



Ultrasound-induced emulsification of subcritical carbon dioxide/water with and without surfactant as a strategy for enhanced mass transport



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ARTICLE INFO

Article history:

Received 21 February 2013

Received in revised form 14 May 2013

Accepted 26 May 2013

Available online 3 June 2013

Keywords:

CO₂

Aqueous media

Subcritical

Emulsion

Mass transport

Hydrolysis

ABSTRACT

Pulsed ultrasound was used to disperse a biphasic mixture of CO₂/H₂O in a 1 dm³ high-pressure reactor at 30 °C/80 bar. A view cell positioned in-line with the sonic vessel allowed observation of a turbid emulsion which lasted approximately 30 min after ceasing sonication. Within the ultrasound reactor, simultaneous CO₂-continuous and H₂O-continuous environments were identified. The hydrolysis of benzoyl chloride was employed to show that at similar power intensities, comparable initial rates ($1.6 \pm 0.3 \times 10^{-3} \text{ s}^{-1}$ at 95 W cm^{-2}) were obtained with those reported for a 87 cm³ reactor ($1.8 \pm 0.2 \times 10^{-3} \text{ s}^{-1}$ at 105 W cm^{-2}), demonstrating the conservation of the physical effects of ultrasound in high-pressure systems (emulsification induced by the action of acoustic forces near an interface). A comparison of benzoyl chloride hydrolysis rates and benzaldehyde mass transport relative to the non-sonicated, 'silent' cases confirmed that the application of ultrasound achieved reaction rates which were over 200 times faster, by reducing the mass transport resistance between CO₂ and H₂O. The versatility of the system was further demonstrated by ultrasound-induced hydrolysis in the presence of the polysorbate surfactant, Tween, which formed a more uniform CO₂/H₂O emulsion that significantly increased benzoyl chloride hydrolysis rates. Finally, pulse rate was employed as a means of slowing down the rate of hydrolysis, further illustrating how ultrasound can be used as a valuable tool for controlling reactions in CO₂/H₂O solvent mixtures.

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1. Introduction

One way of improving the separation between immiscible species following synthetic procedures, is to use a biphasic system involving an organic solvent and an aqueous phase [1,2]. Owing to its low critical point (31.1 °C/73.8 bar) and ease of separation upon depressurisation, subcritical or supercritical CO₂ may be employed usefully as a replacement for the organic phase [3,4]. Moreover, as the solvating power of CO₂ is confined to small, relatively non-polar compounds, the presence of a second aqueous phase can greatly increase the accessibility of hydrophilic species (e.g. salts), that are commonly employed in many organic chemistry reactions. Additionally, the reaction of CO₂ with water leads to the formation of carbonic acid, providing an intrinsic proton source (pH 3–4) [5,6], which may be advantageous for several processes [7].

Due to the presence of a liquid–liquid interfacial barrier, mass transfer rates in biphasic systems may be improved by increasing the interfacial area through the formation of an emulsion [8]. Emulsions of CO₂/H₂O (C/W) have the distinct advantage over conventional organic-phase emulsions of allowing the complete reversal to the biphasic system upon depressurisation of the system. Furthermore, the droplet size and stability of the emulsion may be controlled by specially-designed surfactant molecules allowing greater fine-tuning of the reaction system [9]. Dispersions of CO₂ and water, as emulsions or microemulsions [10], have been developed for a range of synthetic processes, including hydrogenations [11–14], hydroformylations [15], the synthesis of metal nanoparticles [16], and polymerisations [17]. Due to its favourable properties, *pulsed* ultrasound may also be used to generate emulsions and therefore increase the rate of reactions. Ultrasound may also be considered as a tool for green chemistry, by allowing the use of benign conditions and enhancing energy efficiency [18]. The use of ultrasound in high-pressure systems has been reported for several extraction processes in CO₂ [19–21].

A previous study by Timko et al. investigated the hydrolysis of benzoyl chloride in a CO₂/H₂O biphasic system under ultrasonication in a reactor of 87.2 cm³ volume [22]. In the present study, we have designed a 1 dm³ high-pressure ultrasound reactor and used this

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to emulsify a dense CO₂/H₂O solvent system. Scaling up the hydrolysis of benzoyl chloride, we show that the efficiency is not lost at this scale-up factor, highlighting the conservation of the positive physical effects of ultrasound. Emulsion formation and subsequent breakdown was observed by passing the emulsions through a 30 cm³ view cell and monitoring the change in turbidity over time. The advantages of ultrasound were further demonstrated for the hydrolysis reaction in the presence of Tween 80, a bio-compatible surfactant, able to stabilise a uniform C/W emulsion with estimated surface areas greater than 480 m² kg⁻¹. The combined effects of ultrasound and Tween 80 resulted in faster reaction rates than those reported for a pre-formed H₂O in CO₂ (W/C) microemulsion [23], with higher energy efficiency than without the use of surfactant. Overall, the results demonstrate the potential utility of this reactor methodology for industrial applications.

2. Materials and methods

2.1. Reagents

Benzaldehyde (>99%) [100-52-7], benzoyl chloride (>99%) [98-88-4], benzoic acid (>99%) [65-85-0], octane [111-65-9] and polysorbate [9005-65-6] (Tween 80), were purchased from Sigma-Aldrich UK, and used without further purification. Liquid carbon dioxide was obtained from BOC UK, CP grade, and had a given purity of 99.995%.

