

A close-up photograph of a person's face in profile, using a white inhaler. The person's mouth is open, and the inhaler is held between their lips. The background is a solid light blue color.

# Pharmaceutical Aerosols

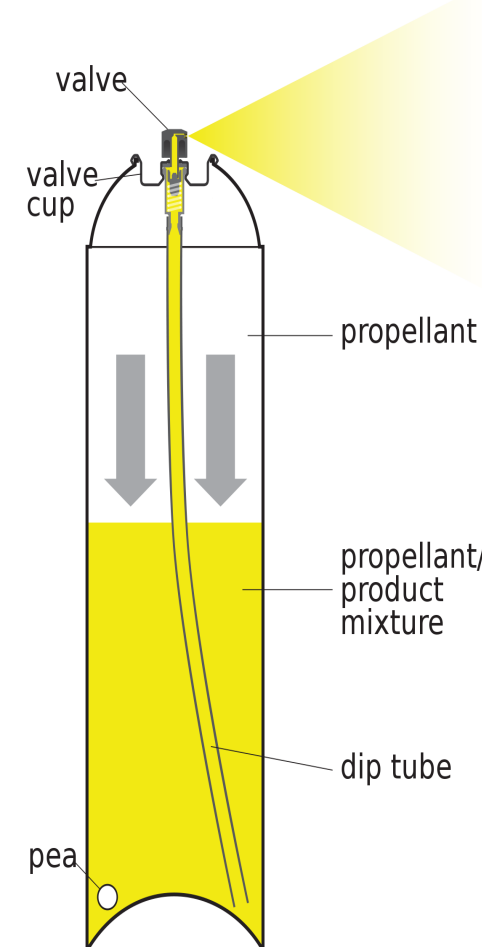
*Mohammed Albarki, PhD.*

# Pharmaceutical Aerosols

- A system that depends on the power of compressed **pre-liquefied gas** to expel the contents from the container.

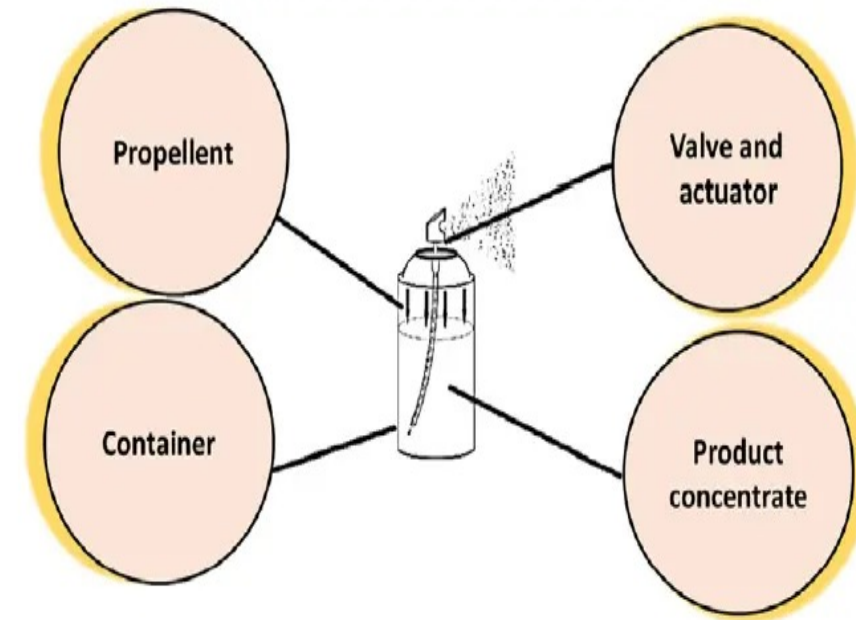
## Advantages:

1. The dose can be **delivered directly** to the site of action such as an inhaler.
2. Relatively **easy to use**.
3. **Rapid** onset of action.
4. **Avoid first-pass** metabolism and degradation in the GI tract.
5. **A lower dose** is used, especially in the case of **steroids**, in which most of the steroid reaches the respiratory tract and less is swallowed.
6. **An alternative route is** when a therapeutic agent may interact chemically or physically with other medicinal needs **concurrently**.
7. Container and valve closure are **tamperproof**.
8. The dose can be adjusted accurately using the **metered valve**.



