

INFLUENCE OF RIPENING AND CREAMING ON THE AGGREGATION RATE OF DODECANE-IN-WATER NANOEMULSIONS. IS CREAMING RATE AN APPROPRIATE MEASURE OF EMULSION STABILITY?

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SUMMARY

The behavior of four oil-in-water (O/W) ionic nanoemulsions composed of dodecane, and mixtures of dodecane with squalene and tetra-chloro-ethylene, is studied. The nanoemulsions were stabilized with sodium dodecyl sulfate (SDS). The

variation of the turbidity and the average radius of the emulsions were followed as a function of time. The results illustrate the shortcomings of characterizing the stability of emulsions by their creaming rate.

According to the Laplace equation (Evans and Wennerström, 1994), the internal pressure of a drop of oil suspended in water is directly proportional to its interfacial tension γ and inversely proportional to its radius (R_i). The interfacial tension originates a difference between the chemical potential of the molecules of oil inside the drop and those of an unbounded bulk oil phase. This additional free energy is equal to $(4\pi R_i^2)\gamma/N_{m,i}$, where $N_{m,i}$ is the number of molecules of drop 'i' ($(4\pi R_i^3)\rho_o N_A/3MW$ (where ρ_o : density of the oil, MW: its molecular weight, and N_A : Avogadro's number (6.02×10^{23} molec/mol)). Hence, the referred difference in the chemical potentials is equal to

$$\Delta\mu \approx \frac{3\gamma V_M}{R_i} \quad (1)$$

where V_M : molar volume of the oil. Eq. 1 indicates that the molecules of oil which belong to drops of different sizes have distinct chemical potentials. This promotes the diffusive transfer of oil through the water phase, a phenomenon known as Ostwald ripening. The theory of Lifshitz, Slesov and Wagner (LSW; Lifshitz and Slesov, 1961; Wagner, 1961), predicts that the ripening rate can be quantified in terms of the linear increase of the cubic critical radius of the emulsion as a function of time:

$$V_{OR} = dR_c^3/dt = 4\alpha D_m C(\infty)/9 \quad (2)$$

where R_c , D_m , $C(\infty)$ and α stand for the critical radius of the dispersion, the diffusion constant of the oil molecules, their bulk solubility in the presence of a planar Oil/Water (O/W) interface, and the capillary length defined as

$$\alpha = 2\gamma V_M / \tilde{R}T \quad (3)$$

where \tilde{R} : universal gas constant, and T : absolute temperature. Finsy (2004) demonstrated that the critical radius of the dispersion is equal to its number average radius (R_a):

$$R_c = R_a = \frac{1}{N_T} \sum_k R_k \quad (4)$$

where N_T : total number of drops. According to LSW, drops with radii smaller than the critical radius decrease in size, and vice-versa. Since the critical radius changes as a function of time, the theory predicts that all drops are constantly dissolving or growing, favoring the development of a self-similar drop size distribution (DSD) at very long times. The predictions of the theory regarding Eq. 2 and the form of the DSD correspond to the so-called 'stationary regime' in which

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a characteristic, left-skewed, drop-size distribution with a cut-off radius of $1.5R_c$ should be attained.

As the drops of any emulsion, the drops of a nanoemulsion are also subject to a constant buoyancy force resulting from the gravity field of the earth and Archimedes' law:

$$F_b = 4\pi R_i^3 (\rho_o - \rho_w) g / 3 \quad (5)$$

This force causes the formation of cream at the top of the container and generates a gradient of concentration in the number of particles per unit volume $n(t)$ along its vertical axis. The velocity of creaming can be easily estimated, if the Brownian movement of the particles is neglected, from the deterministic term of the equation of motion of emulsion stability simulations (ESS; Urbina-Villalba, 2000) as

$$\Delta L = \frac{D_i F}{k_B T} \Delta t \Rightarrow V_g = \frac{\Delta L}{\Delta t} \approx \frac{[k_B T / 6\pi \eta R_i] F_b}{k_B T} \quad (6)$$

where ΔL : height of the container ($\Delta L = r_i(t + \Delta t) - r_i(t)$), $r_i(t)$: position of particle 'i' at time t , η : viscosity of the water phase, and k_B : Boltzmann constant. Using Eq. 5 for the buoyancy force, Eq. 6 yields

$$V_g = \frac{[k_B T / 6\pi \eta R_i] [4\pi R_i^3 \Delta \rho g]}{3 k_B T} = \frac{2 R_i^2 \Delta \rho g}{9 \eta} \quad (7)$$

