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Phenol and benzenoid alcohols separation from aqueous stream using cloud point extraction: Scaling-up of the process in a mixer-settler

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A B S T R A C T

In the present work, the cloud point extraction (CPE) of three organic pollutants (phenol, benzyl alcohol and 1-phenylethanol) with aqueous solutions of biodegradable alkoxyated nonionic surfactants (TERGITOL 15-S-7 and SIMULSOL NW342), is investigated. First, the partial phase diagrams of the water-surfactant binary systems are established. Then, the effects of organic pollutants and sodium chloride on the cloud point (T_c) are determined. Extraction efficiency is evaluated by the following responses: percentage of solute extracted, E (%), residual concentrations of solute and surfactant in dilute phase ($X_{s,w}$ and $X_{t,w}$, respectively) and volume fraction of coacervate at equilibrium (ϕ_c). Three-dimensional empirical correlations are used for fitting the experimental results. The comparison between experimental and calculated values allows model parameter identification. Based on these data, CPE was implemented in a continuous mixer-settler device. The feasibility of a multi-stage crossflow process for the purification of samples containing phenol using SIMULSOL NW342 was tested. Six stages were required to reduce the pollutant concentration below the allowed level (0.3 ppm), which proves the efficiency of CPE in the treatment of wastewaters.

1. Introduction

From the wide variety of organic pollutants, phenol, benzyl alcohol and 1-phenylethanol were chosen for this study. Phenol is considered as one of the major water pollutants. Even at very low doses, it is still very dangerous because of its persistence, toxicity, ecotoxicity and bioaccumulation [1–5]. The Environmental Protection Agency (EPA) and the French legislation limit its allowed concentration in surface waters to 0.3 mg/L [6,7]. For phenol removal, solvent extraction, adsorption, but also ion exchange, polymerization, electro-coagulation, membrane-based separations and biological methods have been found effective [1,4,8–10]. In soap, perfume and flavor industries, benzyl alcohol is employed as such in bar soap fragrances and in the form of its esters. It is also used in the polymer industry and in the manufacture of car tires. Its photocatalytic degradation has been investigated [11]. 1-Phenylethanol is mainly a coproduct of the oxidation step of ethylbenzene, whose hydroperoxide is used to convert propylene to its oxide. It is then valorized through dehydration to styrene. It can be burnt in a chemical incinerator equipped with a post combustion and epuration system [12].

The laws and regulations on wastewater treatment are becoming increasingly strict. Therefore, there is a strong trend to develop efficient

methods for the removal and/or recovery of toxic species in the environment [13]. Among others, Cloud Point Extraction (CPE) appears to be a relatively simple and ecologically safe technique. In fact, the aqueous solutions of most polyethoxylated nonionic surfactants become cloudy and start to separate into two phases, coacervate and dilute phase, as soon as temperature rises above their cloud point, T_c [14]. This phenomenon is the basis of the CPE process [15–20]. This latter avoids the use of an organic solvent, produces small sludge volume and requires low energy consumption. This process is very efficient for treating water containing various contaminants including dissolved or dispersed organic matter [21–40]. This method of water purification was also applied to the extraction of metal ions using diverse appropriate chelates [15,41–44] and without chelates [45–49]. Associated to the cloud point value, the main factors in surfactant selection are its biodegradability, toxicity and ecotoxicity. The use of CPE offers an interesting alternative to conventional extraction systems. This technique allows moving toward Green Chemistry. Many advantages were claimed to CPE compared with conventional liquid-liquid extraction: CPE is an efficient and selective process that works continuously, saves energy and can be scaled up [50–54]. On the basis of this finding, the batch CPE of phenol, benzyl alcohol and 1-phenylethanol from aqueous solution was investigated in the present work. The effects of

Nomenclature

Symbols

BA	Benzyl alcohol
cmc	Critical micelle concentration
E	Extent of extraction (%)
PE	1-phenylethanol
PH	Phenol
T _c	Cloud point (temperature)(°C)