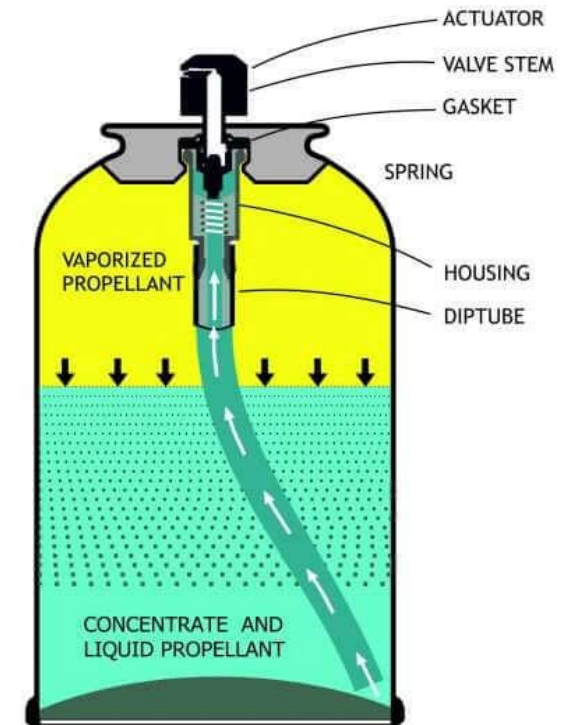
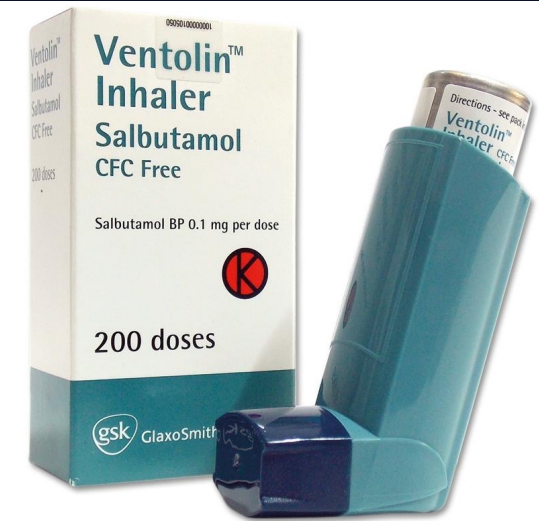
# Component of Aerosol Package

- **Actuator and valve, propellant, drug concentrate, and container.**
1. **Product concentrate** consists of an active ingredient, or a mixture of active ingredients and other necessary agents such as solvents, antioxidants, and surfactants.
  2. **Propellant:**
    - Responsible for developing the proper **pressure** within the container, and it **expels** the product when the valve is opened.
      1. It aids in the atomization or foam production of the product.
      2. Pharmaceutical aerosols usually use a blend of propellants, which allows us to achieve the desired **vapor pressure**.
      3. Act as a solvent or diluent



# Types of Propellants

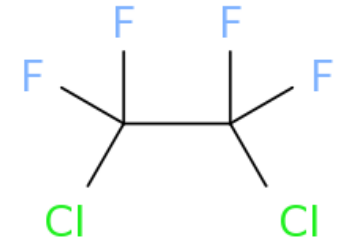
- **Liquified gas:**
  - Chlorofluorocarbons (CFCs)
  - Hydrochlorofluorocarbons (HCFCs).
  - Hydrofluorocarbons (HFCs).
  - Hydrocarbons.
- **Compressed gas** (used for products such as hair preparations).
  - Nitrogen ( $N_2$ )





# Liquified Gases

- Used for **orally** administered aerosols.
- These gases are **turned into liquid** by **decreasing** the temperature below boiling point and/or increasing the pressure.
- When in a sealed container, the liquified gas will be in **two** phases which are **liquid and vapor**.
- Has the advantage of maintaining a constant pressure inside the container and effectively dispersing the active ingredients.
- **Chlorofluorocarbons CFC**.
- Defined by the global numerical number which depends on the number of **C**, **F**, and **H**. For example diclorotetrafluoroethane ( $C_2Cl_2F_4$ ) = C 2-1, H 0+1, F 4, so it's called **propellant 114**.
- **Nontoxic**, and **non-flammable**.
- **Limited use nowadays** due to the effect on the ozone layer. And replaced with other types.