2.2. System configuration (Fig. 1)

A high-pressure, cylindrical 1 dm³ stainless-steel vessel (Parr, USA I.D. = 9.5 cm, H = 16 cm) was designed to incorporate an ultrasound probe and manufactured for use in the present study. CO₂ was delivered to the vessel via a high-pressure pump (Thar Instruments), electronically connected to a PC. The amount of CO₂ entering the reactor was recorded via a mass flow meter positioned in-line before the pump. The vessel was insulated by an electric heating jacket, and the temperature was controlled by a temperature controller (Parr 4838) coupled to a J type thermocouple, positioned in an immersed thermowell. The pressure was recorded to within ±0.1 bar using a transducer (Druck PTX 1400; Druck Ltd, Leicester, UK) with dedicated display. Ultrasound was introduced from a 20 kHz VCX-1500W processor (Sonics & Materials, CT, USA), coupled with a titanium alloy horn of tip diameter 2 cm. Tip amplitude (20–100%) was manually selected and related to the energy transferred to the fluid, while the pulse rate could define the ratio of ultrasound ON/ultrasound OFF based on a sonication cycle (e.g. 25%, 1 s ON/3 s OFF). A calibration between delivered and dissipated power was undertaken using the calorimetric method [24]. Power density values reported here (W cm⁻³) were obtained by dividing the total energy per pulse time (J s⁻¹) by the vessel volume. When comparing vessels of different sizes, power intensity (W cm⁻²) may also be encountered. Here power intensity values were calculated by dividing the power by the irradiating surface area of the tip. As a consequence of high amplitude settings resulting in a rapid ~3 °C rise in the temperature of the vessel, water cooling was employed to keep the vessel temperature constant during experiments. A dual piston HPLC pump (Jasco PU-2087) with a maximum flow rate of 20 cm³ min⁻¹ was used to recirculate the vessel contents. Connections to the vessel were made by 1/4" and 1/8" stainless steel Swagelok tubing, while connections to the pump were made in 1/16" tubing.

2.3. General method for sonication of the CO₂/H₂O biphasic

The vessel was initially heated until the desired temperature (30 °C) was reached. Water was then added to give a volume be-

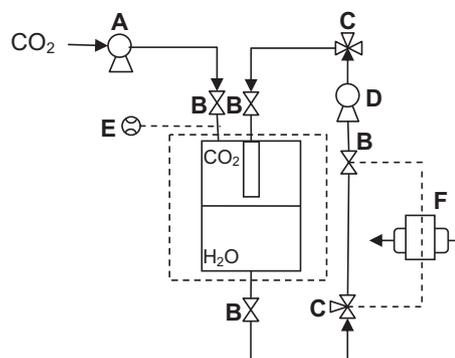


Fig. 1. Schematic showing the experimental setup utilised for sonication of CO₂/H₂O in the 1 dm³ reactor: A: CO₂ pump; B: high-pressure valve; C: 3-way sampling valve; D: circulation pump; E: 6-port valve for injecting compounds into the CO₂ stream; F: bypass of the H₂O recirculating line for visualisation through the 30 cm³ view cell.

tween 400 and 650 mL. CO₂ was supplied to the sealed vessel at a flow rate of ~40 g min⁻¹, until the desired pressure was reached. Following 30 min of equilibration/recirculation, sonication was initiated and continued for a known time.

2.4. Observation of emulsion stability

A 30 cm³ high-pressure stainless-steel view cell I.D. = 2.5 cm, H = 5.7 cm (Parr Instruments, Moline, IL, USA) fitted with two in-line sapphire windows (D = 1.5 cm), was used to visualise the dispersion generated in the ultrasound reactor (Fig. 1F), at 30 °C/80 bar. Heating was provided by two electric heating plates connected to a dedicated controller, and monitored by an immersed J type thermocouple. Emulsion stability was assessed by measuring the increase in light intensity (Visilight I-Led) over time, following termination of a 10-min sonication/circulation period. Time-lapse recordings were carried out using a Photron high-speed camera at 10-s time intervals, and the digital images transferred to a PC, using a dedicated interface. The images were analysed with ImageJ for the sum of the greyscale pixel values (integrated density) inside a defined area, and each time-point value divided by the maximum density recorded in the absence of turbidity. Experiments were also performed under conditions above the CO₂ critical point in order to minimise any effect associated with phase transition.

2.5. Compound injection, sampling and analysis

For greater precision, hydrophobic compounds were added to the CO₂ phase via a 6-port valve positioned in-line with the CO₂ feed, and connected by 1/16" Swagelok tubing. A standard 1 mL bypass loop was used to inject a known amount of each compound. Following equilibration of the reactor contents at 30 °C/70 bar, either benzaldehyde or benzoyl chloride was injected into the flowing CO₂ feed, while final pressurisation to 80 bar was achieved. Aliquots (2 mL) were taken from the recirculating H₂O line at the specified time intervals and analysed immediately by gas chromatography (Shimadzu GC-2010, combined with autosampler and injector AOC-20, and equipped with a DB-5 column and an FID). Four or five aliquots were taken over the course of each experiment at different time points. For the same set of conditions, experiments were repeated three times, generating aggregated data of non-overlapping time points. The final concentration was obtained from a standard calibration line. The H₂O recirculating line originated from a 14-cm 1/4" stainless steel dip tube immersed in the reactor. The sampling point was therefore positioned 8 cm below the probe tip. Aliquots from the CO₂ phase on the other hand were

taken from a line originating from a port at the top of the reactor, and were collected by slowly bubbling into cold acetone. In order to verify that all of the compounds tested were dissolved in the CO₂ phase prior to initiating sonication, visual experiments were conducted by injecting 100 μL of compound into the view cell charged with just CO₂. At the pressure and temperature described in this paper, only a single phase was observed.

