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Flow process conditions to control the void fraction of food foams in static mixers

M. Laporte ^{a,b}, C. Loisel ^b, D. Della Valle ^{b,d}, A. Riaublanc ^c, A. Montillet ^{a,*}

^a LUNAM Université, Université de Nantes, CNRS, GEPEA, UMR6144, BP 406, 44602 Saint-Nazaire, France

^b ONIRIS, BP 82225, 44322 Nantes, France

^c UR1268 Biopolymères, Interactions, Assemblages, INRA, 44316 Nantes, France

^d LUNAM Université, Université de Nantes, CNRS, LTN, UMR6607, BP 50609, 44306 Nantes, France

Previous studies have shown the feasibility of food foam production in a SMX10 static mixer. Nevertheless the void fraction, α , obtained was high (>0.85) compared to the target range in food industry (0.5–0.7). This work aims to reduce α by using a smaller hydraulic diameter of the static mixer (SMX+6) and higher liquid base viscosities. The hydrodynamic study exhibits slight differences in the pressure drop and process shear constants. The foams are processed with a liquid base of whey protein isolates (3%) and xanthan gum (0.35% or 0.6%). The boundaries between “slug flow” and “bubble flow” are determined for the two SMX and represented in a flow pattern map. Only the latter flow regime is suitable as α (0.73–0.96) can directly be controlled from the flow-rates. Working with a SMX+6 mm offers a substantial enlargement of the convenient velocity range that allows attaining lower void fractions compared to the SMX10.

1. Introduction

Food foams such as foams based on proteins are in constant development, as they are appreciated by the consumers and offer a wide range of possibilities to suit a variety of tastes, textures, or nutritional allegations (Campbell and Mougeot, 1999). In most cases for food foams, the foaming processes are carried out in batch in table top mixers. According to Campbell and Mougeot (1999), the void fraction of foams usually ranges from 0.5 for chocolate foams or ice creams to 0.7 for whipped creams (Chantilly). Yet, continuous processes bring significant benefits: high productivity, homogeneous shearing, small size of the equipment, energy saving and easy-to-monitor systems (Thakur et al., 2003a, Al Taweel and Chen, 1996; Al Taweel et al., 2007). However, continuous foaming is relatively poorly understood and the current knowledge is mostly empirical.

There are two types of continuous foaming processes. The first type is “dynamic mixers”, such as rotor–stator systems (Mondomix™, for one of the most widespread use in the industry), scraped-surface heat exchanger, or even the foaming column developed by Djelveh et al. (1998) and Souidi et al. (2012). The second one is “static mixers”, consisting of inserts in the flow section, namely “elements”, generating complex flow structure to create

the mixing. There are different shapes of static mixers but the cross designed ones, of SMX type, outperform the others, especially in the laminar regime (Meijer et al., 2012, Fradette et al., 2006). Static mixers are usually used for liquid–liquid extraction, liquid–gas absorption, heterogeneous chemical reactions, suspensions, emulsification or solid–liquid mixing (Paglianti, 2008). In the suitable foaming conditions, the main asset of static mixers is that the gas incorporated in the foam can directly be controlled from the flow-rates (Talansier, 2009), which is not the case in most of the other foaming processes. For example, the amount of gas incorporated with the Mondomix™ is only 30–80%, which makes it impossible to master the amount of overrun (Thakur et al., 2003b, 2005).

In a previous study displaying the opportunity to produce food foams with a static mixer, it has been shown that foams from egg white proteins can be produced in a SMX10 static mixer for void fractions (i.e. gas volume fractions) above 0.85, i.e. corresponding to dry foams (Talansier et al., 2013). However, as mentioned above, food foams are usually liquid foams with void fractions lower than 0.8. Actually, a key parameter for industrial foams is the stability, whereby smaller bubbles as well as higher void fractions improve the foam stability (Talansier et al., 2013). The challenge of overrun reduction (less than 0.85, compared with the previous runs in the SMX10 with egg white proteins), appears at first sight to be delicate, as it will affect the structural and textural characteristics of a foam with a negative effect on the stability. Literature reports that it is necessary to increase either the shear rates or/and the

* Corresponding author. Tel.: +33 (0)240 172 677; fax: +33 (0)240 172 618.
E-mail address: agnes.montillet@univ-nantes.fr (A. Montillet).

liquid viscosity to compensate for it (Hanselmann and Windhab, 1999; Mary, 2011; den Engelsen et al., 2002).