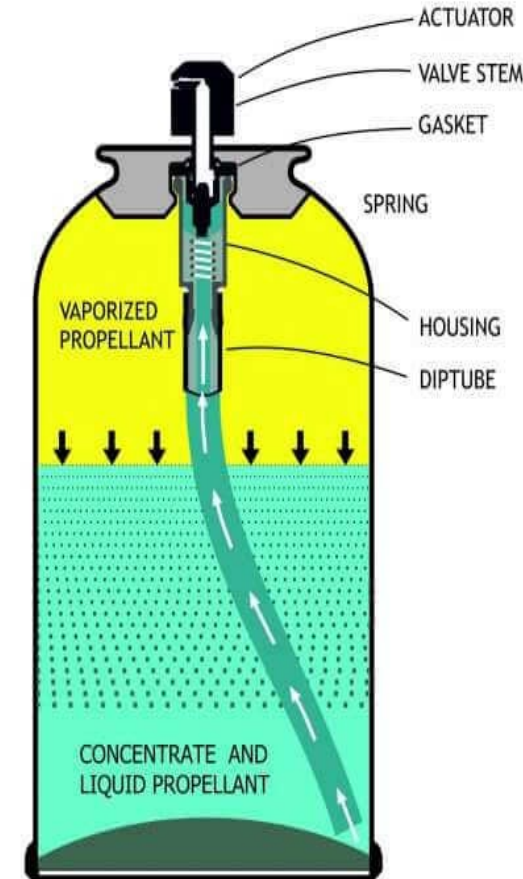
# Hydrocarbons

- Used mainly for **topical** application
- As **compared** to fluorinated hydrocarbons:
  - They are **flammable**, but **less toxic** and more economical, more soluble, and more chemically stable.
  - It has **little** expansion pressure and will produce a **wet spray**.
- Examples:
- Butane, propane



# Propellant (continue)

- **Propellants** may be a single type or a blend of various propellants. The propellant is selected to give the desired **vapor pressure, solubility, and particle size**.
- Vapor pressure can be controlled to the desired pressure by **mixing** another propellant with different properties (different vapor pressure)
- The total vapor pressure of the mixture is the sum of vapor pressures of the mixture component (**Dalton Law**):
- $P_{\text{total}} = P_a + P_b$



# Vapor pressure of Blend of Propellant

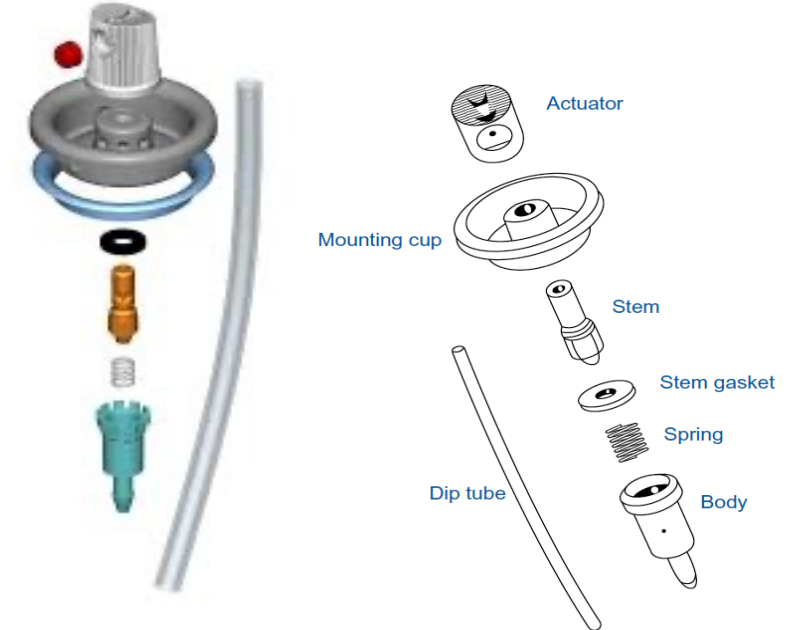
- The partial pressure of each component is calculated according to **Raoult's** law.
- In ideal behavior, this can be represented by the following equations for propellant A and Propellant B:
- $P_a = \left( \frac{n_a}{n_a + n_b} \right) * P_{A^o} = N_A * P_{A^o}$  **Partial pressure for propellant A,**
- $n$ = moles of propellant;  $P_{A^o}$  = vapor pressure of pure propellant A;  $N_A$ = mole fraction of component A
- For **propellant B** the equation will be:
- $P_b = \left( \frac{n_b}{n_a + n_b} \right) * P_{B^o} = N_B * P_{B^o}$  **Partial pressure for propellant B**