2.6. Benzaldehyde mass transport

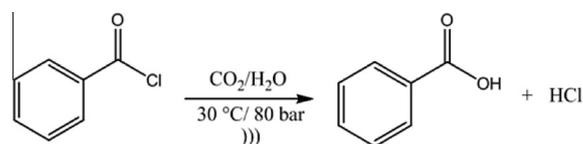
Mass transport experiments were conducted in order to assess the dispersive capacity of ultrasound for this vessel geometry (1 dm³) using a hydrophobic compound, and to potentially compare mass transport rates with other systems reported in the literature. Benzaldehyde was chosen for the mass transport experiments as its solubility in CO₂ has been studied previously [25,26]. Moreover, it contains an important functional group for organic chemistry, namely the carbonyl group. The mass transport of benzaldehyde from CO₂ to H₂O was assessed by the two-film theory, previously described for similar systems by Timko [27], and Tai et al. [28]¹.

Due to its influence on the mass transport, the partition coefficient was determined by measuring the concentrations of benzaldehyde partitioned in both phases, following 4 h of equilibration, and the resulting value was found to match to within 5% that reported in the literature ($K_{CW} = 12$) [29]. For comparison with values reported in the literature, the effective mass transport coefficient $K_{mrc}a^*$ (s⁻¹) was calculated, where 'a*' is the surface area per unit volume (cm⁻¹).

2.7. Kinetics of benzoyl chloride hydrolysis

The hydrolysis of benzoyl chloride represents a useful reaction for modelling kinetics in which a hydrophobic species (the acid chloride) must cross the interface, meet the hydrophilic species (H₂O), and react (Scheme 1). In the present system, the reverse case is also true, whereby H₂O travels to the CO₂ side and reacts with the acid chloride.

Initial rates of benzoyl chloride hydrolysis were analysed under amplitude settings of 20–70%, and were found to follow first-order kinetics, as previously reported [23,30].¹ The reaction was also studied in the presence of the surfactant polysorbate, Tween 80. This surfactant was chosen due to an existing in-depth knowledge of its C/W emulsion properties from the literature [31]. To this end, the same procedure was followed as that described above other than 0.5 or 1.0 wt.% of surfactant (with respect to water) was dissolved in the water prior to charging the vessel. Samples were taken at various time points, and the reaction was quenched by trapping the carboxylic acid hydrolysis product in acetone on ice (1:6 dilution by volume) before analysing by GC. As sonication in the presence of Tween resulted in a greater fraction of CO₂ being dispersed in H₂O compared to when sonication was carried out without Tween, a few considerations had to be made. To investigate whether or not the benzoic acid hydrolysis product had formed *in situ* as a result of ultrasound mixing, or by mixing during sample depressurisation, the same experimental protocol was carried out with an additional visual aid using the view cell. Following 1 min of sonication, ultrasound was stopped and the contents were recirculated until the emulsion reverted to the biphasic. After 1 h, when only a clear solution was observed to pass through the view cell, samples were collected in the usual manner. The sample concentration closely paralleled that obtained from 1 min ongoing sonication, indicating that the benzoic acid had not formed as a consequence of the benzoyl chloride-containing emulsion rapidly



Scheme 1.

breaking down upon depressurisation into the sample vial. The yields were plotted as a function of time at different ultrasound power settings.

3. Results and discussion

3.1. Emulsion appearance and stability

Emulsions originating in the H₂O-continuous phase were assessed first. Approximately 10 min after commencing sonication, turbidity was observed to increase inside the view cell (Fig. 2a and b). Following the ceasing of sonication, this turbidity remained for approximately 30 min, gradually reverting to the biphasic with a characteristic demulsification front (Fig. 2c). Droplets could be observed to collect on the upper part of the inside wall of the window, which increased in size and number over time, indicating that the emulsion was breaking down by CO₂ droplet coalescence.

Turbid emulsions generated by ultrasound were reported as being indicative of droplets with Sauter mean diameters of 9 μm for H₂O droplets in CO₂-continuous (W/C), and 15 μm for CO₂ droplets in H₂O-continuous (C/W) emulsions [22]. In our system, turbidity was observed regardless of whether the H₂O- or CO₂-continuous phase was recirculated through the view cell during sonication (Fig. 3). This allowed measurement of emulsion stability by assessing the decrease in turbidity in the view cell. The data points for the fastest increase observed in light intensity were plotted and found to fit a second-order polynomial function ($r^2 > 0.99$, Fig. 4). This was found to closely match the observed time course for emulsion breakdown visualised by eye. Volume fractions of CO₂ in H₂O were evaluated by the tracer technique,¹ and were found to be between 1% and 6%, depending on the standoff distance of the biphasic line from the probe (Fig. 5). Optimal dispersion of CO₂ in H₂O was achieved when the biphasic line was positioned between 1 cm and 2 cm below the probe surface. This observation was consistent with previous studies in liquids at ambient conditions that showed cavitation bubble velocity and bubble flow rate decrease dramatically in intensity at distances of more than 4 cm below the probe [32]. Emulsification may also result in H₂O droplets dispersing in the CO₂ phase. As a consequence, water collected from the CO₂-continuous phase revealed the H₂O volume fraction to be 6.5% (± 1) following sonication.

