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Mixing dynamics for easy flowing powders in a lab scale Turbula[®] mixer

C. Mayer-Laigle, C. Gatumel, H. Berthiaux*

Centre RAPSODEE, UMR CNRS 5302, Ecole des Mines d'Albi-Carmaux, France

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A B S T R A C T

Powder mixing is one of the most common unit operations in industries handling powders. However, due to a lack of scientific knowledge on the behavior of powders inside a mixer, the optimization of a powder mixing step typically involves an experimental lab work in order to be transposed to a larger scale. To simplify this huge work, improving our understanding of the mixing dynamics remains some of the main industrial issues.

In this work, the mixing dynamics of an easy flowing powder made of lactose and couscous have been studied in the Turbula[®] mixer T2F. According to process parameters, we focused on the definition of mixing modes and we sought to identify the main mixing and segregation mechanisms at play for each mode, based on the analysis of mixing kinetics and autocorrelation functions. This work is a first step toward the definition of scale-up laws. The ultimate aim is to build predictive models to determine process parameters for obtaining the desired quality of mixing according to powder characteristics, flow properties and industrial constraints.

1. Introduction

Powder mixing is a very old and common operation performed instinctively by humans since the dawn of time with as first tools hands, poles, spoons and shovels. The need for mixing growing volumes and the formulation of more complex powders for industrial applications led to the design of numerous mixing devices in order to achieve homogeneity at specific scales. (Bridgwater, 2010). Many process parameters and powder properties influence the quality of a mixture, such as the shape of the vessel, the nature, the speed and the motion of the driving system, the mixing time, the particle size distribution of the powder and its flowability. . . , etc. However, the complexity of granular materials makes the description of the system powder/mixer extremely difficult, especially if it has to be done into all its details, under a mathematical form and for a reasonable time, (Ammarcha et al., 2013). Today, the

choice of an industrial mixing device and its optimization is still essentially based on an experimental work that can be tedious.

To help solving this task and a science, Chemical Engineering has to develop dynamic models to describe the mixing and segregation mechanisms based on powders properties, in particular their flowability, and the accumulated knowledge on the different powder mixing devices for powders of similar characteristics. These models should provide a better understanding of the behavior of the powder inside the device and thereby reduce the number of experiments for the determination of mixing parameters.

This is the aim of this paper. Beginning from an experimental study of the behavior of an easy flowing mixture in a Turbula[®] mixer, we propose a map of mixing modes related to mixing and segregation mechanisms according to mixer's operational parameters.

* Corresponding author. Tel.: +33 563493144.

E-mail address: berthiau@enstimac.fr (H. Berthiaux).

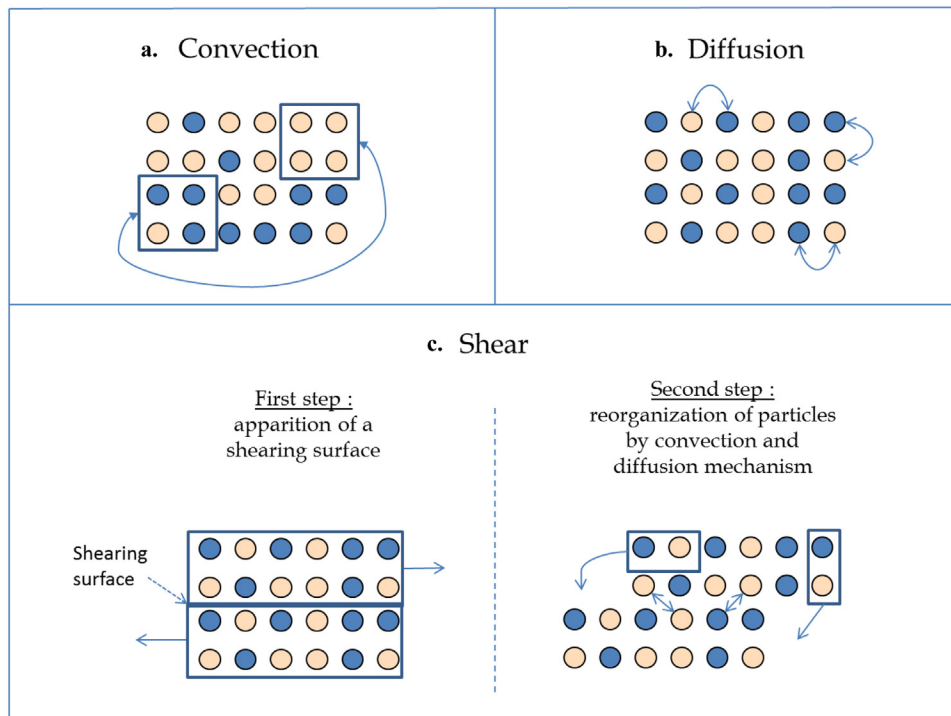


Fig. 1 – Convection, diffusion and shear mixing mechanisms.

2. Overview on powder mixing dynamics in scientific literacy

2.1. Mixing and segregation mechanisms

Mixing powders first means to set in motion the different particles inside the mixer. The external supply of energy needed is either provided by the motion of a vessel, by a blade passing into the bulk, by gravity or by a combination of some of these three ways. Depending on their own chemical and mechanical properties, powders can give very different answers to these solicitations. They can mix, which is generally the goal, or conversely tend to group together. Many mixing and segregation mechanisms have been described. However it is generally agreed that there are three main mixing mechanisms (Lacey, 1954):

- (i) Mixing by convection (Fig. 1a) corresponding to the motion of clumps of particles, generally under the action of a blade. The spatial homogeneity is quickly improved and the surface area between the components increases.
- (ii) Mixing by diffusion (Fig. 1b) which concerns rather the rearrangement of one or a small number of particles. It can occur at the frontier of the segregated zones when a blade is moving into the bulk or by particles/particles or particles/walls impacts. It is a slow process which promotes mixing on the length scale of the particles.
- (iii) Mixing by shear (Fig. 1c) is an intermediate mechanism which occurs when there exists a high velocity gradient between two clumps of particles. A shearing surface appears and leads to the reorganization of particles. Shear mechanism is often difficult to dissociate from convection inside a mixer. (Venables and Wells, 2001).

More than 19 segregation mechanisms have been identified by Mc Carthy (McCarthy, 2009) but the main mechanisms that typically occur during a mixing step are: segregation by

trajectory, segregation by percolation, segregation by elutriation (Williams, 1976), free surface segregation:

- (i) Segregation by trajectory can be observed when particles, having differences in size or density, are cascading inside a rotating drum or when they are projected by a blade: the heavier ones are projected farther than the others.
- (ii) Segregation by elutriation appears when particles are falling. The finer ones can be entrained with the air stream running through the bed of particles.
- (iii) Segregation by percolation is somewhat the opposite of segregation by elutriation. In vibrating conditions due to the motion of the mixer, smaller particles fall in the free spaces left by the coarser one
- (iv) Free surface segregation takes place when particles flow down on an inclined surface (when heap pouring for example). Particles flow along free surfaces all the more far as they are heavy or coarse. This mechanism is often the cause of radial segregation inside a rotating drum.

During a mixing step, the mixture quality is constantly changing through the action of the co-existing mixing and segregation mechanisms. Consequently, the balance between the co-existing mechanisms of mixing and segregation depends on the characteristics of particles, the shape of the vessel and the way energy is brought to the bulk.

