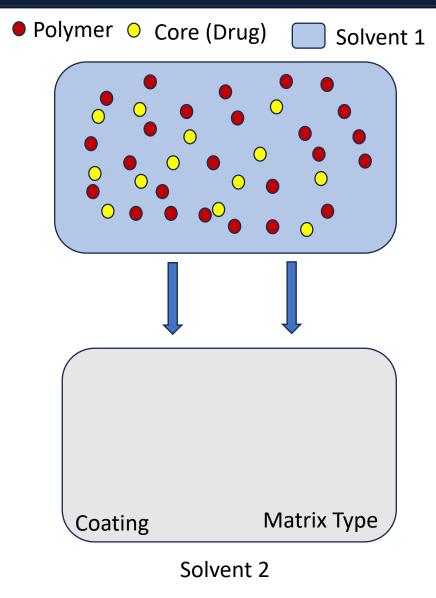
- The process includes preparing two solutions:
- Solution 1: The coating polymer is dissolved in a volatile solvent.
- Solution 2: contains a solvent that is: 1) Immiscible with solution 1. 2) Coating polymer and core material are insoluble in this solvent.
  - Solution 1 is added with agitation to solution 2,
  - Solvent 1 is removed by evaporation with or without the aid of heat. The polymer will shrink on itself and **coat** the core material.



## 3- Solvent Evaporation

- The core material is dissolved or dispersed in a coating polymer solution (in solution 1)
  - 1. If the core material (drug) was (dissolved (soluble) in **solution 1** it will form a matrix-type microcapsule (mixture) with polymer (not coating)

- Note:
- 1. This method is suitable for **liquid and solid core** material



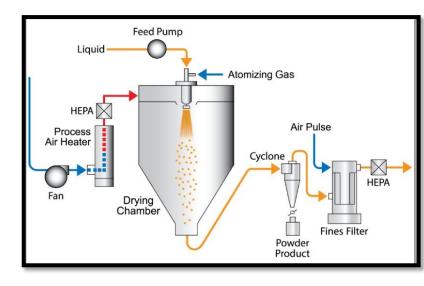
# Spray drying and spray congealing:

- These methods can be used for both **liquid and solid** drugs.
- Both involve:
- 1. Dispersing the core material in a liquid coating material.
- 2. Spraying the core-coating mixture (drug + polymer) into certain environmental conditions achieves rapid solidification of the coating.
- The principal <u>difference</u> between the two methods is how the solidification is achieved.
- 1. **Spray drying** is achieved by rapid evaporation (heating) of a solvent in which the coating material is dissolved.
- 2. Spray congealing is accomplished by thermally congealing (cooling) a molten coating material.



## Spray Drying:

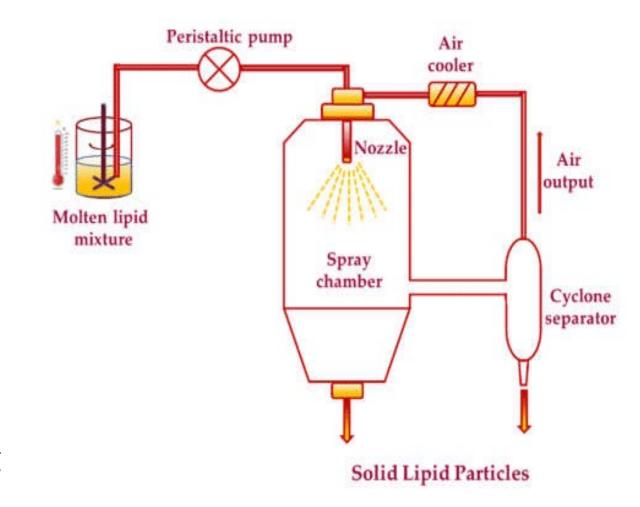
- Spray drying is done by **dispersing** a core material in a coating solution, **in which the core material is insoluble**, and then atomizing the mixture into an air stream.
- The air (**hot air**) supplies the heat of vaporization required to remove the solvent from the coating material, thus forming the microencapsulated product.
- The equipment used for this purpose is the usual spray dryer.