3.2. Mass transport

Following injection of 1 mL benzaldehyde, sample analysis revealed that its concentration measured in the upper part of the

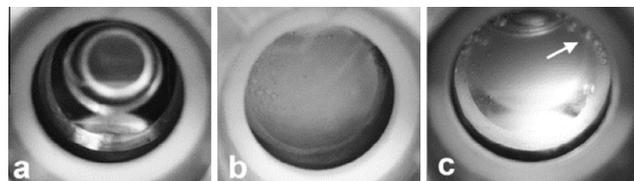


Fig. 2. Turbidity of the recirculating H₂O-continuous phase as observed through the view cell, following sonication in the 1 dm³ high-pressure reactor. (a) Circulation, no sonication; (b) 10 min following the start of sonication (c) approximately 10 min after stopping sonication, CO₂ droplets collect on the upper inside wall (arrow). (Conditions: 30 °C/80 bar, 70% amplitude, 25% pulse rate).

¹ See supplementary data.

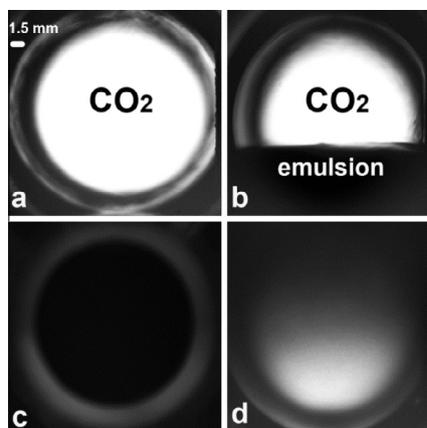


Fig. 3. Darkening of the recirculated CO₂-continuous phase through the view cell, following sonication in the 1 dm³ high-pressure vessel: (a) for image analysis experiments, a light source was located at the rear window to illuminate the contents of the cell; (b) approximately 3 min after the start of sonication, a turbid phase was seen in the cell; (c) after 10 min of sonication/circulation, complete opacity was observed; (d) 5 min after stopping sonication, light transmittance was partially restored due to reversal to the biphasic. Conditions: 30 °C/80 bar, 70% amplitude, 25% pulse rate.

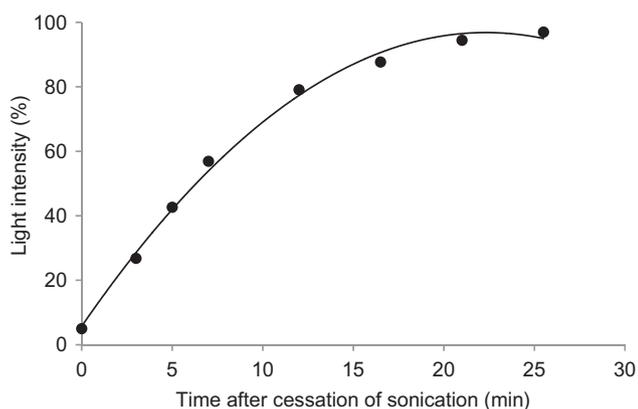


Fig. 4. Increase in light intensity through the view cell following termination of sonication, and reversal from emulsified to biphasic conditions. Conditions: 30 °C/80 bar, 70% amplitude, 25% pulse rate.

reactor during sonication was roughly ten times higher than that in the lower part. Together with volume fraction measurements from the previous section, the results indicated that two distinct environments existed simultaneously during sonication, one CO₂-continuous and one H₂O-continuous in the upper and lower parts of the reactor, respectively. The change of benzaldehyde concentration over time under only circulation ('silent') and under sonication conditions, is presented in Fig. 6. The equilibrium for benzaldehyde mass transport was reached within approximately 2 h under only circulation conditions, whereas it was reached within 1 min following the start of sonication. Analysis of the log plot yielded slopes of 2.7 and $0.023 \times 10^{-2} \text{ s}^{-1}$ for the sonicated and 'silent' case, respectively. Insertion of known parameters ($V_W = 650 \text{ cm}^3$, $V_C = 350 \text{ cm}^3$ and $K_{CW} = 12$), yielded respective $K_{mtc}a^*$ values of $1.9 \times 10^{-3} \text{ s}^{-1}$ and $\approx 0.016 \times 10^{-3} \text{ s}^{-1}$, and calculation of their ratio afforded a sonic enhancement factor of ~ 120 . The value for the sonicated case was within the range for the transfer of most species in liquid–liquid mixtures. Tai et al. studied the impeller-induced mass transport of zinc(II) ions in a 1.3 dm³ batch with a CO₂/H₂O biphasic, reporting values of $0.2\text{--}2.6 \times 10^{-3} \text{ s}^{-1}$ at impeller speeds of 400–1000 rpm at 40 °C/83 bar [28]. Mass transport is furthermore highly dependent on the type of contactor and agitation