2.2. How to evaluate powder mixing dynamics?

Mixing dynamics is related to the flow of particles as a result of external energy supply. Today, the understanding of mixing dynamics is a key point for the design and control of particle mixing processes as evidenced by the increasing number of studies published in the last 5 years (see: Ammarcha et al., 2013; Gosselin et al., 2008; Koller et al., 2011; Aït Aïssa et al., 2010; Fullard et al., 2013; Geng et al., 2013; Vanarase et al., 2013; Sahni et al., 2011; Tjakra et al., 2013).

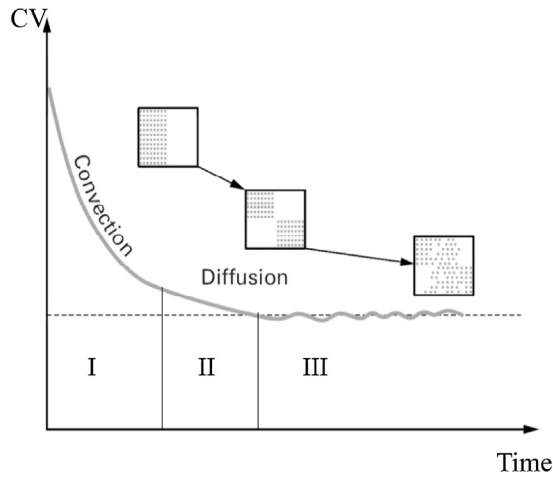


Fig. 2 – Typical mixing kinetics (Rose, 1959).

A recent study (An-Ni et al., 2013) reviewed the different experimental and numerical tools that could be used to investigate the mixing of particulate systems. This work shows the technical advances about non-intrusive methods and numerical simulations to assess mixture quality and understand mixing dynamics in a specific mixer. However these methods are difficult to apply to any industrial system. However by following the time-variation of mixture homogeneity, information about powder behavior inside the mixer can be deduced.

For assessing mixture homogeneity from samples composition measurement, two concepts are mainly used: the intensity and the scale of segregation (Danckwerts, 1953). The first one studies fluctuations among sample's compositions and is generally linked to the coefficient of variation (CV) (Eq. (1)). It qualifies the state of macromixing (Massol-Chaudeur et al., 2002a). A decrease of segregation intensity reflects the fact that the composition of each sample is close to the mean composition.

$$CV = \frac{\sqrt{\frac{1}{N} * \sum_{i=1}^N (x_i - \bar{x})^2}}{\bar{x}} \quad (1)$$

x_i and \bar{x} are the composition of sample i , and the mean composition in the key component respectively.

Mixing kinetic curves (see Fig. 2) are plotted by following the evolution of CV versus time. The mixing time is expressed as a dimensionless number to enable reliable comparison with various mixer sizes. (Delaplace et al., 2007).

A typical kinetic curve begins by a sort of quick decay linked to a rapid improvement of the quality of the mixture thanks to important large scale re-arrangements of the bulk induced

by convection and shear mechanisms. Then the CV decreases slowly as macroscopic motions become less efficient and only rearrangements at a finer scale, generated by the diffusion mechanisms, allow improving the mixture quality. Finally, on the last part of the curve a series of oscillations is likely to be observed reflecting a competition between mixing and segregation mechanisms (Massol-Chaudeur et al., 2002b).

The study of mixing kinetics is a key approach to compare different mixing conditions, choose the more appropriate mixer or optimize mixing conditions. Some authors are going further and try to understand, from the analysis of these data, the behavior of powder inside the blender (Massol-Chaudeur et al., 2002b; Marigo et al., 2011; O. Berntsson et al., 2002; Losada Martin et al., 2004).

The second tool suggested by Danckwerts for assessing mixture homogeneity, is the scale of segregation which describes the state of subdivision of clusters (Fig. 3). It can be seen as the maximum size of segregated regions. Its numerical value is determined by finding the value r_0 for which the autocorrelation function, $R(r)$ among composition x_i of N consecutive samples equals zero (r : distance between two samples).

$$R(r) = \frac{\sum_{i=1}^{N-r} (x_i - \bar{x})(x_{i+r} - \bar{x})}{\sum_{i=1}^N (x_i - \bar{x})^2} \quad (2)$$

Fig. 4 shows the associated autocorrelogram, for which ε corresponds to the limit below which $R(r)$ may be considered as equaling zero, from statistical thoughts (Chatfield, 2003) based on confidence intervals. For example, if a confidence interval of 95% is considered, $\varepsilon = 2/\sqrt{N}$.

In a certain way, r_0 can be seen as a characteristic length of the mixture. A low value of r_0 fits a mixing pattern arrangement at a fine scale, as obtained by diffusion mechanism. Conversely, a high value of r_0 means that there is a strong correlation between sample compositions and the mixture pattern is arranged at a larger scale, as obtained by convection mechanisms.

Although the scale of segregation provides interesting information on the mixture structure, examples of its use in assessing powder mixture homogeneity are scarce. We can mention the work by Schofield (1970) and, more recently, the one by Gyenis (1999). Both have linked the scale of segregation to the mixing mechanisms and particularly to convection through some simulations. They have shown that autocorrelograms bring suitable indices of mixing for predominantly convective mixing mechanism. However, when diffusion becomes the limiting phenomena, autocorrelation data are more difficult to interpret.

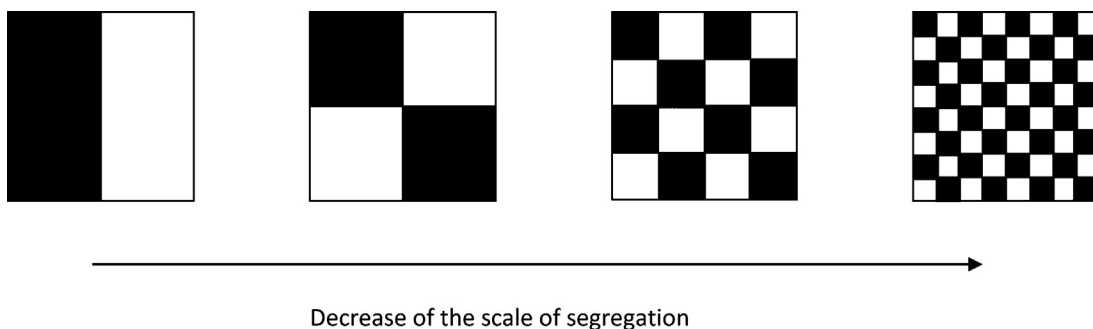


Fig. 3 – Schematic representation of the scale of segregation according to Schofield (1970).

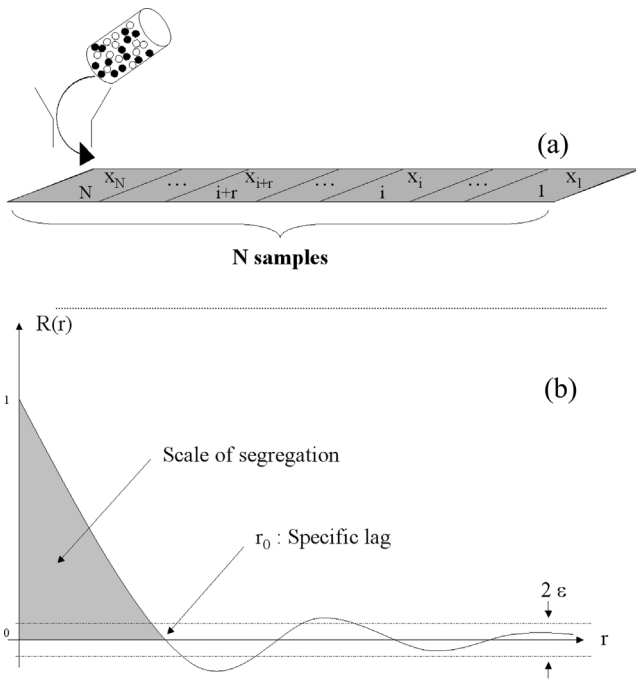


Fig. 4 – Discharge of a mixture onto a conveyor and division into n consecutive samples (a) as in Massol-Chaudeur et al. (2002a). Representation of the related autocorrelogram (b) the Turbula® mixer T 2F holding in the 2L mixing basket container.