V _r	Stirrer speed
X _{s,w}	Mass fraction of solute in the dilute phase after extraction
X _{t,w}	Mass fraction of surfactant in the dilute phase after extraction
X _t	Initial surfactant mass fraction
X _{PH,F} , X _{PH,S} , X _{PH,E} , X _{PH,R}	Mass fraction of solute in the feed, solvent, extract and raffinate phase, respectively
X _{t,F} , X _{t,S} , X _{t,E} , X _{t,R}	Mass fraction of surfactant in the feed, solvent, extract and raffinate phase, respectively
φ _c	Volume fraction of coacervate

temperature, surfactant concentration and decantation time on solute extraction extent were also studied. For this purpose, two polyalkoxylated nonionic surfactants were used: TERGITOL 15-S-7 and SIMULSOL NW342. The initial pollutant concentration used in water was 0.2 wt.%. Using these data, the continuous multi-stage crossflow cloud point extraction was implemented in a mixer-settler. As a model system, phenol was extracted using SIMULSOL NW342. The mixer-settler can be easily arranged in battery for counter-current multi-stage process. Each mixer-settler couple can be considered as a theoretical stage. This equipment offers the advantage to operate with highly unequal phase fractions. This technology is able to operate with high flow rates, and can be useful for wastewater treatment.

2. Materials and methods

2.1. Chemical species

The nonionic surfactants used in this work were obtained from Oxo-alcohol alkoxylation: SIMULSOL NW342 (cmc = 1.52 mmol/L at 15 °C), kindly provided by SEPPIC (Castres, France) and TERGITOL 15-S-7 (cmc = 1.22 mmol/L at 20 °C), a Dow Chemical specialty purchased from Aldrich, are mixtures of primary and secondary alcohol alkoxyates with the alcohol group located at various positions along the carbon chain. Phenol, benzyl alcohol and 1-phenylethanol were purchased from Aldrich and sodium chloride from VWR. The formulas and some properties of the species used in this work are listed in Table 1. Deionized water was used in all cases except for the HPLC analyses, carried out with ultrapure water.

2.2. Methods

2.2.1. Cloud point measurements

Cloud point measurements were carried out using a Mettler FP 900 apparatus. It consists of a FP90 control and operating unit, and a FP81C measuring cell dedicated to cloud point measurements. The cell temperature was measured with a Pt100 sensor; light transmission was measured continuously, while the cell temperature was increasing linearly according to the chosen heating rate. The cloud point corresponds to the temperature at which the limpid phase becomes cloudy, inducing a light transmission decrease.

2.2.2. Experimental conditions

For batch extraction tests, 30 mL of solution, containing the surfactant (at concentrations from 2 to 10 wt.%) and the solute (0.2 wt.%) in demineralized water, were poured into graduated cylinders and heated in a precision oven and kept during 24 h to reach equilibrium. The volumes of both phases (coacervate and dilute) were measured.

The dilute phase was analyzed. The residual pollutant and surfactant concentrations were determined by reversed-phase high-performance liquid chromatography: for the solutes, the conditions were as follows: column RP18 (ODS), 95 bar, mobile phase H₂O/CH₃CN/CH₃OH, 42.5/50/7.5 (v/v), flowrate 1 mL/min.; λ = 260 nm; t = 25 °C. The conditions were slightly different for the surfactant:

mobile phase H₂O/CH₃CN/CH₃OH, 7.5/60/32.5 (v/v). The sensitivity of the evaporative light-scattering detector (DDL 31, EUROSEP Instruments) was optimized by the control of the air flowrate in the atomizer (relative pressure: 1 bar), the temperature of the evaporator (55 °C) and the gain of the photomultiplier (400 mV) [59–61].

The total capacity of the mixer-settler (Fig. 1); tentatively used by our research group in a previous work [21], was 9.5 L. However, the occupied volume was 7 L. The stirrer diameter was 6.5 cm and the mixing tank diameter was 8.5 cm. The cylindrical settler was 98 cm length and 10 cm diameter; the stirring speed could vary from 0 to 900 rpm. For the ternary water/Simulsol NW342/phenol system, the equipment was operated with equal volumes of the feed solution F (0.4 wt.% phenol) and the solvent S (8 wt.% surfactant); the mixture was maintained at 30 °C.