The determination of the conditions that are required to achieve the gas mixing in the main liquid flow, i.e., the flow pattern, is crucial in the industrial process control for both dynamic and static mixers. In a biphasic gas/liquid flow, the two phases distribute themselves into several distinct flow patterns depending upon their flow-rates, properties and pipe geometry (Kleinstreuer, 2003). These are referred to as flow regimes, mainly the annular, slug/plug and bubble regimes that are recognized by visual inspection. There have been several experimental works on the gas-liquid flow patterns in horizontal pipes (Baker, 1954. Mandhane et al., 1974; Wong and Yau, 1997). However, the frontiers between the different regimes cannot be expressed by the same dimensionless numbers, despite the different attempts to establish universal maps (Schicht, 1969; Taitel and Dukler, 1976; Barnea et al., 1980; Spedding and Chen, 1981; Lin and Hanratty, 1987). Different studies oriented toward fluid properties have been published; nevertheless, very few investigations were conducted on the effect of the fluid viscosity in the case of flow through horizontal pipes. Studies concerning high fluids viscosity came up with contradictory conclusions: Matsubara and Naito (2011) stated that the regimes are affected by the viscosity, whereas Weisman et al. (1979) reported only minor changes induced by this parameter. A decrease in the surface tension shifts the frontiers to lower gas velocity (Spedding and Hand, 1997. Wilkens et al., 2006) but this effect vanishes at high gas velocities (Tzotzi et al., 2011)). Therefore, additional work is required for the understanding of the flow pattern maps, especially in a complex geometry such as the SMX, generating elongational effects that are not present in empty plain tubes (Bone, 2005).

In this study, we propose to investigate the effect of the flow conditions and of the liquid base properties in two SMX static mixers characterized by different hydraulic diameters. The experiments are designed to determine the flow process conditions in SMX static mixers enabling formation of foams at lower void fractions. Firstly, a hydrodynamic study is performed through the determination of the respective characteristic constants: K_p , the power constant, and K_s , the shear rate constant evaluated by the Metzner-Otto method (Metzner and Otto, 1957). The hydrodynamic conditions allowing complete and homogeneous gas incorporation and corresponding to the bubble regime are experimentally determined, by varying the gas and liquid flow-rates for each tested solution. Foams are made of whey protein isolates (WPI) as foaming agent and xanthan gum (XG) at different concentrations to modulate the liquid viscosities, elucidating the effect of liquid base viscosities on the flow pattern.

2. Experimental facility and methods

2.1. SMX static mixers

The foams are elaborated by gas injection in a continuous flow of liquid using the static mixers SMX™ or SMX+™ (Sulzer, Winterthur, Switzerland). They have a respective inside diameter of 10 and 6 mm in diameter. One element of SMX consists of stainless steel crossed bars at angles of 45° over flow direction with ratios of length to diameter equal to one. Each element is rotated 90° prior to the previous element. Characteristics of the static mixers geometry are given in Table 1.

2.2. Hydraulic loop

The elements of static mixers are included in the hydraulic loop shown in Fig. 1. A gas injection device has been designed to feed

the static mixer section with air. It is located immediately upstream the first element of static mixer. Its specific geometry ensures that the gas is well dispersed all around the pipe diameter: the gas is introduced in the liquid by an annular dispenser with 12 tubular holes symmetrically located around the section. The injection holes are 0.5 mm in diameter for SMX10 and 0.3 mm in diameter for SMX+6.

The experimental loop includes a screw pump (Lenze™) to feed the static mixer with the liquid, as well as a liquid flow-meter and a gas mass flow-meter (Proline Promag 50 and Promass 80 from Endress + Hauser™). Two pressure sensors, Ceraphant T and Deltabar S (Endress + Hauser™), with adjusted measuring range until 3 bars, are located at the gas inlet and at the bounds of the static mixer to measure the pressure drop across the mixer. Thermocouples (Type K, Omega®) are used to record the temperatures of liquid and foam.

2.3. Fluids: whey protein – xanthan gum solutions

2.3.1. Composition and preparation

The liquid phase consists of Whey Protein Isolate (WPI) (Lactalis Ingredients, Bourbarré, France) with 95 wt% protein as surface-active agent and Xanthan Gum (XG) (Cargill, Bapte, France) acting as a thickener. Two WPI-XG solutions are prepared from 3 wt% WPI, and 0.35 and 0.6 wt% xanthan gum, referred to as WPI3XG35 and WPI3XG06 respectively. A third solution is obtained by adding NaCl (0.2 M) to WPI3XG035 to test the eventual effect of the ionic strength on the conformation of the protein and/or xanthan gum. These concentrations correspond to the ones used in the milk industry (Fox and McSweeney, 2003; Simon, 2001). For these concentrations, the two biopolymers are non-interacting and mutually soluble in the water (Benichou et al., 2007).

