

Chapter 7 Mixing and Granulation

7.1 Mixing and Segregation (Chapter 9)

Mixing vs. segregation

(1) Types of Mixture

* *Perfect mixing*

Random mixing

Segregating mixing

Figure 9.1

(2) Segregation

1) Causes and Consequences of Segregation

- *Particles with the same physical property (size, density and shape) collect together in one part of the mixture.*
- *Usually it occurs during moving, pouring, conveying, processing*
- *Its degree depends on particle-particle interaction**

* *Free-flowing powder or coarse particles → segregating rather than mixing*
Cohesive powder or fine particles → mixing rather than segregating but easily aggregating

2) Mechanisms of Separation *Figure 9-2*

- *Trajectory segregation*

From Chapter 3 in lecture note,

$$\text{Stop distance } s = \frac{\rho x^2 U}{18}$$

- *Percolation of fine particles - Figure 9.3*

Rise of coarse particles on vibration - Figure 9.4

- *Elutriation segregation*

3) Reduction of Segregation

- Make the sizes of the components as close as possible
- Reduce the absolute size of the particles

$$< 30 \mu\text{m with density about } \rho_p = 2000-3000\text{kg/m}^3$$

Critical diameter lowered as the density increases.

- Use of interparticulate forces

Add a small amount of liquid (Use of liquid-bridge force)

- Make one of the components very fine (less than 5 μm)

Ordered mixing*

Figure 9.5

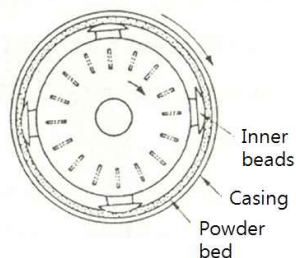
- Avoid to promote the segregation
- Use continuous mixing for very segregating materials

4) Equipment for Particulate Mixing

- Mechanisms of Mixing and Types of Mixer (9.5.1 and 9.5.2)

Diffusive mixing	<ul style="list-style-type: none"> - Random walk phenomenon - Essential for microscopic homogenization - Not suitable for segregating particles 	Tumbling mixers, Figure 9.6
Shear mixing	<ul style="list-style-type: none"> - Induced by the momentum exchange of powders having different velocities - Semi-microscopic mixing 	High-velocity rotating blade Low velocity-high compression rollers,
Convective mixing	<ul style="list-style-type: none"> - Circulation of powders by rotating blades - Beneficial for batch mode, not for continuous mixing - Suitable for segregating particles 	Ribbon blender, Figures 9.7,9.8 Fluidized-bed mixer

* Ordered mixture by dry impact blending method



5) Assessing the Mixture

For Binary mixture(2 components)

If $y_i(i=1,2,\dots,N)$: composition of the key component in the i -th sample,

Sample mean

$$\bar{y} = \frac{1}{N} \sum_{i=1}^N y_i$$

* True mean?

Standard deviation, σ (standard variance, σ^2)

- Estimated standard variance(S^2)

$$S^2 = \frac{1}{N} \sum_{i=1}^N (y_i - \bar{y})^2$$

- Theoretical Limits of variance

Upper limit: true standard deviation for a completely unmixed system, σ_0

$$\sigma_0^2 = p(1-p)$$

Lower limit: true standard deviation of random binary mixture, σ_R

$$\sigma_R^2 = \frac{p(1-p)}{n}$$

where $p, 1-p$: fractions of two components in the whole mixture

Degree of Mixing (Mixing indices)

The ratio of mixing achieved to mixing possible

$$\text{Lacey} : \frac{\sigma_0^2 - \sigma^2}{\sigma_0^2 - \sigma_R^2}$$

$$\text{Poole} : \frac{\sigma - \sigma_0}{\sigma - \sigma_R}$$

Worked Example 9.1, 9.2, 9.3

7.2 Size Enlargement - Granulation (Chapter 11)

* Size enlargement - agglomeration of particles

cf. coagulation

* *Why enlarge the particles?*

- *To reduce dust hazard*
- *To reduce cake and lump formation*
- *To increase flow properties*
- *To increase bulk density for storage*
- *To increase nonsegregating mixtures*
- *To provide defined metered quantity of active ingredients*
- *To control surface-to-volume ratio*

* *How enlarge the particles?*

- *Granulation: agglomeration by agitation (relative motion of particles)*
- *Machine granulation : compaction(tabletting), extrusion*
- *Sintering: thermal, final densification*
- *Spray drying: starting from droplets followed by its drying*
- *Prilling (freeze drying)*