In practice, the scale of segregation concept has been used to study static mixing using on-line image analysis (Muerza et al., 2002) but very few other papers have been published on the subject. Indeed, the determination of the scale of segregation (or the r_0 associated) requires exhaustive sampling. The collection of accurate data is somewhat a tedious task and authors (Kukukova et al., 2009; Hogg, 2003) concluded that this measure is difficult to be performed for a real case, and that such a tool cannot be used as a routine operation. Nowadays, thanks to on-line analytical techniques, the study of a larger number of samples in a short time looks now possible. Autocorrelation could therefore appear now as much viable tool for a better understanding of mixing process and mixture structure than it was a decade ago.

3. Equipment & methods

3.1. The Turbula® mixer

The Turbula® mixer has found a growing use both in industry and research since its release on the market at the end of the nineties. This device consists of a container which is placed in a mixing basket set by 2 stirrups at 2 rotary axes (see Fig. 4). One of them drives the whole system by rotating at an adjustable speed. The model T2F considered in this work has a mixing chamber that can hold a container of up to 2 L and the rotational speed can be varied from 22 to 96 rpm. The specific dimensions of the mixer are summarized in Table 1.

3.2. Powder mixture studied

We have studied the behavior of a blend composed of 85% in weight of lactose and 15% of couscous. Particle size distributions have been obtained by LASER diffraction in the dry mode (Malvern Mastersizer), Carr and Hausner indexes have been

Table 1 – Specific dimensions of the Turbula mixer T 2F.

Turbula® T 2F, Lab scale	
Volume of the container	2L
Radius of the stirrup	130 mm
Diameter of the vessel	130 mm
Length of the vessel	200 mm

Table 2 – particle size distributions and flow properties.

	Lactose	Couscous	Mixture of lactose/ couscous
Volume mean diameter	65.4 μm	≈ 1 mm	
Span ($D_{90} - D_{10}$)/ D_{50}	1.952	0.723	
Carr index	38.10%	7.3%	24.47%
Hausner index	1.61	1.08	1.32
Flowability	Bad	Very good	Normal

calculated from the bulk and tapped densities measured by an Erweka SVM 22 volumometer. These provide information on the bulk compressibility and the flow properties of the powder (Carr, 1970; Hausner, 1967). All these characteristics are summarized in Table 2.

Lactose is a cohesive powder whereas couscous is free flowing one. The flowability of the mixture is intermediate between that of each powder and can be qualified as “easy flowing” powder (Demeyre, 2007). In this context, due to a wide difference in particle sizes between each type of powder, it is expected that a segregation phenomena will take place during mixing.

3.3. Experimental protocol

Lactose and couscous powders (see Table 3) have been blended within the Turbula® mixer T2F in a 2 L stainless steel container with a 50% filling ratio.

Couscous and lactose were weighed directly into the mixing container to prevent possible losses during handling. Once the input speed of the engine axis has been chosen, the mixing operation is started for a defined mixing time. Different mixing times and rotary speeds have been explored in order to build mixing kinetics and autocorrelograms.

The quantitative characterization of mixture quality requires the prior set-up of a sampling procedure (number and size of samples, withdrawing method), as well as the determination of a reliable method of measurement which must be adapted to the nature of the powders, together with the intended use of the final product. In order to limit a possible perturbation of the sampling tools, the powder mixture has been discharged through a funnel on a conveyor belt. An inclined piece of stiff cardboard allows the formation of a powder layer of constant width and thickness. This method has been developed by Massol-Chaudeur et al. (2002a).

Couscous particles have been colored in black by iodine adsorption to increase the contrast between the two constituents, being lactose a white powder. This color difference is

Table 3 – Mixture compositions.

	Lactose	Couscous	Mixture
Mixing ratio	85%	15%	100%
Mass of constituents	357 g	63 g	420 g

easily detected by a current image processing technique. The measurement chain consists of a linear CCD camera placed over the conveyor belt which transports the loose materials, as well as of a computer (see Fig. 6) for which a specific Labview® program has been developed to determine the composition of the mixture. Each pixel has a value between 0 and 255, according to the gray level defined by the sensor. A binarization procedure is achieved by defining a threshold value and attributing the value 0 or 1 to the pixels whose gray level is below or above the threshold. This procedure allows differentiation between the products (lactose or couscous). A calibration curve links the number composition of black pixels to the weight composition of couscous in the sample.

The camera captures images of the powder mixture on a grayscale and the images are built from a series of 200 rows (see Fig. 7). The mixture is not spread over the entire width of the conveyor, so that the edges of the image are cut to retain only the part of it that contains powder. This portion of the image is called “region of interest”. The sample size roughly corresponds to 0.6 g of mixture. That way the whole mixture is sampled, calculating the autocorrelation function of the sample composition. The whole procedure is summarized in Fig. 5.

4. Simulation of Turbula® motion and definition of mixing regimes

Turbula® mixers have been used for numerous studies in scientific literacy and it would take too long to provide an exhaustive list. As examples, we can cite the work by Sundell-Bredenberg and Nyström (2001) which study the homogeneity of highly diluted micronized drug or those by Sommier (2000) and Sommier et al. (2001) who assess the mixture homogeneity for non-cohesive powders or cohesive powders differing in particle size (Obadele et al., 2012).

However, the behavior of powders during mixing in the Turbula® is still little understood. Recently, the powder motion inside this mixer has been investigated with the use of DEM simulation by Marigo et al. (2010, 2011, 2012). This modeling work focused on the simulation of granular mixing of simple particles (spheres) compared to experimental results obtained using Positron Emission Particle Tracking (Marigo et al., 2013). Similar trends are observed, especially in terms of bed transition behavior as a function of operating speed for mono-sized particles. However differences do exist between experimental and simulated values. In order to better fit the experimental results, these authors concluded by stressing the need to adjust coefficients in the simulation.