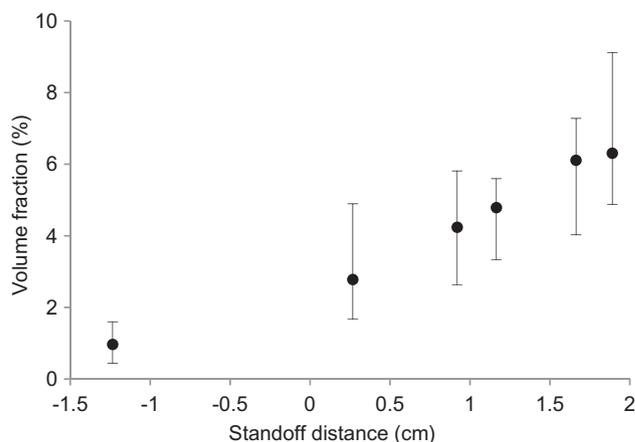


Fig. 5. Volume fractions of CO₂ dispersed in H₂O following sonication. The standoff distance signifies the position of the biphasic line with respect to the probe. The negative values denote conditions in which the biphasic line was above the probe surface (i.e. probe immersed in H₂O). The position of the biphasic line was adjusted by changing the initial H₂O volume in the reactor, which was (from left to right): 800, 700, 650, 630, 600 and 580 mL H₂O. The number of experimental points for each standoff distance was between 3 and 7. No tracer was detected in the H₂O phase after 3 h of only recirculation. Conditions: 30 °C/80 bar, 70% amplitude, 25% pulse rate.

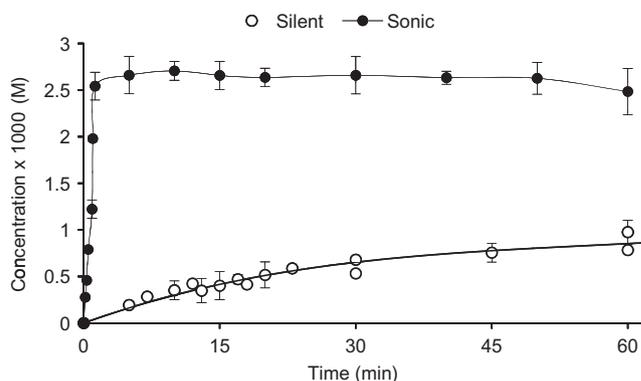


Fig. 6. Benzaldehyde concentration in samples from the H₂O-continuous phase for 'silent' and sonicated (70% amplitude, 25% pulse rate) environments. Conditions: 1 mL benzaldehyde ($C_{CO} = 0.028 \text{ M}$), CO₂ = 25 wt% (220 g CO₂, 650 g H₂O), 30 °C/0 bar.

employed. For example, the effective mass transport coefficient ranged between 0.016 and $1.6 \times 10^{-3} \text{ s}^{-1}$ for a stirred tank, or as high as 30.6 s^{-1} in a microchannel [33]. In a different study, the effective gas–liquid mass transfer coefficient increased from ~ 0.1 to $10 \times 10^{-3} \text{ s}^{-1}$ when 500 rpm stirring was complemented with ultrasound in a 1 L vessel [34]. Separation of K_{mtc} into its H₂O and CO₂ components² (K_W and K_C respectively) revealed an estimated $K_W \approx 1.5 \pm 0.5 \times 10^{-4} \text{ cm}^2 \text{ s}^{-1}$. This value was found to be approximately half that reported for a 87.2 cm³ vessel with a higher interfacial area to volume ratio, under gentle circulation ($a/V_W = 3.7$ higher [22]), and was within an order of magnitude for the transfer of most organic species in water. Hydrophobic compounds such as benzaldehyde are always expected to transfer through CO₂ faster than through H₂O, highlighting how pulsed ultrasound may be employed to enhance mass transport rates of such species through the interface, by almost two orders of magnitude.

² Once K_{mtc} was obtained from $K_{mtc}a^*$ (by inserting ' a^* ' = 71 cm^2 and $V_W = 650 \text{ cm}^3$), k_W and k_C were derived from the relation: $K_{mtc} = 1/(1/K_C + K_{CW}/K_W)$, and solved for $K_C/K_W \geq 9$.

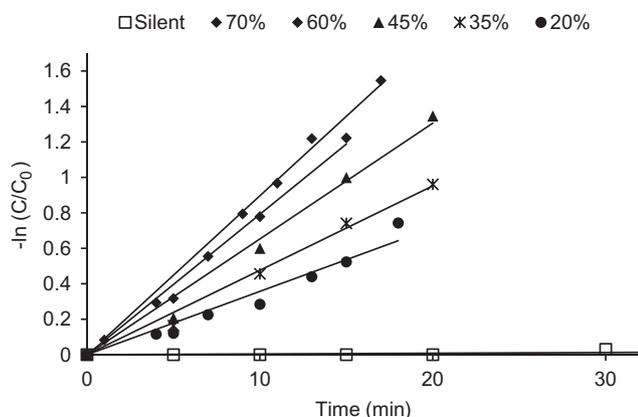


Fig. 7. Initial rate first-order plots for the hydrolysis of benzoyl chloride, under silent conditions or with sonication at different amplitude settings ($r^2 \geq 0.96$). Each value in the legend represents the amplitude setting. Conditions: $C_{CO} = 0.025$ M, $W_{CO_2} = 25\%$ (220 g CO_2 , 650 g H_2O), 30 °C/80 bar, pulse rate 25%.