(1) Interparticle Forces (11.2)

1) Van der Waals Forces

- *Between two spheres*

$$W = - \frac{A}{12z} \frac{x_1 x_2}{x_1 + x_2}$$

where A : Hamaker constant

z : separation

2) Forces due to Adsorbed Liquid Layers

- *Overlapping of adsorbed layers*
- *Dependent on area of contact and tensile strength of the adsorbed layers*

3) Forces due to Liquid Bridges

For pendular state Figure 11.1

$$F = 2 r_2 + r_2^2 \left[\frac{1}{r_1} - \frac{1}{r_2} \right]$$

- * Strong granules in which the quantity of liquid is not critical...
- * Granule strength continuously decreases in funicular, capillary and droplet states.

4) Electrostatic Forces

- * Contact electrification:
 - Friction caused by interparticle collision → Transfer of electrons between bodies

5) Solid bridges

- Crystalline bridges
- Liquid binder bridges
- Solid binder bridges

6) Comparison and Interaction between Forces

- Humidity vs. van der Waals forces, interparticle friction, liquid bridges and electrostatic forces

Figure 11.2 Tensile strength for various bonding mechanisms

(2) Granulation (11.3)

- Agitation: distribute liquid binder and impart energy to particles and granules for relative motion to meet together...

1) Granulation Rate Process (11.3.2)

- i) Wetting
 - Rate of penetration of liquid

$$\frac{dz}{dt} = \frac{R_p \cos \theta}{4 \mu z}$$

Washburn equation

where R_p : average pore radius, depending on particle size and packing density \Rightarrow packing..

θ : dynamic contact angle

μ : viscosity of liquid, depending on the binder concentration

ii) Growth *Figure 11.3*

- Nucleation - shatter
- Coalescence - breakage
- Layering - attrition
- Abrasive transfer

Define $Stk \equiv \frac{\rho_g V_{app}^2}{16 \mu}$ *Box on p274 Ennis and Litster(1997)*

$$Stk^* = \left(1 + \frac{1}{e}\right) \ln\left(\frac{h}{h_a}\right)$$

where e : coefficient of restitution

h_a : surface roughness of granules

- Noninertial regime: $Stk < Stk^*$
 - all collisions effective for coalescence
 - rate of wetting controls
 - independent of liquid viscosity, granule size and kinetic energy of collision
- Inertial regime: some Stk exceeds Stk^*
 - the proportion of successful collision decreases
 - dependent on viscosity, granule size and kinetic energy
- Coating regime: average Stk exceeds Stk^*
 - granule growth is balanced by breakage
 - growth continues by coating of primary particles onto existing granules

iii) Granule consolidation

- increase in granule density by closer packing density
- squeeze out liquid

2) Simulation of Granule Growth (11.3.3)

Rate of increase of number of granules in size interval v to $v+dv$	=	Rate of inflow of granules in size interval v to $v+dv$	-	Rate of outflow of granules in size interval v to $v+dv$	+	Rate at which granules enter size range v to $v+dv$ by growth	-	Rate at which granules leave size range v to $v+dv$ by breakage
(11.7)		(11.8)				(11.9) + (11.10) + (11.11)		

$$\begin{aligned}
 \frac{\partial n(v,t)}{\partial t} = & \frac{Q_{in}}{V} n_{in}(v) - \frac{Q_{out}}{V} n_{out}(v) + \frac{\partial G(v)n(v,t)}{\partial v} + B_{nuc}(v) \\
 & + \frac{1}{2} \int_0^v \beta(u, v-u, t) n(u, t) n(v-u, t) du - \int_0^\infty \beta(u, v, t) n(u, t) n(v, t) du \\
 & - \frac{\partial A(v)n(v,t)}{\partial v}
 \end{aligned}$$

3) Granulation Equipments (11.3.4) Table 11.1

- *Tumbling granulator* Figure 11.4
 - *Tumbling inclined drum and pan*
 - *Operate in continuous mode*

- *Mixer granulator*
 - *Rotating agitator*
 - *From 50 rpm (horizontal pug mixer-fertilizer) to 3000 rpm (vertical Schugi high shear continuous granulator-detergent, agricultural chemicals)*

- *Fluidized bed granulators*

- *Bubbling or spouted bed Figure 11.5*
- *Operate in batch or continuous mode*
- *Good heat and mass transfer*
- *Mechanical simplicity*
- *Combine drying stage with granulation*
- *Produce small granules*
- *Running cost and attrition rates : higher*