2.2.3. Extraction parameters

In order to find the optimal conditions of the two variables: wt.% surfactant (X_t), and temperature (T), allowing to obtain the best possible extraction results, we have worked out the best compromise between the four “responses”, Y (E, X_{s,w}, X_{t,w} and φ_c), defined as follows:

- The extraction yield E (%):

$$E(\%) = \frac{m_{s(in)} - m_{s(w)}}{m_{s(in)}} \times 100 \quad (1)$$

Where m_{S(in)} and m_{S(w)} represent the mass of the solute in the initial solution and in the dilute phase, respectively.

- The volume fraction of coacervate, i.e. the ratio of the volume of the coacervate, V_c, to the total volume (V_c + V_w), V_w being the volume of the dilute phase:

$$\phi_c = \frac{V_c}{V_c + V_w} \quad (2)$$

- The weight percentage of solute in the dilute phase:

$$X_{s,w}(\%) = \frac{m_{s(w)}}{m_w} \times 100 \quad (3)$$

with m_w: mass of dilute phase,

The weight percentage of surfactant in the dilute phase:

Table 1
Chemicals: formulas and properties.

Name (abbreviation)	Formula	Aqueous solubility at 25 °C (g/L)	log P [58]
SIMULSOL NW342 (Oxo-C ₁₀ E ₃ P ₄ E ₂)	C ₁₀ H ₂₁ -(OCH ₂ -CH ₂) ₃ -(O-CH ₂ -CH(CH ₃)) ₄ -(OCH ₂ -CH ₂) ₂ -OH		
TERGITOL 15-S-7 (C ₁₁ -15E _{7,3})	C ₁₅ H ₃₁ -(O-CH ₂ -CH ₂) _{7,3} -OH		
Phenol (PH)	C ₆ H ₅ OH	82.8 [55]	1.5
Benzyl alcohol (AB)	C ₆ H ₅ CH ₂ OH	40 [56]	1.05
1-Phenylethanol (PE)	C ₆ H ₅ CH(CH ₃)OH	20 (at 20 °C) [57]	1.42

Fig. 1. Mixer-settler device.

Fig. 2. The cloud point of nonionic surfactants.

Fig. 3. Effect of solutes on the cloud point of SIMULSOL NW342 and TERGITOL 15-S-7.

$$X_{t,w}(\%) = \frac{m_{t(w)}}{m_w} \times 100 \quad (4)$$

with m_{bw} : mass of surfactant in dilute phase.

Those four criteria can be sufficient to evaluate the performances of a single contact extraction.

3. Modeling of results

By using the two variables: wt.% surfactant (X_t), and temperature (T), the extraction results of the three solutes at 0.2 wt.% by the two surfactants (TERGITOL 15-S-7 and SIMULSOL NW342) were expressed by four responses (Y): percentage of extracted solute (E), residual concentrations of solute and surfactant in dilute phase $X_{s,w}$, and $X_{t,w}$, respectively, and coacervate volume fraction at equilibrium (ϕ_c). For each parameter determined by considering central composite designs, the results were analyzed using an empirical fitting method. [36,62,63].

Fig. 4. Effect of NaCl on the cloud points of SIMULSOL NW342 and TERGITOL 15-S-7.

The experimental values could be used to determine the quadratic polynomial model constants, which were adjusted. The models were checked by plotting computed data against experimental results. The nonlinear system formed by the equations:

$$Y = a_0 + a_1 X_t + a_2 T + a_{12} X_t T + a_{11} X_t^2 + a_{22} T^2 \quad (5)$$

was solved to give the slope and the regression coefficient (R^2) closest to unity. Even if the quadratic equations for E , $X_{s,w}$, $X_{t,w}$ and ϕ_c allow to obtain a good correlation with the experimental values, one has to keep in mind that the model validity is restricted to the field of study and a physical significance cannot be given to the portion of horizontal planes corresponding to the maximal values of the response.

Coefficient identification and data treatment of response surfaces were carried out using the STATISTICA software (8.0.360.0).