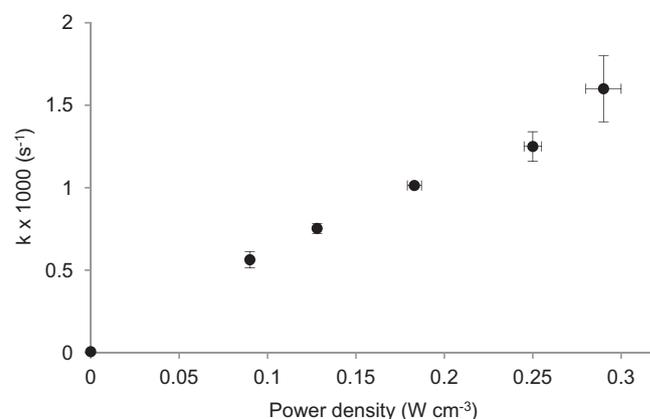


Fig. 8. Graph showing the relationship between the rate constant for benzoyl chloride hydrolysis, and ultrasound power density. Conditions: $C_{CO} = 0.025$ M, $CO_2 = 25$ wt.% (220 g CO_2 , 650 g H_2O), 30 °C/80 bar, pulse rate 25%.

3.3. Scale-up feasibility by kinetics of benzoyl chloride hydrolysis

Hydrolysis represents an important reaction and is employed in the preparation of many pharmaceutical compounds [35]. A range of amplitude settings was used to study the hydrolysis of benzoyl chloride and for the calculation of the respective rate constants (Fig. 7).

Comparison with a smaller, 87.2 cm³ reactor reported in the literature revealed that initial rates were similar for both systems at approximately equal power intensities (power intensity here was 95.5 W cm⁻² with the 1500 W sonifier and a probe tip surface area of 3.14 cm², whereas for the smaller vessel the intensity was 104 W cm⁻² with a 400 W sonifier and a probe tip surface area of 0.5 cm², Table 1, Entries 1, 2). In the study by Timko et al., no sonochemical effects by ultrasound were observed, inferring that faster hydrolysis rates were due to increased interfacial areas between CO_2 and H_2O . Further demonstration of the conservation of efficiency, was provided by observing an almost equivalent rise in the rate constant over the same range of power densities, for both vessel dimensions (rate constant increased roughly 3 times in the range 0.1–0.3 W cm⁻³, Fig. 8) [30].

Scale-up of sonochemical processes is usually problematic due to their dependence on localised cavitation events (immediately below the irradiating probe), and wide variation in the energy dissipation in the bulk volume [24]. Scale-up operations may also be hindered by the fact that the progressive erosion and shortening of the probe may eventually lead to reduced efficiency and increased process costs [36]. However, ultrasound in high-pressure, and especially near-critical, fluids, displays distinct advantages compared to sonication of fluids under ambient conditions, including

‘soft’ cavitation (therefore limiting erosion of the probe tip), and increased biphasic interfacial disruption (better dispersive and micro-mixing capacity) due to the low surface tension, high vapour density and low liquid density of CO_2 [37,38]. Furthermore, in the vicinity of a heterogeneous surface, the acoustic wave leads to the generation of physical and mechanical phenomena such as acoustic streaming and microjetting, which may be beneficial regardless of any sonochemical effect taking place [39,40]. As a consequence, two immiscible liquids will generate an emulsion, which may not be solely localised beneath the irradiating surface once formed, but as in the present case, will occupy larger fractions of the vessel volume. Factors governing ultrasound efficiency for heterogeneous systems include distance of the biphasic line from the probe (here with 650 cm³ H_2O 75 wt.%, this was 1 cm below the ultrasound probe), probe surface area, and vessel geometry. An aluminium foil test [41] revealed that significant erosion took place at 1 cm below the probe, indicating the presence of cavitation events with a circular pattern and diameter (~1.2 cm) approximately equal to that of the probe (Fig. 9).

Ultrasound-induced cavitation in liquid CO_2 has proven to be beneficial in a number of applications, including the preparation of polymers, drug nanoparticles, and ceramic materials [42–44]. In this system, hydrolysis occurred in two distinct environments, a H_2O -continuous phase existing below the probe, where benzoyl chloride-containing CO_2 droplets were constantly being formed and reacted with surrounding H_2O , and a CO_2 -continuous phase in which H_2O was constantly fluxing to react with benzoyl chloride (Scheme 2).

Table 1

Comparison of benzoyl chloride hydrolysis rates between various emulsion systems.

Entry	Conditions	$k_{hyd} (\times 10^3 s^{-1})$
1	C/W–W/C Ultrasound ^a Silent (This work)	1.6 ± 0.3
2	C/W W/C Ultrasound ^b	1.8 ± 0.2
3	C/W Ultrasound + Tween 80 (This work) ^c	6–16
4	Pre-formed W/C microemulsion ^d	0.064
5	Pre-formed O/W microemulsion ^e	4.14

^a US 0.3 W cm⁻³/95 W cm⁻², 1 dm³ US reactor, 30 °C/80 bar CO_2/H_2O .

^b US 0.6 W cm⁻³/105 W cm⁻², 87 cm³ US reactor, 30 °C/80 bar CO_2/H_2O [22].

^c Measured from the initial linear portion of the plot.