Dispersions are obtained using the following procedure. Batches of 20 L are prepared in a cylindrical mixing vessel (Guedu, Lavergne, France). Powders of WPI and xanthan are dispersed in water, at a controlled temperature (40 °C) and at pH 7, by the addition of 0.1 M NaOH solution, for better protein solubilization (Guimarães Pelegrine and De Moraes Santos Gomes, 2008). Stirring is carried out at low speed (50 rpm) to prevent formation of foam, during five hours, and then the batches are kept at 4 °C to ensure complete hydration of the polymers

2.3.2. Viscosity and surface tension

The viscosity of the WPI-XG mixtures is determined for each batch using the AR-1000N rheometer (TA Instruments, USA) with cone-plate geometry (60 mm diameter, 4° cone angle) at 25 °C. Shear rate $\dot{\gamma}$ ranged between 0.2 s⁻¹ and 2800 s⁻¹. The viscosity of the solutions is affected by the XG concentration, and the addition of sodium chloride decreases the viscosity. These solutions exhibit a shear thinning behavior as seen in Fig. 2 and can be fitted with a power law for shear rates between 0.3 and 700 s⁻¹.

$$\mu = K\dot{\gamma}^{n-1} \quad (1)$$

where τ is the shear stress, K is the flow consistency index and n the structure index.

The interfacial tension σ of the test solutions (3 wt% WPI) was measured with a K12 tensiometer (Krüss GmbH, Germany) using the Wilhelmy's plate method at 20 °C during 3 h. With a constant WPI concentration, the interfacial tension is 43 ± 1 mN m⁻¹ at long time (equilibrium value) for all the solutions. Neither xanthan gum nor sodium chloride changes the interfacial tension. The composition and properties of the solutions are gathered in Table 2.

Table 1
Characteristic values of Sulzer SMX10 and SMX+6.

Characteristic values	SMX10	SMX+6
Number of elements, n_e (-)	25	10
Pipe diameter, D (m)	0.010	0.006
Total length, L (m)	0.25	0.06
Hydraulic diameter, D_h (m) from Sulzer	0.00265	0.0018
Porosity, ε (-)	0.77	0.77

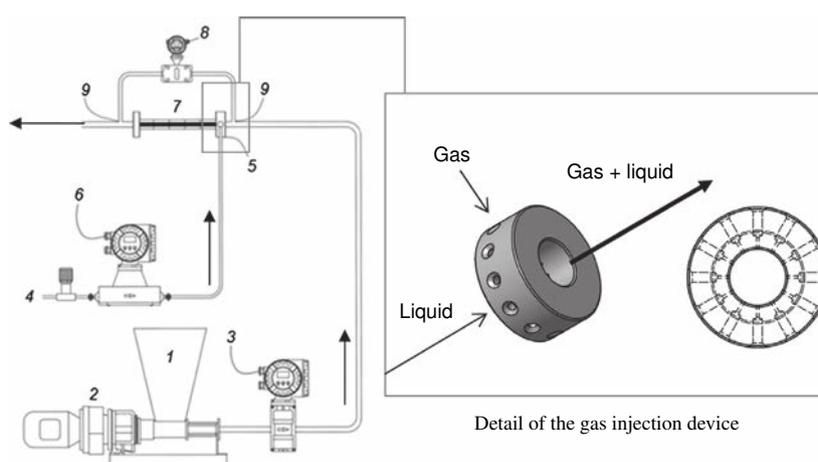


Fig. 1. Experimental rig (1) liquid feed; (2) pump; (3) volumetric liquid flow meter; (4) gas supply; (5) absolute gas pressure sensor, (6) mass gas flow meter; (7) static mixer; (8) differential pressure sensor; (9) temperature sensors. The gas injection device is zoomed on the right of the figure.

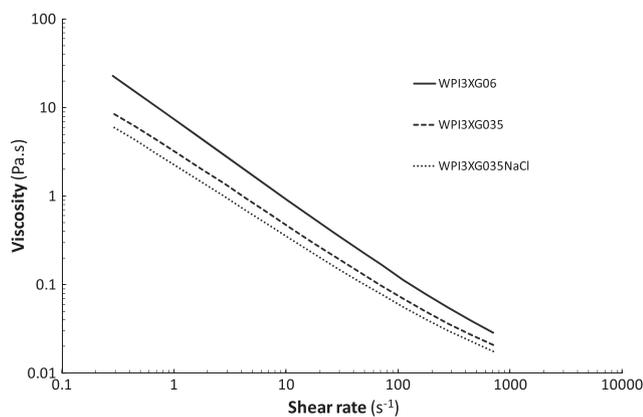


Fig. 2. Flow curves of the WPI-XG liquid solutions, 20 °C.