4. Results and discussion

4.1. Binary and pseudo-binary phase diagrams

Before the extraction test, it is necessary to plot the phase diagrams of the following systems: water/surfactant and water/surfactant with a fixed solute content (pseudo-binary systems) as a function of temperature. One can notice in Fig. 2 that the cloud points of TERGITOL are higher than those of SIMULSOL NW342. Thus, the critical temperature of this latter ($t_c = 17^\circ\text{C}$) is lower than the ambient one, while that of TERGITOL 15-S-7 is above 34°C . This might have been difficult to predict, since the surfactants do not belong to the same series, and TERGITOL 15-S-7 possesses longer hydrophobic tail and hydrophilic head at the same time. Thus, the effect of the four propylene oxide groups of SIMULSOL NW342 prevails over the shortening of the aliphatic chain. The cloud-point curves are more sensitive to surfactant concentration in a narrow range of lower values.

4.1.1. Effect of organic solutes

The effect of the three organic solutes phenol, benzyl alcohol and 1-phenylethanol, in the range 0–2 g/L, on the cloud points of SIMULSOL NW342 and TERGITOL 15-S-7 has been investigated, as an example of solute effect. For $X_t = 4$ wt.%, the results are shown in Fig. 3. One can notice that very small concentrations of the three additives have no effect on the cloud point. However, at higher concentration, those compounds lower the cloud points of both nonionic surfactants. In each case, the trend is similar: phenol shows the strongest effect, whilst benzyl alcohol the lowest. Besides, the effect of solutes is relatively higher at lower surfactant concentration.

The influence of solutes on the cloud point is governed by their hydrophobic affinity, namely their octanol-water partition coefficients, expressed as $\log P$ (Table 1). Although phenol is more soluble in water

Fig. 5. Three-dimensional isoresponse curves smoothed by a quadratic model, $E(\%) = f(X_i, T)$, (Eqs. (6), (7), (14), (15), (22) and (23)).

than aromatic alcohols, it may be considered as more hydrophobic and interacting more strongly, through hydrogen bonding, with ethylene oxide units. This was already noticed elsewhere [36,64].

The cloud point curve lowering caused by phenol, has also been observed with other surfactants [18,21,22]. This phase diagram relative to a given solute can be used as a qualitative guide for the extraction

process.

4.1.2. Effect of sodium chloride

Electrolytes can lower or raise the cloud point of nonionic surfactant systems [65]. Fig. 4 shows the decrease of the cloud point for 2 and 4 wt.% of each surfactant, by addition of sodium chloride. It can be

Fig. 6. Three-dimensional isoresponse curves smoothed by a quadratic model, $X_{s,w} = f(X_t, T)$, (Eqs. (8), (9), (16), (17), (24) and (25)).

observed that NaCl affects TERGitol 15-S-7 more than SIMULSOL NW342. This phenomenon is known as the salting-out effect, caused by weakening of hydrogen bonds between the polar head groups of the nonionic surfactants and water molecules. This observation has been made with many other micellar systems [65–68]. Therefore, it is possible to adjust the cloud point at room temperature by a simple control

of surfactant and salt concentrations, and consequently reduce the heating energy cost in the extraction process. Salt addition can also have an application in the extraction of heat sensitive products.

Fig. 7. Three-dimensional isoresponse curves smoothed by a quadratic model, $X_{l,w} = f(X_t, T)$, (Eqs. (10)–(12), (19), (26) and (27)).

4.2. Extraction results

The CPE of 0.2 wt.% of each solute (phenol, benzyl alcohol or 1-phenylethanol) with SIMULSOL NW342 or TERGITOL 15-S-7 was performed. The results were obtained for initial surfactant concentrations, (X_t) ranging from 2 to 10 wt.%, and in temperature ranges 26–42 °C

with SIMULSOL NW342 and 3–46 °C with TERGITOL 15-S-7.

The equations relative to the different solutes (pollutants) are:

- Phenol

$$E_{NW342} = 35.32 + 12.24 x + 0.15 T - 0.11 xT - 0.39x^2 + 0.4410^{-2}T^2 \quad (6)$$

Fig. 8. Three-dimensional isoresponse curves smoothed by a quadratic model, $\phi_c = f(X_t, T)$, (Eqs. (12), (13), (20), (21), (28) and (29)).