^d CO_2 /PFPE/ H_2O microemulsion 35 °C/276 bar [23].

^e Isooctane/AOT(dioctyl sodium sulfosuccinate)/ H_2O microemulsion, 25 °C/ ambient pressure [50].

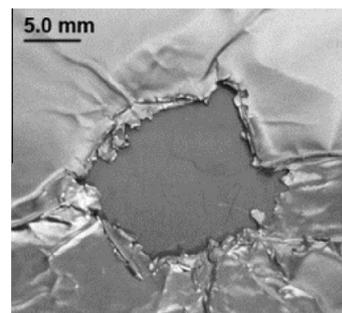
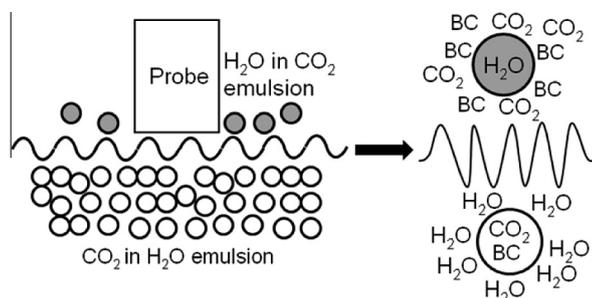


Fig. 9. Image of aluminium foil erosion following 1 min sonication in CO_2/H_2O . Four layers of foil were wrapped around a stainless steel plate positioned 1 cm below the probe. Conditions: CO_2/H_2O 50 v%, 30 °C/80 bar, 70% amplitude, 25% pulse rate.



Scheme 2. Simplified diagram showing the simultaneous formation of CO₂ in H₂O and H₂O in CO₂ environments by ultrasound (left). The reaction may proceed in both environments, by the hydrophobic compound encountering a water droplet in the bulk CO₂ phase, or by the compound-carrying CO₂ droplet interacting with the bulk water phase (right, BC = benzoyl chloride).

As expected, the reaction proceeded faster under sonication compared to silent conditions, by effectively increasing the CO₂/H₂O interfacial area. Sonication with 70% amplitude afforded the maximum yield in 30 min (Fig. 11, dashed lines)³. In the absence of sonication, the hydrolysis of benzoyl chloride was extremely slow ($t_{1/2} \approx 27$ h). The sonic enhancement factor for hydrolysis rates relative to the silent case was therefore almost 230. When assessing reaction rates in binary solvent systems, hydrophobic interactions between the solvent and the reagent must also be taken into account; in this respect, ultrasound would alter the solvent structure and solvation [45], thus affecting the activation energy and entropy values for the reaction. Using different alcohol–water compositions, Salmar et al. showed that the solvent affected the rate constants depending on the alcohol/water fraction used, by hydrophobic stabilisation of either the transition complex or the ground state of the reagent [46].

The results from this part of the study highlight how the physical effects of ultrasound may be applied to include the efficient processing of high-pressure fluids on the litre scale, and may provide scope for identifying specific reaction pathways that may benefit from emulsification of H₂O-containing heterogeneous systems [47]. Furthermore, in this system, the reaction between CO₂ and H₂O formed a spontaneous acidic environment (pH ≤ 4), potentially acting as an intrinsic catalyst that tends to neutral pH following CO₂ venting.

3.4. Emulsion formation using Tween 80

Sonication in the presence of 1.0 wt.% Tween 80 generated a uniform, white emulsion (Fig. 10). Analysis of samples taken from the upper, CO₂-continuous section of the reactor, following injection of 1 mL octane (a hydrophobic tracer that preferentially partitions in CO₂ [29]), showed a 2.5-fold increase in the concentration of octane during sonication that reverted to the biphasic when sonication was ceased.

Additionally, the quantity of water present in these samples disappeared gradually following termination of ultrasound. Quantification of the water mass collected, in combination with octane concentrations, suggested that the emulsion consisted of $35 \pm 2\%$ CO₂ and $70 \pm 5\%$ H₂O by volume, which gradually reverted to a separated biphasic approximately 1 h after ceasing sonication. Comparison of samples from the upper and lower sections of the reactor showed similar compositions, indicating uniformity throughout the reactor.

The hydrolysis reaction was next assessed in the presence of Tween.

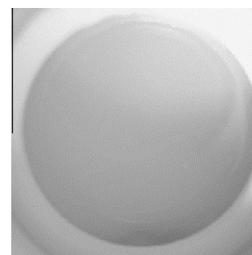


Fig. 10. Tween 80-stabilised (1.0 wt.%) emulsions generated by ultrasound. Conditions: CO₂ = 25 wt.% (220 g CO₂, 650 g H₂O), 30 °C/80 bar, after 10 min of ultrasound at 70% amplitude, 25% pulse rate.