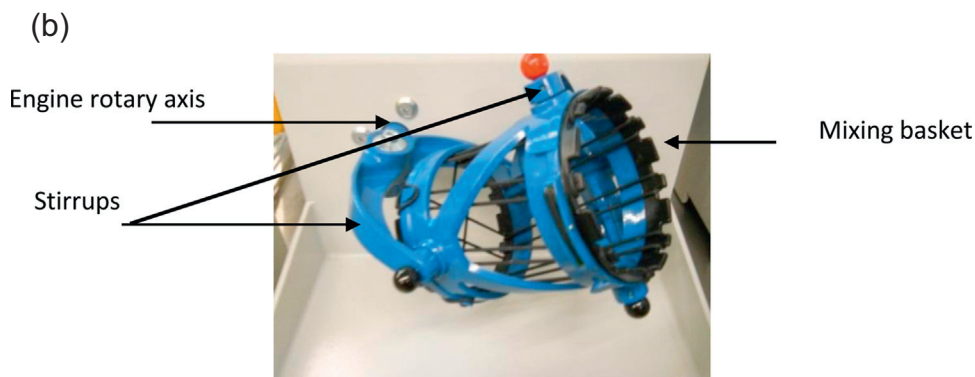
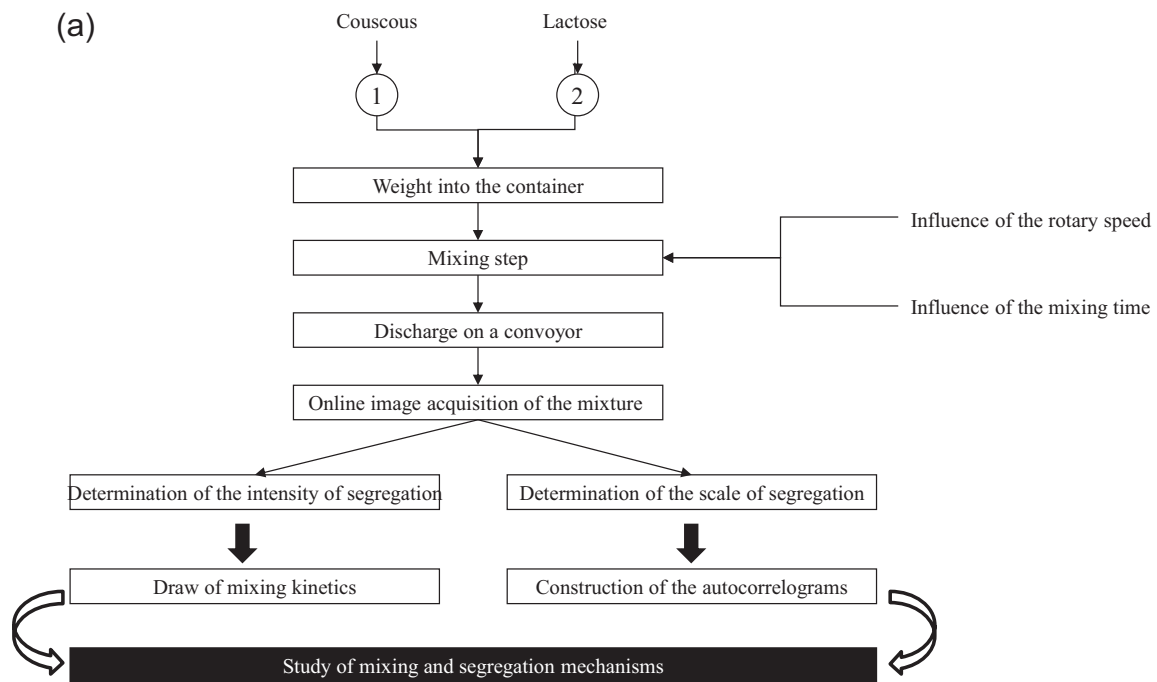


Fig. 5 – Experimental procedure and general methodology.

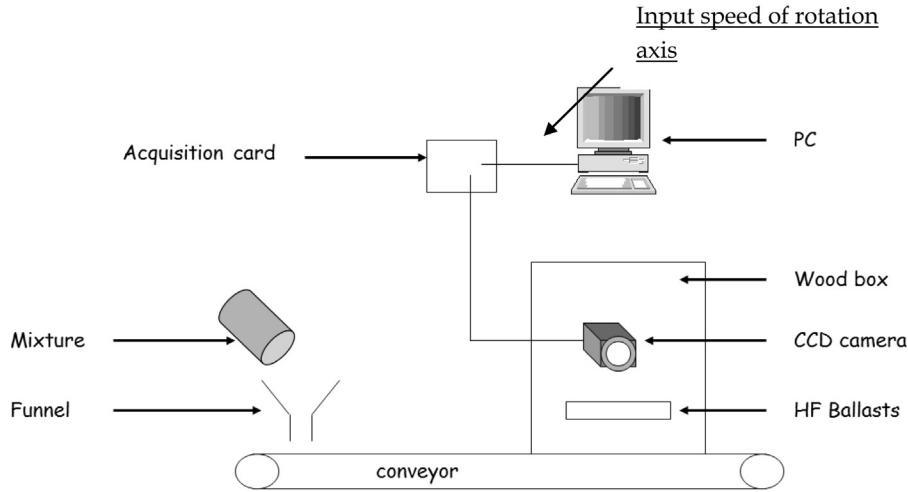


Fig. 6 – Experimental set-up for assessing mixture homogeneity.

In the present work, we choose to use a chemical engineering approach, seeking to describe the mixing process as a whole through dimensionless numbers. This approach, which is well known for processes involving fluids, has been sporadically used for granular materials. Indeed, on one hand it is hard to determine the impact of each factor influencing the flow of a given powder and on the other hand, these factors are difficult to quantify. He's work (He et al., 2007) is a good illustration of this daunting task, even with the use of a simple mixing device (rotating drum) and a model powder (particles of glass beads).

4.1. Turbula® motion

This mixer is known for its quickness to obtain homogeneous mixtures for a wide variety of granular systems. Its operation is based on the interaction of rotation, translation and inversion motions as per the geometric theory according to Schatz. The particle bulk moves across the vessel through a three-dimensional unsteady periodical behavior (Marigo et al., 2013).

The Turbula® has been modeled in the present work with the CAO soft Solidworks®. The mixer has been simplified and only the elements influencing the movement of the vessel has been designed (see Fig. 8). Each element has been built independently before being assembled by specifying the mechanical (pivot) link between them. Mixer motion has then been simulated and broken down (see Fig. 9) in the course of a period of revolution (T) which corresponds to a complete revolution of the engine axis.

In Fig. 9, the white arrows provide an indication of the direction of rotation of the vessel at the various stages of the movement. The main motion of the vessel is, of course, revolution. Inversions occur when the vessel turn over for the initial position, at $i = 0.3 * T$, $t = 0.51 * T$, $t = 0.87 * T$ and $t = 1 * T$. We

can also notice translation for t values around $0.18 * T$, $0.44 * T$, $0.67 * T$ and $0.95 * T$.

4.2. Definition of mixing regimes inside the mixer

It has been suggested to describe the powder mixing system in a simplified way using several dimensionless numbers as power number (N_p) and/or Froude number (Fr) (Sato et al., 1979; Entrop, 1978). For rotating drum, Mellmann (2001) defined different flow regimes (Fig. 9) according to the transversal motion of the powder. For example, he showed that rolling, cascading or cataracting promote mixing by convection for free flowing powder.

The transitions between flow regimes (featured on the diagram in Fig. 11) are based on the Froude number and the filling rate, calculated in rotating drum as follows:

$$Fr = \frac{\omega^2 * R}{g} \quad (3)$$

and

$$f = \frac{1}{\pi} * (\alpha - \sin \alpha \cos \alpha) \quad (4)$$

where α is the half angle of the section occupied by the powder bed (Fig. 12)

The determination of the Froude number from Eq. (3) is uneasy in the Turbula®, due to the non-circular mixing path (Kushner and Moore, 2010).