$$E_{15-S-7} = -292.27 - 10.14 x + 17.12 T + 0.15 xT - 0.69x^2 - 0.22T^2 \quad (7)$$

$$X_{s,w,NW342} = (144.49 - 23.89 x - 0.88 T + 0.19 xT + 0.87x^2)10^{-3} \quad (8)$$

$$X_{s,w,15-S-7} = 1.30 + (58.42 x - 60.58 T - 1.93 xT + 1.76x^2 + 0.78T^2)10^{-3} \quad (9)$$

$$X_{t,w,NW342} = 31.53 - 0.42x - 1.40T - (1.41 xT - 13x^2 - 1.91T^2)10^{-2} \quad (10)$$

$$X_{t,w,15-S-7} = 11.68 + 1.80 x - 0.38T - (3.87x.T - 1.28x^2 - 0.50T^2)10^{-2} \quad (11)$$

$$\phi_{CNW342} = (94.815 + 40.64 x - 4.52 T - 1.15 xT + 1.98x^2 + 0.08T^2)10^{-3} \quad (12)$$

$$\phi_{C15-S-7} = -1.14 + 0.18 x + (55.08 T - 4.17 xT + 5.21x^2 - 0.62T^2)10^{-3} \quad (13)$$

- Benzyl alcohol

$$E_{NW342} = -69.41 - 1.71 x + 6.12 T + 0.13xT + 0.08x^2 - 0.08T^2 \quad (14)$$

$$E_{15-S-7} = 305.13 + 10.61 x - 12.46 T - 0.01 xT - 0.41x^2 + 0.14T^2 \quad (15)$$

$$X_{s,w,NW342} = 46.90 + 1.40 x + 1.10 T + 0.03 xT - 0.08x^2 - 0.01T^2 \quad (16)$$

Fig. 9. Batch cross-flow extraction results after $t = 135$ min.

$$X_{s,w,15-S-7} = 0.31 + (24.62 x - 9.82 T - 0.71 xT + 0.21x^2 + 0.14T^2)10^{-3} \quad (17)$$

$$X_{t,w,NW342} = 24.75 - 0.87x - 0.84 + (16x^2 \pm 1.04T^2)10^{-2} \quad (18)$$

$$X_{t,w,15-S-7} = 64.19 + 0.74 x - 2.71 T - (1.25x \cdot T - 1.15x^2 - 3.09T^2)10^{-2} \quad (19)$$

$$\phi_{CNW342} = 0.21 + (13.53 x - 8.75 T - 1.35 xT + 5.83x^2 + 0.15T^2)10^{-3} \quad (20)$$

$$\phi_{C15-S-7} = 3.51 + 0.21 x - 0.16 T - (0.21 xT + 0.44x^2 - 0.19T^2)10^{-2} \quad (21)$$

- 1-Phenylethanol

$$E_{NW342} = -92.21 + 9.09 x + 6.26 T + 0.14 xT - 0.83x^2 - 0.09T^2 \quad (22)$$

$$E_{15-S-7} = -369.29 + 21.19 x + 16.52 T - 0.26 xT - 0.41x^2 - 0.17T^2 \quad (23)$$

$$X_{s,w,NW342} = 0.42 - (13.24 x + 14.91 T + 0.46xT - 1.82x^2 - 0.21T^2)10^{-3} \quad (24)$$

$$X_{s,w,15-S-7} = 1.21 + 0.01 x - 0.05 T - (0.07 xT - 0.12x^2 - 5.40T^2)10^{-2} \quad (25)$$

$$X_{t,w,NW342} = 33.46 + 1.22 x - 1.59 T - (6.37 xT - 17x^2 - 2.37T^2)10^{-2} \quad (26)$$

$$X_{t,w,15-S-7} = 61.21 + 0.82 x - 2.59 T - (1.50x \cdot T - 1.40x^2 - 2.96T^2)10^{-2} \quad (27)$$

$$\phi_{CNW342} = 0.18 + (56.47 x - 13.37 T - 1.56 xT + 1.46x^2 + 0.26T^2)10^{-3} \quad (28)$$

$$\phi_{C15-S-7} = 0.46 + 0.28 x - (30.17 T + 4.58 xT + 1.25x^2 - 0.42T^2)10^{-3} \quad (29)$$

4.2.1. Extraction efficiency (E)

The influence of surfactant concentration and temperature on the extraction percentage of organic solutes has already been studied [36]. As expected, the extraction efficiency increases with non-ionic

surfactant concentration. On the other hand, an excessive increase of temperature above the cloud point can have negative effect on separation results.