2.4. Foam void fraction

In the perfectly dispersed regime, assuming a no-slip velocity between the two phases, the void fraction α is determined from the input liquid and gas volumetric flow-rates, respectively Q_{liquid} and Q_{gas} , as follows:

$$\alpha_{theoretical} = \frac{Q_{gas}}{Q_{gas} + Q_{liquid}} \quad (2)$$

The experimental void fraction is deduced from the amount of gas incorporated in the foam. It is calculated from the weight, W_l , of the liquid solution filling a receptacle of given volume and that, W_f , of the foam filling the same receptacle. The deviation is 0.5%.

$$\alpha_{experimental} = \frac{W_l - W_f}{W_l} \quad (3)$$

Table 2
Composition and properties of liquid solutions at 25 °C.

Reference of the liquid solution	WPI (wt%)	XG (wt%)	NaCl 0.2 M (wt%)	ρ_l (kg m ⁻³)	K (Pa s ^{n}) Eq. (1)	n (-) Eq. (1)	Difference between experiments and model (%)	σ_l (mN m ⁻¹)
WPI3X06	3	0.60	0	1036	7.2 ± 0.4	0.12 ± 0.01	5.6	43 ± 1
WPI3X035	3	0.35	0	1034	3.1 ± 0.2	0.21 ± 0.01	4.9	43 ± 1
WPI3X035NaCl	3	0.35	1.2	1046	2.2 ± 0.2	0.24 ± 0.01	5.6	43 ± 1

3. Results and discussion

3.1. Hydrodynamic conditions

3.1.1. Pressure drop in the SMX static mixers

The power consumption in monophasic steady-state flow can be quantified in terms of the Fanning friction factor f , which is related to the only Reynolds number. The friction factor, f , is defined as:

$$f = \frac{\Delta P}{2\rho U_s^2} \frac{D}{L} \quad (4)$$

where D is the tube diameter, L the total length of the static mixers, ΔP , the pressure drop, U_s the superficial velocity, i.e. the average velocity in the empty duct. ρ and μ are respectively the density and the viscosity of the fluid.

The Reynolds number, Re , is defined as:

$$Re = \frac{U_s \rho D}{\mu} \quad (5)$$

The evolution of the friction factor versus Reynolds number can be represented by Eq. (6). K_p and C are two constants which only depend on the geometry shape, and hence are supposed to be invariant in a scale-up of static mixers:

$$f = \frac{K_p}{2Re} + C \quad (6)$$

The pressure drop measurements are carried out with sunflower oil and a solution of glycerol 25 w/w% (see properties in Table 3) to determine K_p and C values. Fig. 3 displays the friction factor curve obtained for these fluids versus the Reynolds numbers. The values of K_p and C , including their mean relative errors, are provided in Table 4 for the two SMX mixers. These values are in agreement with the literature values (Hirschberg et al., 2009). It is observed that the pressure drop factor in the SMX+6 is lower than for the standard SMX10 at a given Reynolds number, due to a slightly different geometry. Therefore, there is a certain compensating effect on the pressure drop increase at a given flow-rate, when switching from the SMX10 to the SMX+6.

3.1.2. Determination of process shear rate in the SMX static mixers

Metzner and Otto (1957) established a simple method to evaluate the shearing level in stirred tanks, based on a scale law of the process shear, $\dot{\gamma}_{app}$. The Metzner Otto constant, K_s , has to be determined experimentally in order to evaluate the apparent shear rate $\dot{\gamma}_{app}$ from the rotation frequency, N , in the stirred tank:

$$\dot{\gamma}^n = K_s N \quad (7)$$

A similar law can also be used to determine the effective shear rate in static mixers, replacing the number of rotations per unit time, N , by the ratio of superficial fluid velocity and pipe diameter, U_s/D as proposed by Alonso et al. (2000):

$$\dot{\gamma}_{app} = K_s \frac{U_s}{D} \quad (8)$$

Table 3
Properties of fluids at 20 °C.

Solution	ρ (kg m ⁻³)	μ (Pa s) Eq. (1)		
Guar 0.5 wt%	1005	Non Newtonian ($10 < \dot{\gamma} < 2000$ s ⁻¹)	$K = 1.29$ Pa s ^{n}	$n = 0.48$
Guar 0.35 wt%	1003	Non Newtonian ($10 < \dot{\gamma} < 2000$ s ⁻¹)	$K = 0.32$ Pa s ^{n}	$n = 0.65$
Sunflower oil	910	Newtonian	$K = 0.0670$ Pa s	$n = 1$
Glycerol 25 wt%	1060	Newtonian	$K = 0.0043$ Pa s	$n = 1$
Water	1000	Newtonian	$K = 0.0010$ Pa s	$n = 1$