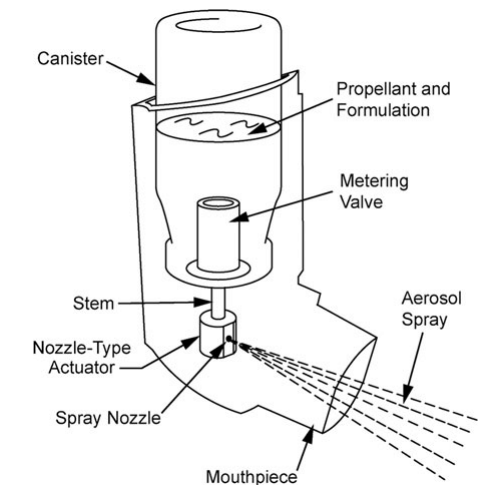
# Component of Aerosol Package

## 3. Valves

- It is a multifunctional part of aerosol dosage forms that can be easily opened and closed and delivers the content in the desired **form and amount**.
- The primary purpose is to regulate flow from the container.
- **Types:**
- **Metered valves:** contain a chamber whose size determines the amount of the medication dispensed.
- **Continuous spray valve:** no metered chamber.



**Figure 33-7.** Continuous-spray aerosol valve, showing subcomponents used for sprays, foams, and semisolids.



# Component of Aerosol Package

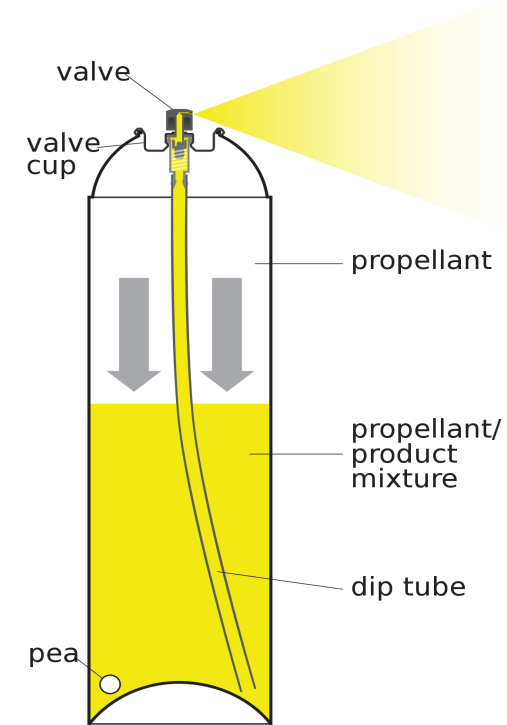
## 3. Actuators

- To ensure that aerosol product is delivered in the **proper and desired form**.
- Different types of actuators:
  1. **Spray**: mostly used for pharmaceutical applications. If the product contains a **low** amount of propellant, these sprays will deliver the product as a stream rather than a spray.
  2. **Foam**: consists of relatively **large** orifices and a large **chamber** that allows the product to **expand**.
  3. **Solid** stream: essentially similar to foam type and used for dispensing **semisolid** products such as ointment.
  4. **Special applications**: special design to deliver medication to the site of action such as throat, nose, or vaginal tract.



# Types of Aerosols Systems

- **First Type:** Solution system/**two**-phase system.
- **The simplest** and most widely used system that composed of **vapor** and **liquid phases**.
- In this system, the active ingredient is **soluble** in the propellant
- The liquid phase consists of a **solution** of **active ingredients** in liquid **propellant** or a mixture of liquid propellant and a solvent.
  - The solvent is **miscible** with the propellant.
- The amount of propellant may vary from 5% for foam to to 95% for inhalation products.
  - This amount will affect the vapor pressure of the system which is the main factor that will **determine** the size of the particle produced and the type of spray (fine spray, wet mist, or foam).



# Types of Aerosols Systems

- **Second Type: Water-based system (three-phase system):** propellant, water, vapor
- A relatively large amount of water can be used to replace all or part of the non-aqueous solvent used in aerosols.
- The active ingredient is dissolved in water and then formulated as an emulsion of water droplets in the propellant
  - Droplets of (Water+ drug) are the internal phase, and propellant is the external phase.
- Water is not miscible with liquefied gas propellants.
- Vaporized propellant will disperse the active ingredients into minute particles.
- This system is useful for topical pharmaceutical aerosols in that it allows greater use of liquid components not miscible with the propellants.
- Depending on the formulation, the product will be emitted as a spray or foam.