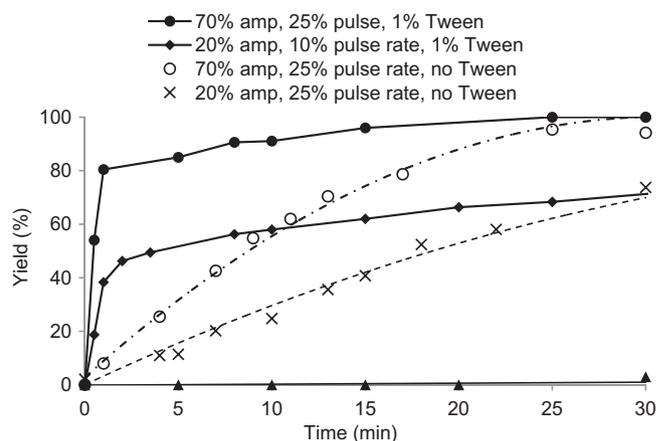


Fig. 11. Benzoyl chloride hydrolysis in Tween-stabilised (1.0 wt.%) C/W emulsions under different ultrasound conditions. Similar rates and yields were obtained with 0.5 wt.% Tween 80. Conditions: C₀ = 0.025 M, CO₂ = 25 wt.% (220 g CO₂, 650 g H₂O), 30 °C/80 bar.

Studies revealed that a combination of ultrasound and Tween was an effective environment for producing fast conversions of benzoyl chloride to benzoic acid (Fig. 11, solid lines).

Tween possesses an amphiphilic structure, and has been shown to stabilise C/W emulsions exclusively (even at 90 wt.% CO₂ for concentrated emulsions), with mean droplet diameters of 0.5–5.4 μm depending on the shear rates through a sand pack [31]. Confirmation of the C/W nature of the emulsion was obtained by injecting benzoyl chloride into the pre-formed emulsion, without further sonication, and analysing samples from both the upper and lower sections of the reactor. If the emulsion proved to be CO₂-continuous (W/C), substantial dissolution of either benzoyl chloride, benzoic acid, or both, would have occurred. The samples, however, contained neither species implying that following injection, benzoyl chloride encountered a H₂O-continuous phase, and due to its density (1.21 g cm⁻³), resided at the bottom of the vessel without reacting. Tween-stabilised hydrolysis rates under ongoing sonication showed two distinct stages, the first achieving ~80% yield almost instantly, and a second showing slower conversion, reaching 100% yield after approximately 30 min. The rate of the Tween-stabilised hydrolysis could be controlled by manipulating the pulse rate. Pulsed ultrasound has distinct advantages over continuous ultrasound, such as an increase in the number of spatial reactive sites and residual pressure amplitude during the OFF stage of the cycle, and enhanced energy efficiency by decreased power draw [48,49]. Pulse rate may therefore be regarded as a valuable variable for process optimisation. Using 10% pulse rate and 20% amplitude, 100% yield of benzoic acid was reached within ~120 min, significantly slower than when higher pulse rates were

³ For all yields after 30 min, see supplementary data.

employed. Observation through the view cell revealed that turbidity was of a lower intensity and lagged by several minutes in reaching a maximum, suggesting that the emulsion consisted of a lower surface area during low pulse rate. At steady-state, assuming the highest mean CO₂ droplet diameter of 5 μm (as reported under conditions of shearing), and 35% volume of the dispersed phase, the total emulsion interfacial area was estimated⁴ at 420 m² or 480 m² kg⁻¹.

Under the same conditions without Tween, surface areas for ultrasound-generated C/W and W/C emulsions were reported as 20 and 100 m² kg⁻¹, respectively, making the Tween-stabilised emulsion surface area at least four times higher [22]. However, surface area may not independently account for the large increase observed in hydrolysis rates. Additional ultrasound-induced effects such as convective forces on existing droplets, may define mass transport and reaction rates, with droplet size playing an important factor in droplet dynamics. In support of this, benzoyl chloride hydrolysis in a pre-formed W/C emulsion was much slower (Table 1, Entry 4).

Another advantage of using a surfactant is improved energy efficiency, as demonstrated by experiments using different pulse rates. After 5 min of sonication at 20% amplitude, inclusion of surfactant yielded a fivefold higher concentration in benzoic acid with a 2.5-times lower energy requirement (2700 J with 10% pulse rate compared to 6750 J with 25% pulse rate for sonication with and without surfactant, respectively). A combination of low pulse rate and surfactant may therefore offer a useful processing strategy for minimising energy expenditure without compromising the yield.

4. Conclusions

This study highlights two important aspects regarding the utilisation of ultrasound in high-pressure systems. First, processes that rely on the physical effects of sonication and lead to dispersion of heterogeneous phases may be safely scaled-up (over 11-fold in the present study) without loss of efficiency, due to less dependence on cavitation and the sonolytic effect which may quickly dissipate from the irradiating surface in larger vessels. Second, the effect of ultrasound may be complemented by including low quantities of a bio-compatible surfactant (as low as 0.5 wt.%), resulting in complete emulsification of the biphasic mixture, which maintains a defined C/W nature and uniformity throughout the reactor. It is therefore easy to envisage how the nature of the environment may be best suited to different applications either: (i) ongoing reaction/separation (without Tween two distinct environments co-exist, allowing preferential partitioning in the CO₂ and H₂O phases in real time), or (ii) complete emulsification forming only one type of environment (with Tween, achieving greater uniformity and surface area).

Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at <http://dx.doi.org/10.1016/j.ultsonch.2013.05.017>.

⁴ Surface area values were calculated according to the formula: $a = 6\phi / d_{32}$, where 'a' is the surface area (m²), ϕ is the volume fraction (%) of the dispersed phase (or here simply volume, m³), and d_{32} is the Sauter mean diameter of the droplet (m). The specific surface area (m² kg⁻¹) was obtained with respect to the total emulsion mass (0.87 kg).

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