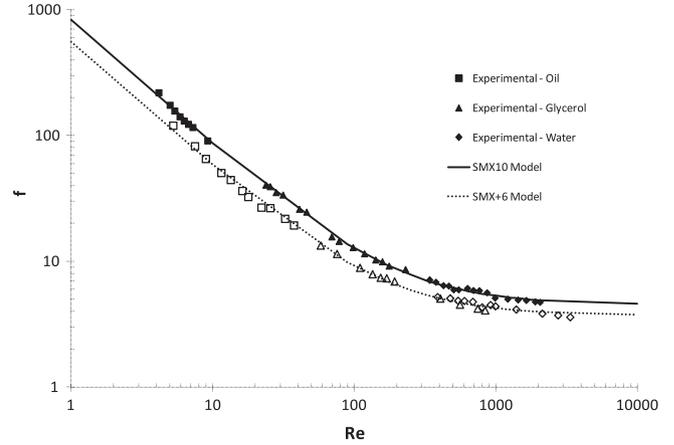


Fig. 3. Experimental data (symbols – hollow for SMX+6 and full for SMX10) and power characteristic model (lines – dotted for SMX+6 and solid for SMX10).

Table 4
Characteristic hydrodynamical constants of SMX10 and SMX+6.

SMX	K_p	C	Experiment-model deviation (%)	K_s
SMX+6	1100	3.7	3.7	16 ± 1
SMX10	1650	4.5	3.3	24 ± 2

The method consists of determining the shear rate through the viscosity μ of a non-Newtonian fluid following a given Ostwald-de Waele rheological behavior (Chhabra and Richardson, 2008):

$$\mu = K \dot{\gamma}_{app}^{n-1} \quad (9)$$

By representing the experimental variation of friction factor versus generalized Reynolds number (Eq. (10)) in laminar flow ($Re < 50$), one obtains a friction factor curve that is shifted from the reference Newtonian curve by a factor K_s^{n-1} . The gap between the two curves, i.e. the factor K_s^{n-1} , can be determined in log scale.

$$Re_g = \frac{U_s^{2-n} \rho D^n}{K} \quad (10)$$

Then, merging the curves into a single calibration curve is achieved using the effective Reynolds number:

$$Re_{eff} = \frac{U_s^{2-n} \rho D^n}{K K_s^{n-1}} \quad (11)$$

Here, the solutions of guar gum, 0.35 wt% and 0.5 wt% are used as non-Newtonian fluid; their rheological properties are listed in Table 3. Experimental results are shown in Fig. 4, where the friction factor f is plotted as a function of the effective Reynolds number, Re_{eff} . The curves match in the range of Reynolds number below 15 for $K_s = 16$.

Concerning the SMX10, previous experiments allowed to determine a value of K_s equal to 24 (Bone, 2005). It can be observed that the value of K_s corresponding to the SMX+6 is lower than that of

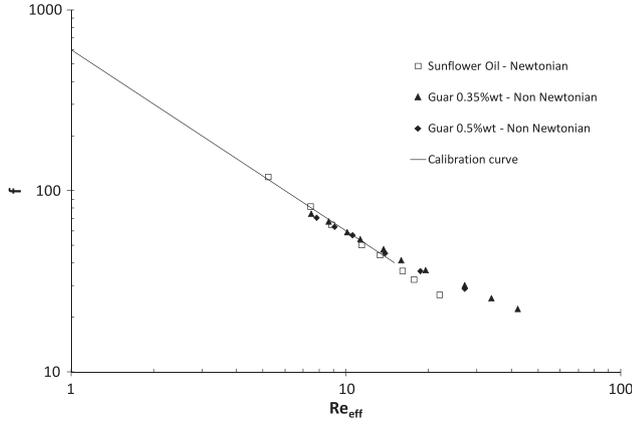


Fig. 4. Friction factor versus effective Reynolds number in the SMX+6 with flow of sunflower oil (hollow symbols) and guar solutions (full symbols).

SMX10. This obviously confirms the fact outlined in Section 3.1.1 with the lower pressure drop constant. This implies that, at a given flow-rate per unit surface, a lower level of shear is achieved in the SMX+6. However, the process shear rate $\dot{\gamma}_{app}$ is higher in the SMX+6 for a given flow rate due to the smaller section.

3.1.3. Dimensional process shear rate

The apparent shear rate in foaming condition needs to be evaluated as it is a strategic parameter for mixing and gas bubbles breakup. However, its determination from Eq. (8) is only valid in creeping flow. The dimensional range of process shear rate is presented in Fig. 5 as a function of Reynolds numbers for the two SMX, as far as the Reynolds number value indicates the reliability of the shear rate model in the range of operating conditions. The curves are plotted for two fluids of different properties, one is a monophasic oil (glycerol solution 25 wt%), and the other is a typical foam of 0.8 void fraction, by considering the density, ρ_f , as $(1 - \alpha) \rho_l$ and the velocity, U_s , is the sum of liquid U_l and gas U_g velocities. The viscosity used in the Reynolds number is of the order of magnitude taken from our experiments, about 0.1 Pa.s nearly constant for the whole flow rates range. This diagram shows that even with the highest flow-rates (gas and liquid) encountered in this study, the flow is laminar, which justifies the use of the process viscosity calculation.