According to the quadratic model (Eqs. (6), (7), (14), (15), (22) and (23)), the most favorable extraction conditions are expected within the region of higher surfactant concentration (*i.e.* 10 wt.%, Fig. 5). Indeed, if TERGITOL 15-S-7 is known for its high solubilization ability and extraction efficiency [25,69,70], its performances are found similar to those of SIMULSOL NW342 ($E_{\max} \geq 90\%$).

4.2.2. Concentration of residual solute ($X_{s,w}$)

The three-dimensional isoresponse curves of the studied properties fitted to the quadratic model (Eqs. (8), (69), (16), (17), (24) and (25)) are given in Fig. 6. These figures show that the percentage of residual solute in the dilute phase, $X_{s,w}$, decreases as X_t increases. On the other hand, its dependence upon tempera

ture is not monotonous, although the model gives a good illustration of experimental results. In general, a slight temperature rise has a positive effect on the extraction.

4.2.3. Residual concentration of nonionic surfactant ($X_{t,w}$)

Even if the nonionic surfactants used in this study are known for their good biodegradability properties and lack of ecotoxicity [71,72], the concentration of residual surfactant ($X_{t,w}$) still remains a very important parameter. The high loss of surfactant in the dilute phase can compromise the process reliability. Indeed, the presence of another contaminant in the dilute phase is sufficient to make the process useless. The behaviour of ($X_{t,w}$) according to X_t and T is shown in Fig. 7, (fitting by the quadratic model Eqs. (10), (11), (18), (19), (26) and (27)). These figures show that the residual concentration of surfactant decreases at low surfactant concentration and rises according to temperature. The highest concentration cannot be reached with a unique surfactant concentration for the three solutes. It is equal to 10 wt.% and the temperatures are 26 °C and 38 °C for SIMULSOL and TERGITOL, respectively.

4.2.4. Volume fraction of coacervate (ϕ_c)

This factor is very important to show the effectiveness of this technique. In Fig. 8 (fitting with Eqs.(12), (13), (20), (21), (28) and (29)), it is shown that a temperature rise allows to lower the volume fraction of the coacervate. It is clear that, for all solutes studied, the values and the evolution of the volume fraction of coacervate are practically the same, evidencing solutes do not affect significantly the coacervate volume, under these conditions. According to the shape of the phase diagram, the overall composition decreases when moving away from the right branch of the demixing curve. As a consequence, it is advantageous to use less surfactant in order to have a smaller coacervate volume.

4.3. Batch cross-flow cloud point extraction

In a batchwise preliminary study of a cross flow extraction process, a 0.2 wt.% phenol solution was extracted in test tubes under the following conditions: $X_t = 4\%$, $t = 35$ °C. With a 135 min contact

Fig. 10. Effect of stirring speed on: left: surfactant concentration in dilute phase (raffinate) and coacervate (extract); right: solute concentration in dilute phase (raffinate) and coacervate (extract); System: water/4% SIMULSOL NW342/0.2% phenol, $T = 35$ °C.

Fig. 11. Extraction parameters of the system: water/4% SIMULSOL NW342/0.2% phenol at 30 °C; a) Volume fraction of coacervate (ϕ_c); b) Extraction efficiency (E).

Table 2
Continuous crossflow cloud point extraction parameters.

Input		Output	
$F = 19.5(\text{g/min})$	$S = 19.5(\text{g/min})$	$E = 5.6$	$R = 33.4$
$X_{PH,F} = 0.078$	$X_{PH,S} = 0$	$X_{PH,E} = 0.054$	$X_{PH,R} = 0.024$
$X_{t,F} = 0$	$X_{t,S} = 1.560$	$X_{t,E} = 1.551$	$X_{t,R} = 0.009$

duration (equilibrium state supposedly reached, each stage corresponding to a theoretical plate), the number of extractions necessary to lower the residual phenol concentration to the allowed value is 6 (Fig. 9). In fact, at the last output, $X_{s,w} = 0.26 \text{ mg/L}$. For a 15 min contact, the number of stages rises to 9.