Relying on the definition of Froude Number, we sought for a suitable expression of Fr . Actually, Fr expresses a ratio between inertial force and G-force. Inside the container, inertial forces are mainly centrifugal ones and Fr could be written as in Eq. (5)

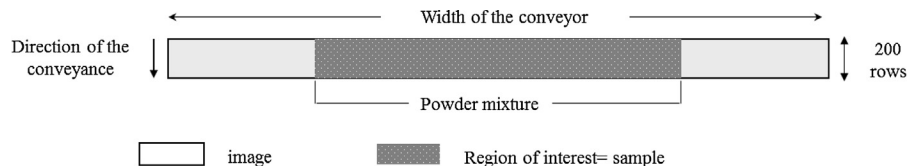


Fig. 7 – Defining the size of the image used for data processing.

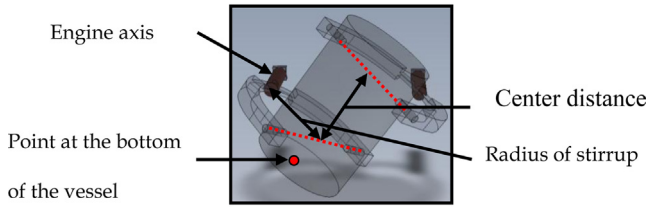


Fig. 8 – Schematic representation of a Turbula® mixer.

where m is the mass of an infinitesimal volume of powder, a its acceleration and g the acceleration of gravity.

$$Fr = \frac{\text{centrifugal forces}}{\text{gravity forces}} = \frac{m * a}{m * g} = \frac{a}{g} \quad (5)$$

Maximal accelerations for a reference point at the bottom of the tank has been extracted from Solidworks® simulations. We assume that this point sum-up velocities and accelerations in the powder bulk. Indeed, this corresponds to the bottom of the powder bed inside the container. During mixing, the powder is “reversed”, so that particles that were at the bottom of the bed will be at the top of it and vice versa. The values of Fr calculated from the different available input speeds of the engine axis are gathered in Table 4.

The calculated Froude number ranges from 0.5 to 4.5 and three flow regimes emerge depending upon centrifugal forces or G-forces are predominant.

Table 4 – Froude number in the Turbula mixer T 2F.

N: speed of the input motor axis (rpm)	Fr	Main force inside the mixer	Flow mode corresponding to Melmann's description
22	0.2		
32	0.5	G-forces	Cascading
46	1	Balance between G-forces and	Catacting
67	2.2	centrifugal forces	
96	4.5	Centrifugal forces	Collisional

We rank the Turbula® mixer as a rotating drum as far as powder bulk is only driven by the motion of the vessel, though this motion is somewhat different from that of classical horizontal drums. For a given Froude number, the forces responsible for the mixing pattern obtained are almost the same in the Turbula® or in a rotating drum, and we assume that particles could have similar overall movement (avalanche, cataract...) but of different amplitudes for the same flow mode.

According to the description by Melmann, and for a filling ratio of 50%, (see Figs. 10 and 11) we assume that:

- For a Froude number less than 1, particles are in a cascading flow regime.



Fig. 9 – Breakdown of the Turbula motion with Solidworks®.

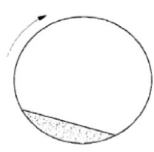
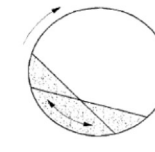

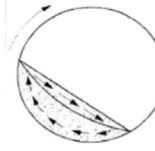

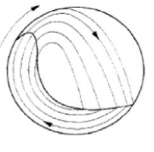
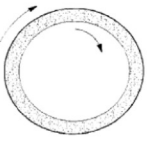
<u>Slipping</u>	 sliding	 surging	
<u>Avalanching</u>	 slumping	 rolling	 cascading
<u>Catacting</u>	 catacting	 centrifuging	

Fig. 10 – Flow mode in a rotary drum according to Mellmann (2001).

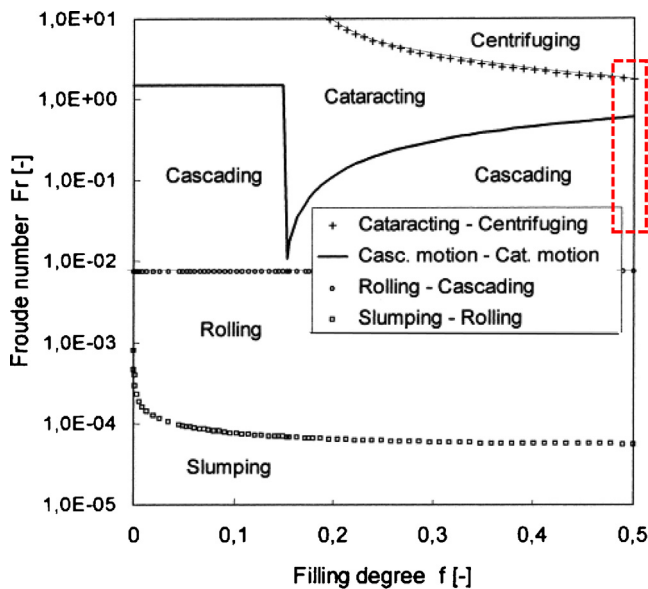


Fig. 11 – Transition between flow regimes in a rotary drum according to Mellmann (2001).

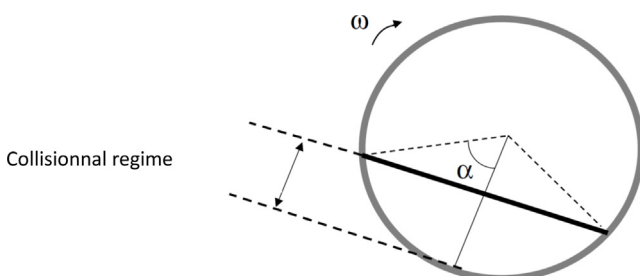


Fig. 12 – Half-angle of the section occupied by the powder bed in a rotating drum mixer.

- For a Froude number between 1 and 2.5, particles are in a catacting flow regime.
- For a Froude number of 4.5, flow is limited to catacting and centrifuging regimes, and can be qualified as collisional.

5. Results and discussion

Intensity and scale of segregation have been assessed over mixing time. Figs. 13 and 14 show the typical mixing kinetic curves, for which the intensity of segregation (coefficient of variation) is plotted against the number of revolutions of the mixer. The curves have been cut into two parts, one for the shorter mixing times (less than 16 revolutions) and one other for longer mixing times.

A reproducibility study has been carried out, being 3–5 measurements realized for each number of revolution. In the early stages of mixing, the relative variation on the CVs can rise up to 30%. As an example, for 3 revolutions at 32 rpm, the CVs vary between 3.63% and 4.77%. For longer mixing times, variations between the CVs are less important (below 7%). It seems that the random error decreases with the number of revolutions, leading to a reproducible pattern inside the mixture. The way the mixer is filled leads to a certain stack of particle. Depending on particle shape and their arrangement, the state of compaction of the powder bed can be different from one test to another. For shorter mixing times, the pattern obtained present important variations due to the rearrangement of the powder bed linked to the initial state of compaction. A high number of revolutions lead to the aeration of the powder and the rearrangement of the product is then more reproducible. Thus, the way in which the vessel has been filled has a weak impact on the pattern obtained for long mixing times.

However, these variations have in practice no impact for the comparison of the mean CV value as far as confidence intervals do not overlap each other between 3, 4 or 6 revolutions at the different speeds (22–96 rpm). For readability, only the average of the measurements has been represent on the graphs.