# Types of Aerosols Systems

- **Third Type:** Suspension or dispersion system
- This system helps to overcome problems resulting from the use of **cosolvent**.
- This involves the **dispersion of the active ingredient in the propellant**.
  - **Surfactant** may be added to decrease the settling of the particle.
- These systems are developed primarily for use for **oral inhalations**.
- The **physical** stability of an aerosol dispersion can be **increased** by
  1. Control of **moisture** content (moisture will increase drug solubility → and cause particle growth).
  2. Use of derivatives of active ingredients that have **minimum solubility** in the propellant (but must have the same pharmacological activity).
  3. Reduction of the initial **particle size**.
  4. Adjustment of the **density** of propellant and/or suspension so they are equalized.
  5. Use of **dispersing** agents.

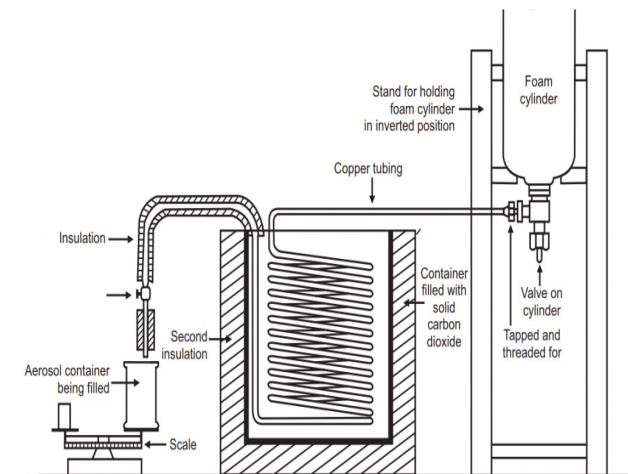
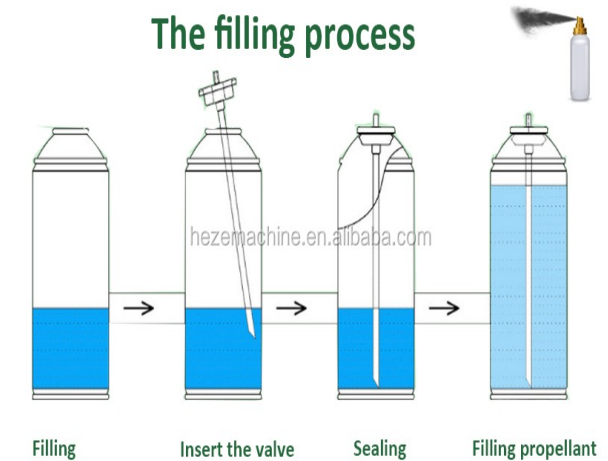
# Types of Aerosols Systems: Foam System

- **Fourth Type:** Foam aerosols consist of a **three-phase** system in which the liquid propellant, which normally does not exceed 10 to 15% by weight, is emulsified (internal phase) with the drug-containing liquid.
- So we have propellant (**internal phase**), and drug dissolved in a solvent (**external phase**)
- When the valve is depressed, the emulsion is forced through the nozzle, and in the presence of warm air and at atmospheric pressure, the entrapped propellant reverts to a vapor and whips the emulsion into a foam. → the product is emitted as a liquid and then converted to foam



# Manufacturing of Pharmaceutical Aerosols: Filling

- **Pressure filling apparatus**: propellant (**liquified gas**) is pressurized into the aerosol container. The container already **contains the product** concentrate and the valve attached **before** the filling process. This is carried out at **room temp**.
- **Cold filling apparatus**: simple and fast filling method which includes **cooling** the content to about  $-30^{\circ}\text{F}$ .
  - The cooled contents (drug and propellant) are added to a cooled container and **then** a valve is **placed**.
  - However, this is **suitable only** for **non-aqueous** content because aqueous contents will **freeze** at that temperature.