4.4. Continuous cross-flow cloud point extraction

Fig. 10 shows that high mixing speed induces a slight increase of surfactant and solute concentration in coacervate phase. Thereby, surfactant and solute concentration decrease in the dilute phase. Indeed, good phase separation was observed with this system at 35 °C. However, according to Ingram et al. [54], the increase of agitation speed greatly raises the surfactant concentration in the raffinate (dilute phase). This was also observed with our system at very high stirring speed (over 800 rpm) and high extraction temperatures ($T > 40 \text{ °C}$).

As seen previously, the cloud point extraction offers good results. However, those data were obtained after 24 h of settling time. Hence, for scaling-up of the CPE process, it is interesting to study the kinetics of CPE. Therefore, separation time was evaluated by following the evolution of the coacervate volume fraction (ϕ_c) and extraction extent (E). The feed and solvent solutions were introduced simultaneously in the “mixer” at $V_r = 300 \text{ rpm}$. One can notice in Fig. 11 that ϕ_c increases rapidly in the early hours, then stabilizes on a plateau after about 3–4 h of settling time. The evolution of the extraction percentage shows equilibrium after 3 h. At this time the maximum of solute was extracted. The parameters of continuous cross-flow CPE are summarized in Table 2. The volume fraction of coacervate was 0.143. The composition of the dilute phase (raffinate) was 0.070% phenol and 0.028% SIMULSOL NW342, the composition of coacervate phase (extract) was 0.93% phenol and 27.4% SIMULSOL NW342. Let us remember that at equilibrium (as determined in test tube), the extraction percentage

reached 73%, while, using the mixer-settler, a value of 69% was obtained. This result agrees with the hypothesis that a mixer-settler stage may be considered as a theoretical stage.

For a multi-stage contact with thermostatic pilot implemented previously, all stages were operated at the same temperature (30 °C) for the system water/4 wt.% SIMULSOL NW342/0.2% phenol. After the first stage contact, the raffinate phase was contacted with a new solvent quantity (Fig. 12).

The cross-flow stage number N needed to achieve a specified raffinate composition, X_n , is given by the following formula:

$$N = \frac{\text{Log}\left(\frac{X_f}{X_n}\right)}{\text{Log}\left(\frac{KS}{F+1}\right)} \quad (30)$$

The partition coefficient $K = 4.7$ and extraction solvent ratio (S/F) are kept constant. Upon seeing the first three stages results, it was assumed that the remaining stages would show the same behaviour ($E \approx 69\%$). Those results allowed us to estimate the mass balance of each stage. Thereby, the stage number which allows less than 0.3 ppm of phenol in raffinate, was equal to 6 (Fig. 13).

Coacervate regeneration for surfactant reuse, by pH changes and salt precipitation, has been addressed and operated successfully with very similar systems [21,36].

5. Conclusion

Cloud point extraction (CPE) with nonionic surfactants has become a very attractive separation technique in recent years. The objective of this study was to develop a simple, practical CPE with selected biodegradable commercial nonionic surfactants. Two alkoxyated species, SIMULSOL NW342 and TERGITOL 15-S-7 were chosen to extract phenol, benzyl alcohol and 1-phenylethanol. Those organic solutes lower the surfactant cloud points, phenol showing the strongest effect. A reduced number of experiments, thanks to a central composite design, gives an insight on the trends followed by the extraction percentage, the residual solute and surfactant concentrations in the dilute phase, and the coacervate volume fraction, but the choice of the optimal conditions requires a compromise between all these parameters. An in-depth analysis of the isoresponse surfaces leads us to recommend: $X_{NW342} \approx 4\text{--}6 \text{ wt.}\%$; $30 \text{ °C} < T < 35 \text{ °C}$, and $X_{15-S-7} \approx 4\text{--}6 \text{ wt.}\%$;

Fig. 12. Cross-flow extraction process for N stages.

Fig. 13. Stage number for continuous cross-flow extraction for the system: 4 wt.% SIMULSOL NW342/0.2% phenol at 30 °C in the mixer-settler.

40 °C < T < 44 °C, as the most favorable initial surfactant concentration and temperature ranges. A batch cross flow CPE experiment with SIMULSOL NW342 and phenol yielded encouraging results. Therefore, a continuous cross flow CPE was developed and showed that the allowed residual pollutant concentration could be reached after 6 stages. Our results thus show that SIMULSOL NW342 can be chosen in a first approach of a multistage crossflow CPE process.

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