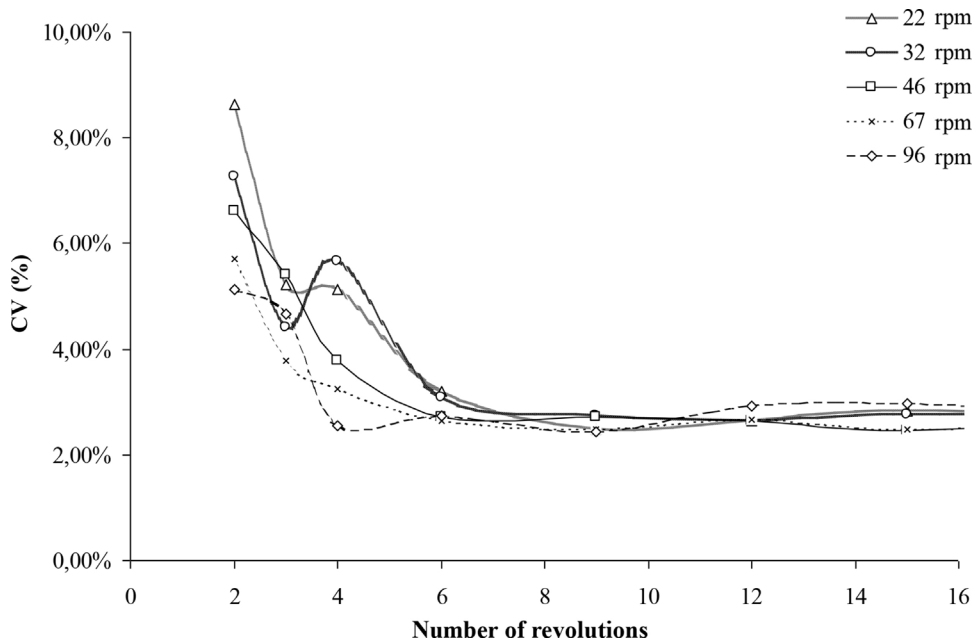


Fig. 13 – Mixing kinetics obtained in the Turbula® T 2F in the early stage of mixing at different rotational speeds.

Autocorrelation functions have also been plotted for 2, 6 and 500 revolutions at the different input speeds. To facilitate comparison and discussion, curves have been grouped according to the flow regimes defined in the previous section (see Figs. 15–17). The limit below which $R(r)$ can be considered to equal zero is featured by dashed lines on the graph. Numerical values of r_0 , the scale of segregation, have been determined for each case and are summarized in Table 5.

Table 5 – Values of the scale of segregation (r_0) obtained for the different input speeds at 2, 6 and 500 revolutions.

Input speed of the motor axis (rpm)	R_0		
	2 revolutions	6 revolutions	500 revolutions
22	169	282	ND ^a
32	251	62	40
46	116	110	ND ^a
67	59	58	ND ^a
96	53	48	257

^a ND: Not determined.

5.1. Discussion at the shorter mixing times: two revolutions

Each mixture is realized following the experimental protocol described in Section 2. When the boot order is given, the mixer accelerates to reach the selected speed. There is no braking system on the mixer, so that it continues to move for a certain time after the stop order. When the number of revolutions is

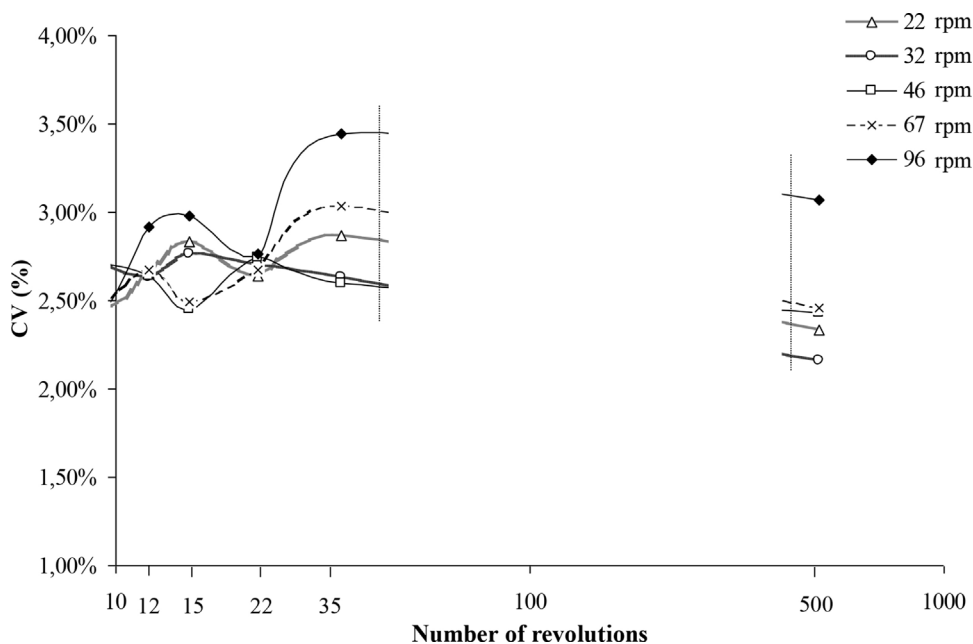


Fig. 14 – Mixing kinetics obtained for longer mixing times in the Turbula® T 2F.