3.1.4. Energetical efficiency

Some insights in the mixers energy efficiency can be provided by the ratio of the apparent shear rate $\dot{\gamma}_{app}$ to the specific energy E (per unit mass) required for the foaming. This specific energy is

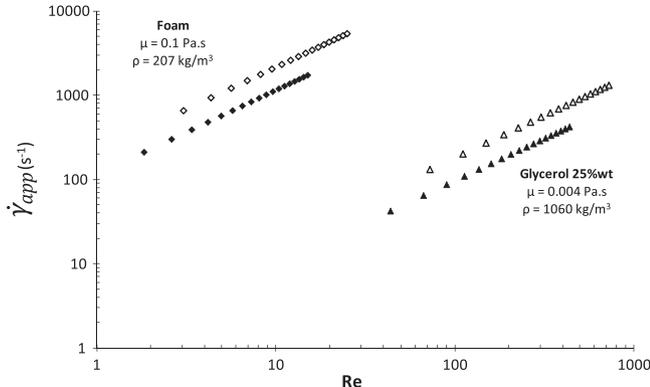


Fig. 5. Evolution of apparent shear rates in SMX+6 (hollow symbols) and SMX10 (full symbols) respectively with flow of foam (\diamond , $\alpha = 0.8$ and $\mu = 0.1$ Pa.s) and flow of a glycerol solution (\blacktriangle) versus Reynolds numbers.

defined as the ratio of the pressure drop, ΔP , to the foam density, ρ . From Eqs. (4), (6), and (8), the SMX efficiency is expressed as:

$$\frac{\dot{\gamma}_{app}}{E} = \frac{\rho K_s D}{\mu K_p L} \quad (12)$$

This efficiency factor, as shown in Eq. (13), only depends on the K_p and K_s values and on the shape ratio, allowing a comparison between the two mixers for a given processed fluid but independently of the operating conditions. The ratio of the respective efficiency factors for the two devices reads:

$$\frac{\left(\frac{\dot{\gamma}_{app}}{E}\right)_{SMX+6}}{\left(\frac{\dot{\gamma}_{app}}{E}\right)_{SMX10}} = \frac{\left(\frac{K_s D}{K_p L}\right)_{SMX+6}}{\left(\frac{K_s D}{K_p L}\right)_{SMX10}} \quad (13)$$

The efficiency related to the SMX+6 is higher than the one in the SMX10 by a factor of 2.85. This gives an advantage for using the SMX+6 in the sense that the energy consumption is less for reaching the same shear rate. It also means that the length of the SMX+6 used in this study (0.06 m) could be increased up to 0.15 m while still maintaining the same efficiency as opposed to the SMX10 of 0.25 m length. This could be taken into account according to the effect of the residence time towards the bubble breakup and the final bubble foam diameter.

3.2. Flow pattern map

The experiments aimed to determine the flow transition are carried out in the SMX10 and in the SMX+6, with a reference method to avoid any hysteresis effect. For a given value of the liquid flow-rate in the range [8–50] L/h, the gas flow-rate is decreased monotonically starting from the maximum value, where the slug flow is generally observed. The tested range of gas flow-rate is [0.02–0.45] Nm³/h. Once the gas plugs disappear, the corresponding gas flow-rate is recorded as well as pressure drop and gas injection pressure. Limits between the regimes are visually determined at the outlet of the static mixers.

The flow pattern map coordinates ($U_{s,l}$, $U_{s,g}$) presented in Fig. 6, are established from eight flow-rates pairs (Q_l , Q_g) and for three liquid solutions. In the present study, the plotted frontier corresponds to the transition between the two flow regimes observed within the experimental range, which are the “dispersed” regime that allows a complete incorporation of the gas, and the so called “slug flow” regime that occurs at high gas flow-rates. Other types of existing flow patterns (plug, annular, stratified, etc.) are not observed over the present range of conditions. Theoretical foam void fractions are mentioned on the maps by contours of iso-values; they are straightforwardly deduced of the flow-rates, but of course the validity is limited to the bubble regime area. Some representative experimental values taken at the frontiers are gathered in Table 5. Operating with a set of velocities located at the left of the limit will ensure that there is no gas loss in the process, and hence a better process yield. Increasing the gas velocity beyond the frontier results in a gas waste, then the foam void fraction cannot be controlled. The limits obtained for SMX10 and SMX+6 seem to be consistent with the previous hydrodynamic study: for a given liquid matrix the dispersed regime is reached for a higher gas flow-rate in the case of a smaller hydraulic diameter in the SMX+6. Thus, reducing the SMX diameters does not induce radical changes in the flow map frame, so that no particular issues are brought to light in the scale-down. It can also be observed that the limit shifts toward lower gas velocity with increasing liquid phase viscosity. Most experiments have been made in triplicate in the case of SMX10 and are at least doubled in the case of SMX+6 (ranges of deviation are shown for each flow condition in Fig. 6). Repeat experiments show a 5% standard deviation for the