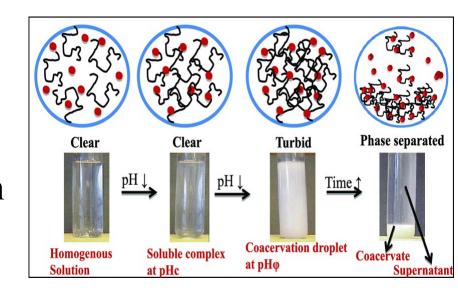
## Spray congealing:

- The core material is dissolved in a **hot** coating solution.
- Cold air is used to solidify the coating polymer on the core material.
- Waxes, fatty acids, and certain polymers which are **solids at room temperature** but meltable at high temperatures, apply to the spray congealing technique.
- Note: spray drying and spray congealing can be used for solid and liquid core materials



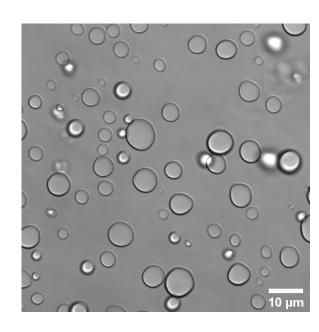
## Coacervation-Phase separation:

- It can be used for **both liquid and solid** drugs.
- It consists of three steps:
- Step 1: Formation of three immiscible phases:
- Which are the
  - Liquid manufacturing vehicle phase.
  - Coating polymer phase.
  - Core material phase
- To form the three phases, the polymer is **dissolved** in the liquid manufacturing phase (**phase 1+ phase 2**), then the core material is **dispersed** into that liquid (**phase 3**).
  - Until this point we have **two phases** (polymer + vehicle) and dispersed core material.

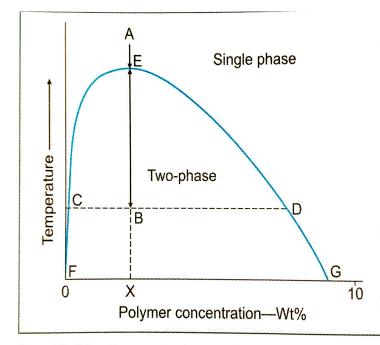


# Step 1 (continue)

- To form three immiscible phases: the solubility of the polymer (the coating material) in the solvent is altered to form a third immiscible phase.
- This polymer-rich phase will contain **most** of the polymer in the solution and it is called a **coacervate**.
- The process of coacervation-phase separation (phasing out the polymer) is formed by utilizing one of the methods:
- by changing the **temperature** of the polymer solution.
- or by adding salt.
- or adding a **nonsolvent (another solvent)**.
- or adding another polymer.



- Step 2: Deposition of the coating.
- It consists of depositing the liquid coating material on the core material under controlled agitation by: for example further change in temperature.
- Deposition of the liquid polymer coating around the core material occurs if the coating polymer is adsorbed at the interface between the core material and the immiscible solvent phase, and this adsorption phenomenon is a prerequisite to effective coating.
- Step 3: Rigidizing the coating.
- It involves rigidizing the coating, usually by thermal techniques, to form the microcapsules.



ig. 15.10: General phase diagram-coacervation duced thermally

# **Example on Phase Separation Technique**

- Ethyl cellulose (a water-insoluble coating polymer) is applied to N-acetyl-p-aminophenol (paracetamol) powder (core material) by utilizing the temperature characteristics of the polymer in the cyclohexane (solvent).
- Ethyl cellulose is insoluble in cyclohexane at **room temperature** but soluble at elevated temperatures. The ethyl cellulose and cyclohexane mixture is **heated** to form a homogeneous (one-phase) solution.
- The aminophenol is **dispersed** (as an insoluble powder) in the solution by stirring.
- Allowing the mixture to cool with continuous stirring results in coacervationphase separation of the ethyl cellulose from cyclohexane and microencapsulate the core material.
- Allowing the mixture to cool further to room temperature causes gelation and solidification of the coating.
- The microencapsulated product can then be collected from the cyclohexane by filtration or centrifugation.