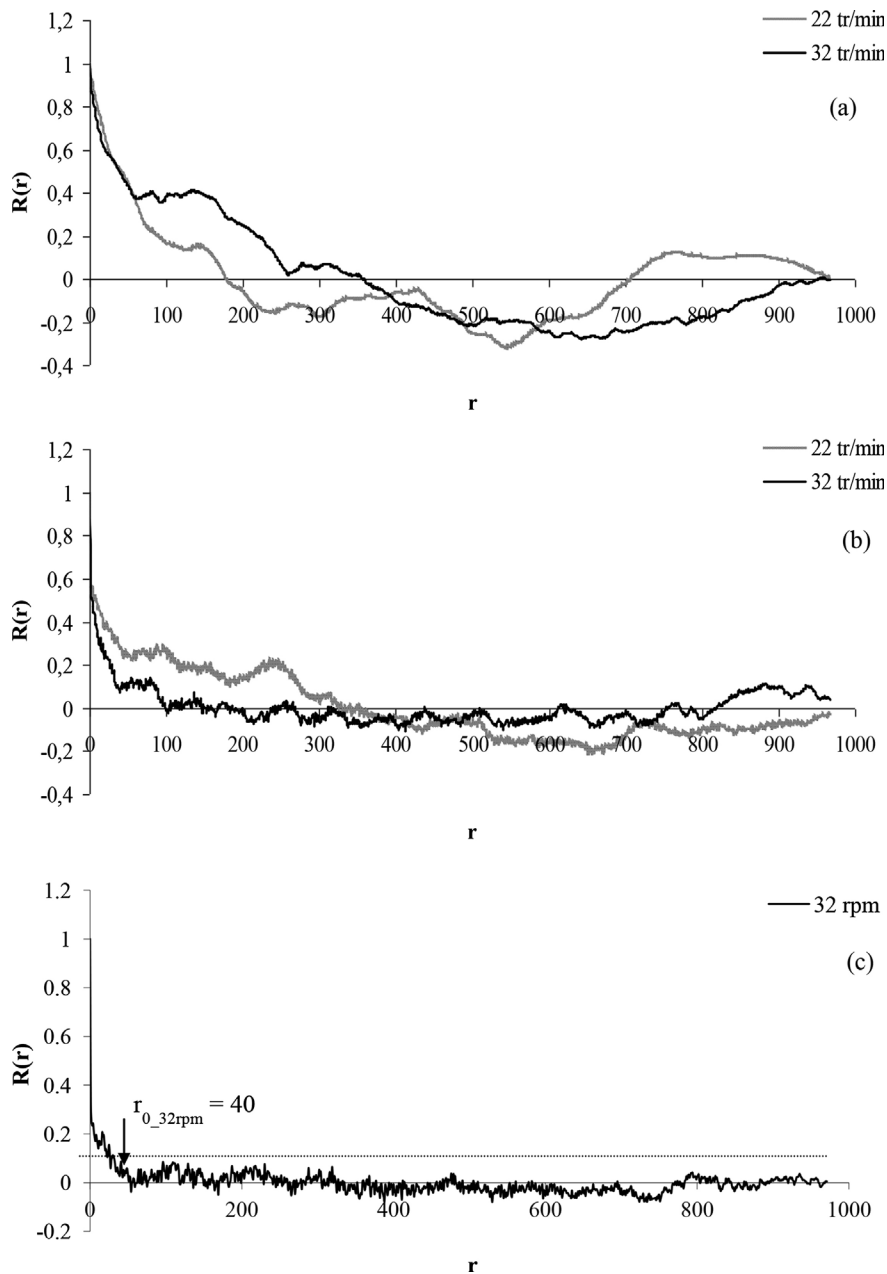


Fig. 15 – Autocorrelograms obtained at 2 (a) 6 (b) and 500 (c) revolutions for low input speed ($Fr < 1$).

small, the acceleration and deceleration periods will have a higher impact than for a longer number of revolutions. For example, at 96 rpm, the period during which the mixer would rotate at a constant speed simply does not exist for a 2 revolutions test.

In the mixing kinetics, the input engine speed has a strong influence on the value of the CV: the higher speeds lead to the lowest CV. This reflects a rapid improvement of the mixture quality at a macroscopic scale as it can be obtained by convection mechanisms. It seems that acceleration and deceleration in the first stage of the mixing enhances the convection mechanisms.

The autocorrelograms obtained for 2 revolutions (Figs. 15a, 16a and 17a) are quite similar for each mixing mode. Curves seem to be pseudo periodic with a succession of positive and negative autocorrelations. In the powder bed, this could be linked to an alternation of areas for which the concentrations of all the samples in the area are close but the

mean concentration of two neighbor areas are very different. For $Fr < 1$, the period of the autocorrelograms being large, we assume that the pattern looks more as represented in Fig. 17a with many large size areas in comparison with the autocorrelograms obtained for higher speeds (Figs. 15a and 16a), for which the mixing pattern is probably better illustrated by Fig. 17b and c.

The pseudo-period of each curve is almost equal to the scale of segregation, r_0 , and decreases as the input speed increases as it also leads to an improvement of the mixture quality at a microscopic scale.

For a Froude number less than 1, we assume that the powder flow inside the mixer is quite similar to the cascading regime in a rotating drum (see Section 4). This means that the powder sticks to the wall following the vessel in its revolution until it reaches an angle from which particles slightly take off and fall in avalanche on the “active layer”, where the mixing is achieved. On the top of the active layer, particles are in a

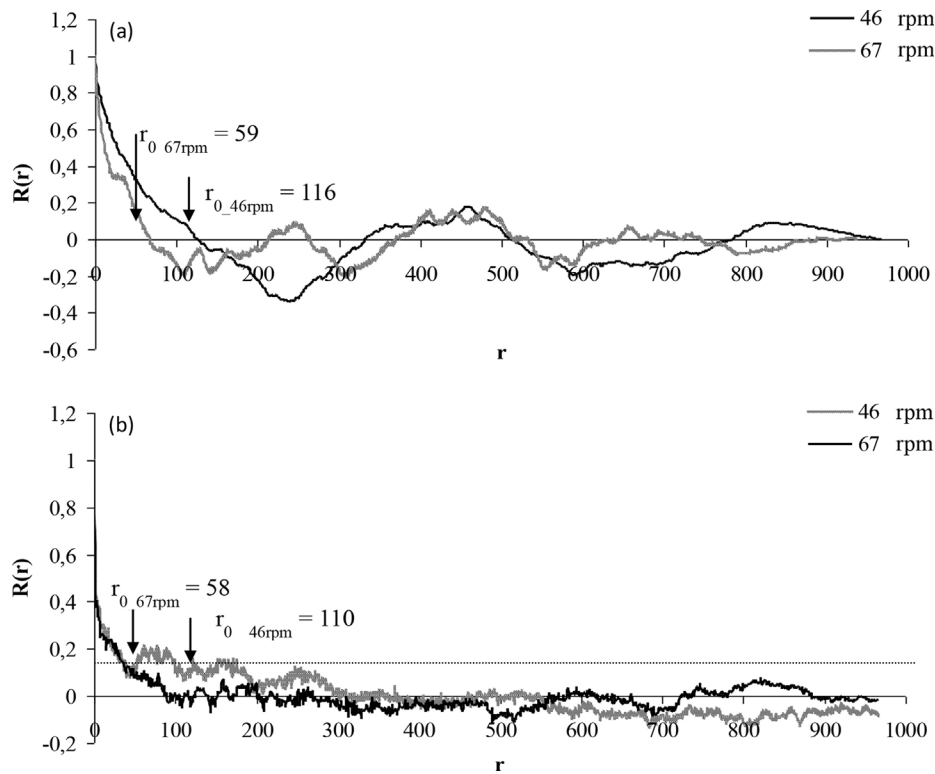


Fig. 16 – Autocorrelograms obtained at 2 (a) and 6 (b) revolutions for intermediate input speed ($1 < Fr < 2.5$).

rolling motion (passive layer). Particles fall in large amounts which could explain the large periods that are obtained for the autocorrelograms.

When the input speed increases (cataracting regime: $1 < Fr < 2.5$), some particles or small groups of particles are detached from the powder bed by centrifugal forces and are projected into the air which could explain the alternating of smaller areas of quite far compositions. In addition, the projection of particles also promotes the efficiency of diffusion mechanisms and explains a smaller scale of segregation.

For the highest speed ($Fr = 4.5$), the flow regime is almost collisional, with the generation of bed vibration and collision between the particles that are becoming more individualized. However, during the first revolutions of the mixer, acceleration and deceleration periods are predominant, and the existence of avalanche and cataract flows could explain the shape of the autocorrelogram, as well as a scale of segregation almost equal to that obtained at 67 rpm.

5.2. Discussion at intermediate mixing times: number of revolution between 2 and 35

In the cascading regimes ($Fr < 1$) and for input speeds of 22 and 32 rpm, the CV (Figs. 13 and 14) decreases more quickly than for higher speeds. It then slightly rises between 3 and 4 revolutions before decreasing again, drawing a “wave” on the curve. The total decrease for the CV between 2 and 6 revolutions is almost 50%.

At these speeds, mixing is most likely achieved thanks to successive powder avalanches. During the rotation of the vessel, particles can roll on the wall of the vessel or on the free surface formed by the powder bed. Because of the size difference between lactose and couscous particles, free surface segregation is likely to appear. This mechanism comes into competition with mixing by convection and creates the wave that is observed. For longer mixing time (up to 35 revolutions)

the CV decreases more slowly. As far as the scale of segregation decreases, shear and diffusion mechanisms are becoming more efficient for the realization of an intimate mixture.