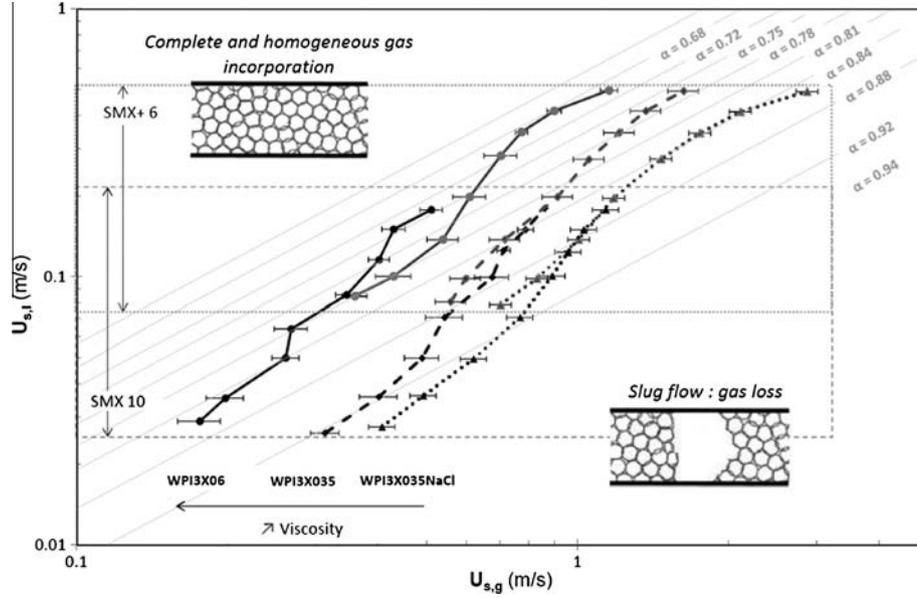


Fig. 6. Limits between dispersed and slug flow regimes in SMX10 (black) and SMX+6 (grey) for foams from WPI3XG06 (●), WPI3X035 (◆) and WPI3X035NaCl (▲) solutions.

Table 5
Experimental conditions for few runs.

SMX	Q_l (L/h)	Q_g (Nm ³ /h)	f (calculated from Eq. (8))	Re	$\dot{\gamma}_{app}$ (s ⁻¹)	α_{max}
<i>WPI3XG035NaCl</i>						
SMX10	8	0.116	333	2.3	360	0.94
	28	0.254	66	12	800	0.90
	50	0.325	34	25	1100	0.87
SMX+6	8	0.072	102	6	870	0.90
	28	0.143	37	18	1740	0.83
	50	0.295	16	46	3100	0.85
<i>WPI3XG035</i>						
SMX10	8	0.090	389	1.9	300	0.92
	28	0.195	73	11	690	0.87
	50	0.248	38	22	970	0.83
SMX+6	8	0.058	101	6.1	760	0.87
	28	0.110	30	23	1510	0.79
	50	0.170	18	41	2370	0.77
<i>WPI3XG06</i>						
SMX10	8	0.050	372	2.1	220	0.86
	28	0.104	86	9.2	520	0.80
	50	0.144	42	20	770	0.74
SMX+6	8	0.038	84	7.4	580	0.82
	28	0.074	27	25	1330	0.72
	50	0.121	16	47	2180	0.71

gas flow-rate at the transition, while there is less than 1% standard deviation regarding liquid flow-rates uncertainty.

From these results, an attempt is made to gather the combined effects of the liquid solution viscosity and the operating conditions on the regime transition and the maximum void fraction attainable in the bubble regime. The global capillary number Ca is introduced, corresponding to the ratio between a viscous stress scale $(U_{s,l}\mu_l)/D_h$ and a surface forces scale σ_l/D_h :

$$Ca = \frac{U_{s,l}\mu_l}{\sigma_l} \quad (14)$$

In Fig. 7, the evolution of this maximum void fraction versus the corresponding capillary number seems to correlate the results for all the tested solutions. It clearly shows that the maximum void fraction decreases with increasing liquid viscosity, for the two SMX. The data merge onto a single curve for each SMX, the lower

level for the SMX+6 being explained by the lower K_s in this device, compared to the SMX10. Concomitantly, for higher capillary numbers (high liquid viscosities), the dynamic conditions are more favorable for the bubble breakup, that contributes to the foam stability (Zaki et al., 2002). It is important to note here that the upper limit of the void fraction remains compatible with the food foams specifications. The correlation proposed in Fig. 7 is also valid concerning the data from Talansier (2009) and obtained in SMX10 respectively with egg white and with glucose syrup 60 wt% mixed with Tween 20 0.1 wt%. It might be emphasized that these solutions exhibited different values of the interfacial tension than that of the presently tested solutions.