Cold filling

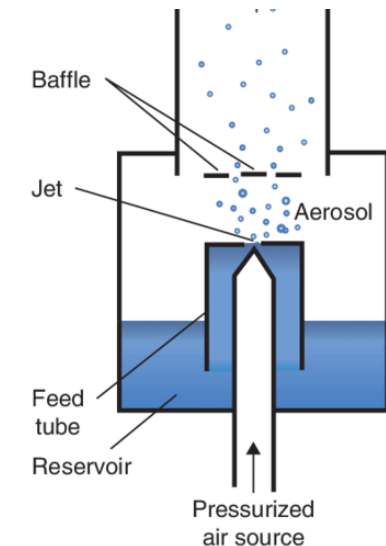
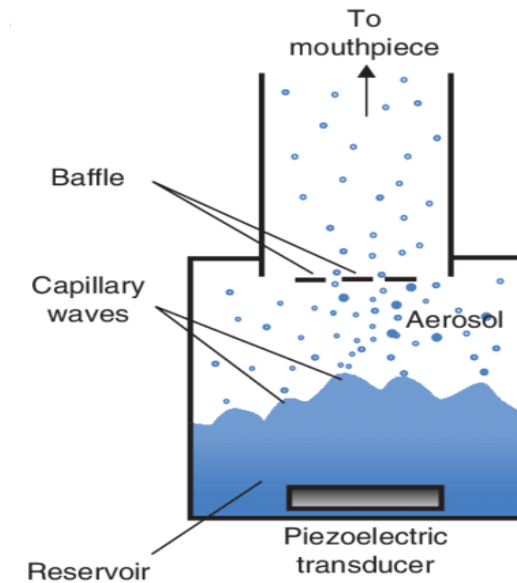
# Quality Control of Pharmaceutical Aerosols

- It includes the testing of :
  1. **Propellant:** need to check specifications like vapor pressure and purity.
  2. **Valves, Actuator, and Dip tube:** test solutions of known components are prepared and filled in the aerosol package to be tested. Then product is actuated and the **delivered dose** is weighed and compared with others.
  3. **Containers:** sample containers are collected and examined for defects.
  4. **Weight checking:** The weight of filled containers is periodically checked.
  5. **Leak testing:** passing full container into a water bath heated to 130°F.
  6. **Spray testing:** is done for **all** aerosols to ensure the **dip tube** is empty of pure concentrate or pure propellant. This will ensure that the patient will get the required dose the first time.



# Aerosols for Pulmonary Drug Delivery

- **Nebulizers**: dosage form for dispensing a **non-pressurized** liquid formulation.
- They are either electric or pneumatic
  1. **The electric (ultrasonic)** type produces **high-frequency sound waves** in the liquid which will form small droplets at the surface and these droplets are moved by **inhalation**.
  2. **Hydrodynamic nebulizers (jet Neb.)**: compressed air will generate a droplet that is delivered to patients.



# Advantages and Disadvantages

- **Advantages:**
  - They can deliver **larger volumes** that cannot be delivered using the metered dose inhaler.
  - Also, **no** high oropharyngeal impaction which will lower the side effects.
- **Disadvantage:**
  - Nebulizers are **bulky** and require a fixed **power source**.
- However, the newer design allows for the manufacturing of a portable nebulizer such as Respimat®



# Aerosols for Pulmonary Drug Delivery

- **Metered dose inhaler:** a **pressurized** formulation in which the **valve contains a metered chamber** to deliver the right dose each time. The preparation is usually a suspension.
- **Advantage;**
  1. Delivers a relatively **accurate amount** of drug each time.
  2. Easy to handle multidose and can be operated by patients.
- **Disadvantage:**
  1. Only a **fraction** of the dose reaches the lung because the **velocity** of the impact makes the particle stick to the mouthpiece and oropharynx.
  2. Requires some **patient education** for proper use that the patient should inhale while pressing the actuator which may make it a little difficult to use.
- **Note:** newer modifications such as a **spacer** reduced this problem.



**Spacer**

# Aerosols for Pulmonary Drug Delivery

- **Dry Powder Inhaler:**
- They **depend** on **inhalation** to draw the drug from the inhaler to the lung.
- The micronized drug is mixed with a carrier.





# Aerosols for Intranasal Drug Delivery

- Intranasal route offer several **advantages** include the relatively **rapid onset**, **easy** to use, and avoid **first pass** metabolism.
- Product are available for local effects such as Beconase<sup>®</sup> or for systemic effect such as desmopressin.