In the cataracting regime ($1 < Fr < 2.5$), for input speeds $N = 46$ and 67 rpm, the above-mentioned “wave” is no longer observed and mixing kinetics are looking more regular. In this mode, particles are projected, so that trajectory segregation is likely to occur, being certainly enhanced by the reversal of the vessel. However this can be offset or exceeded by mixing

Table 6 – Mixing and segregation mechanisms taking place in a Turbula T2F for easy flowing mixtures, according to our observations.

Flow regime	Speed of the engine axis (rpm)	Assumption about the behavior of the mixture for shorter mixing time	Assumption about the behavior of the mixture for longer mixing time
$Fr < 0.5$ cascading	22	Competition between mixing by convection and free surface	Lower value of CV probably due to mixing by diffusion
$0.9 < Fr < 2.5$ cataracting	32	segregation	Balance between mixing by diffusion and segregation (free surface or trajectory)
	46	Stronger mixing by convection. Segregation by trajectory	
$Fr = 4.5$, collisional	67	Mixing by diffusion is probably leading	Segregation by percolation
	96		

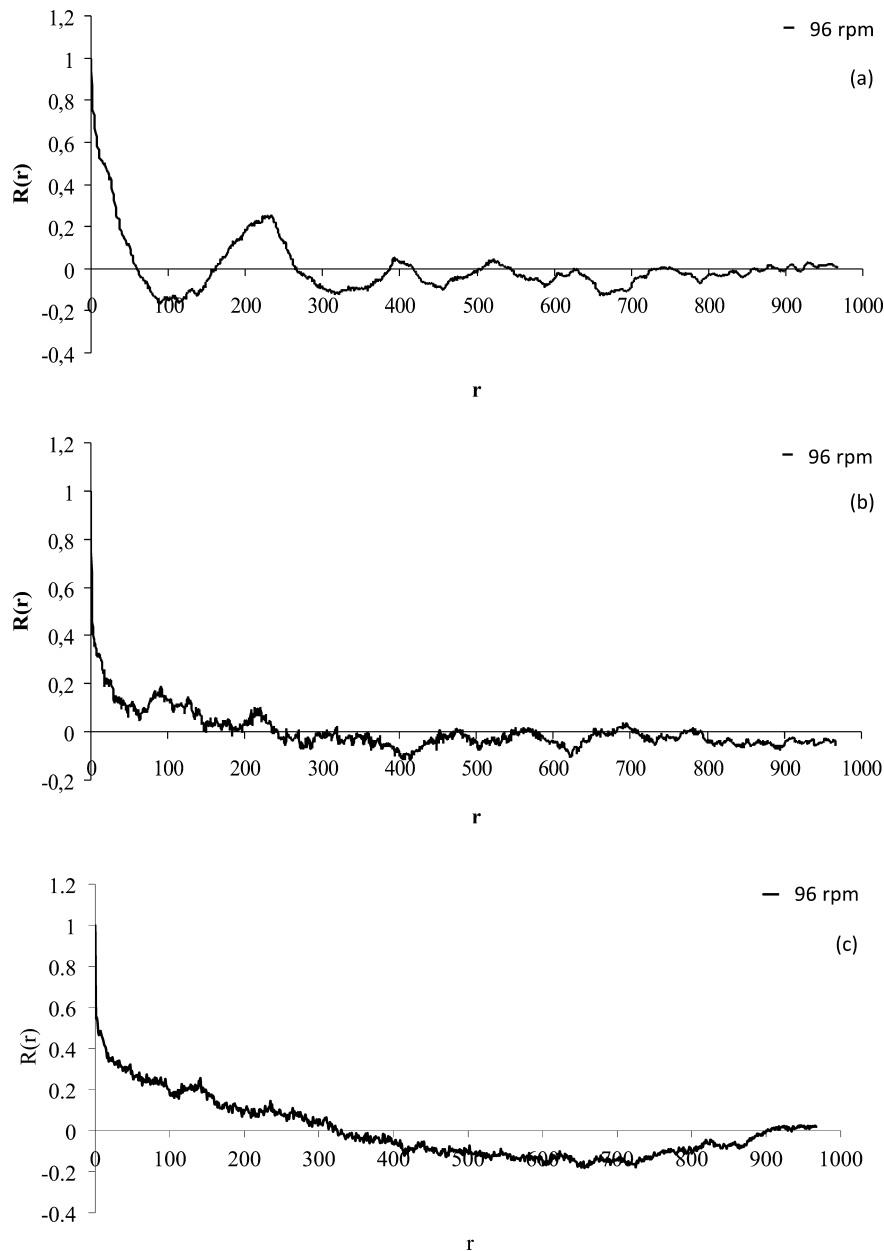


Fig. 17 – Autocorrelograms obtained at 2 (a), 6 (b) and 500 (c) revolutions for a high input speed: 96 rpm ($Fr = 4.5$).

by convection which is more important for this speed and the decrease of the CV is therefore monotone.

In the collisional mode ($Fr = 4.5$), the kinetics (Fig. 13) are showing a slight wave with a stagnant CV value between 3 and 4 revolutions. Inside a rotating drum, for very high speed corresponding to a collisional mode ($Fr = 5$, and filling rate = 50%), it has been observed that the mixture is experimenting a sort of vibration that drives to segregation by percolation. During the early stage of mixing, this comes into competition with mixing by convection, and then with mixing by diffusion for longer mixing times. In the powder system used in this work, the difference between lactose and couscous particle sizes enhances the effect of segregation by percolation, meaning that the resulting mixture is less intimate.

In Figs. 15–17, for all speeds except 22 rpm, the autocorrelogram is flattened. There is no succession of negative and positive correlations: the low contrast between sample's compositions is probably due to the influence of diffusion and shear mechanisms. However, the value of r_0 (Table 5) for the cataracting and collisional regimes did not change, while a

reduction of almost 50% of the CV has been observed in all the kinetics. Thus, it seems that diffusion and shear mechanisms do not allow improving the mixture quality at a microscopic scale because of a competition with segregation mechanisms (by trajectory or percolation)

5.3. Discussion at high mixing times (500 revolutions)

For 500 revolutions of the mixer (see also Fig. 13), the CV obtained at 22 and 32 rpm are the lowest values experienced. These observations are consistent with the work of Marigo et al. (2011) on DEM simulation of glass beads mixing in the same equipment. At 96 rpm, the value of the CV obtained is the highest one, and the mixture obtained is less intimate. The associated autocorrelogram (Fig. 17c) shows a slight positive correlation, then a slight negative correlation which does not exist at 6 revolutions (Fig. 17b). This reflects a certain order in the mixture that is probably linked to segregation by percolation, as discussed previously.

6. Conclusion

Through this work, we have been trying to understand the behavior of an easy flowing mixture constituted of 85% of lactose and 15% of couscous inside the Turbula® mixer T2F. Our analysis is based both on a kinematic simulation of the motion of the mixer and on an experimental work. The former indicates that 3 mixing modes based on the Froude number can be defined by analogies to what has been described in horizontally rotating drums. The experimental work showed that the Turbula® mixer allows a quick achievement of high quality mixtures, whatever the speed of the mixer.

From the study of mixing kinetics and autocorrelation functions, we identified the main mixing and segregation mechanisms that are summarized in Table 6, also depending on the input speeds.

Finally it must be noted that for an easy flowing powder, the cascading regimes seems to be the best to obtain intimate mixtures, which is in accordance to the data obtained by DEM simulation by Marigo et al. (2012).

In further developments, we plan to extend the study to cohesive powder systems. In addition, we would like to test the behavior of the same type of powder in Turbula® mixers of greater sizes (T10B and T50A), in order to define scale-up laws.

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