In the case of the SMX+6, it is possible to reach higher capillary numbers and then to produce finer bubbles and more stable foams on a limited extent. In that sense, the initial objective of reducing the void fraction with a smaller hydraulic diameter and higher

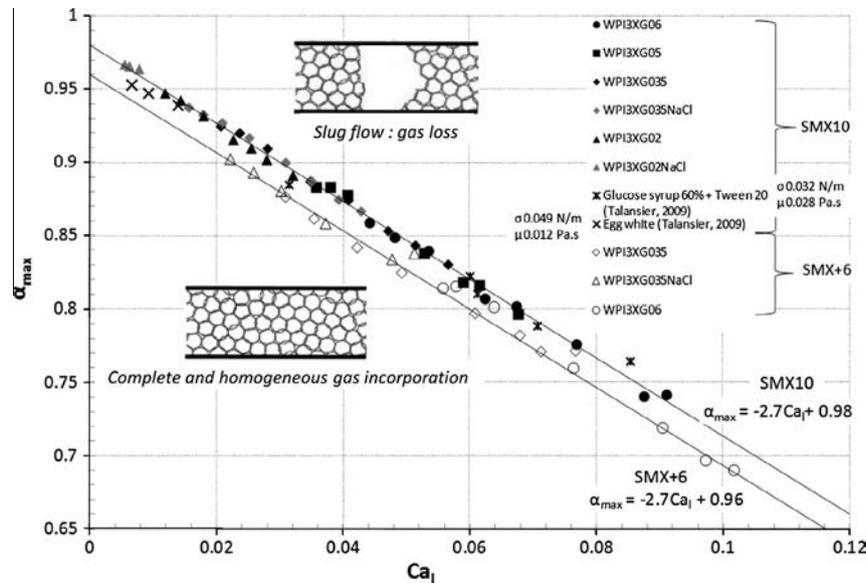


Fig. 7. Maximum void fraction, α_{\max} , available for different liquid capillary numbers, Ca_l in the SMX+6 (hollow symbols) and in the SMX10 (full symbols).

fluid viscosities can be fulfilled, with the drawback of a certain limitation of the void fractions. The chart shown in Fig. 7 can be used to quantify this feature. For a complete knowledge of the process, this representation can be further developed with the lower limit of void fractions corresponding to convenient foam structure, and provide a sizing tool for the foaming process in the SMX. It might be emphasized that literature models, referred to in Section 1, fail to predict the flow regime transitions in the case of the presently tested two-phase gas-liquid systems. In particular, the models of Taitel and Dukler (1976) and Baker (1954), which are the most quoted in the literature for horizontal tubes, give contradictory predictions that are not in agreement with the present experiments in the SMX. Two reasons may explain the observed discrepancies: literature models concerns only plain tubes and secondly have not been built from data obtained with fluids like these tested in this work. As a conclusion, experiment still remains necessary to bound process conditions.

4. Conclusions

WPI and XG foams were produced in SMX10 or SMX+6. The hydrodynamic studies allowed determining the values of the power constant K_p and that of process shear rate coefficient K_s . The values showed a pressure drop reduced with the SMX+6. This means the effect of processing with higher flow-rates in SMX+6 will not increase appreciably the pressure drop.

The respective effect of the gas and liquid flow-rates, of the hydraulic diameter and of the liquid solution viscosities on the dispersed and plug-flow regimes transition were also experimentally investigated. The two-phase flow experiments were performed over a wide range of gas and liquid flow-rates with three liquid solutions of different viscosities. The flow map highlighted a significant flow-rate range in which gas incorporation is fully achieved. The viscosity of liquid solution strongly affects the position of the frontier between flow regimes. The limit shifts towards smaller superficial gas velocities with increasing viscosity as clearly shown with the introduction of a liquid capillary number. Therefore, the maximum void fraction attainable is lower with higher liquid viscosity. The difference in the SMX geometry, characterized by a different hydraulic diameter, does not significantly affect the shape of the flow pattern frontiers: it can be observed that the frontiers are

rather continuous in the explored domain of flow-rates. The maximum void fraction attainable is slightly lower with the SMX+6, but remains sufficient for food foams applications.

This study allowed determining suitable foaming conditions with control of foam void fraction. This work constitutes a basis, for, in a next step, focus on process-product interactions and resulting foam properties like bubble size, rheology and stability.

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