Assuming that the height of the container is $\Delta L = 10\text{cm}$ (typical size of the sample vessel in a spectrophotometer), a dodecane drop located at the bottom of the container will require either 17s ($R_i = 100\mu\text{m}$) or 205 days ($R_i = 100\text{nm}$) to reach the top, depending on its size.

Notice that the potential energy of the buoyancy phenomenon does not affect the chemical potential of the oil molecules, which is constant and independent of R_i :

$$\frac{\Delta G_b}{N_{m,i}} = \left(\frac{4\pi R_i^3}{3} \right) (\rho_o - \rho_w) g (h_o - h) \left(\frac{4\pi R_i^3}{3} \right) \rho_o N_A / 3MW = (1 - \rho_w / \rho_o) g (h_o - h) MW / N_A \quad (8)$$

where h_o and h are the height of the liquid column and the position of the drop along its vertical axis, respectively.

According to Eq. 8 the ripening rate is not influenced by gravity. However, since the Ostwald ripening

process changes the radii of the particles by molecular exchange, the creaming velocity changes due to Ostwald ripening. A similar phenomenon occurs if a set of individual drops flocculate irreversibly to form an aggregate. The total mass of the aggregate is substantially higher than that of their drops and, therefore, the influence of the buoyancy force on the displacement of the cluster is larger. This can be appreciated if the radius of drop i is substituted by the hydrodynamic radius of an aggregate of size k ($R_k = \sqrt[3]{k} R_o$) in Eq. 5.

Moreover, since the relative movement of the drops, either by convection (orthokinetic flocculation) or by diffusion (perikinetic flocculation), depends on particle size, the flocculation rate also does. If the drops coalesce after flocculation, the critical radius of the emulsion increases and larger drops will dissolve. On the other hand, if the ripening occurs in the absence of coalescence, the DSD of the emulsion still changes favoring the occurrence of a variety of aggregates with distinct flocculation rates.

The examples given in the last two paragraphs illustrate the complex situation that might occur if a pair of destabilization processes occurs simultaneously. Fortunately, the effect of the gravity field can be disregarded if the difference between the density of the phases is small (see $\Delta \rho$ in Eq. 7). Similarly the ripening process could be diminished if the chosen oil is sparingly soluble in the water phase (see $C(\infty)$ in Eq. 2). When these conditions are met, the process of coagulation proceeds as described by Smoluchowski (1917). According to this author, the number concentration of aggregates of k primary particles existing at time t , $n_k(t)$, results from a balance between the aggregates produced by the collisions between clusters of smaller sizes i and j (such that $i+j=k$) and the aggregates of size k lost by the collisions with clusters of any other size:

$$\frac{dn_k(t)}{dt} = \frac{1}{2} \sum_{i+j=k-1} k_{ij} n_i(t) n_j(t) - n_k(t) \sum_{i=1}^{\infty} k_{ik} n_i(t) \quad (9)$$

The kernel of Eq. 9 is the set of coagulation rates between aggregates i and j $\{k_{ij}\}$. The evaluation of these rates is very difficult. Fortunately, the rate of doublet formation (k_{11}) is accessible from turbidity measurements. By definition, the turbidity of a dispersion (τ_{exp}) is equal to

$$\tau_{\text{exp}} = (I/L_c) \ln(I_0/I) \quad (10)$$

where I_0 : intensity of the incident light, and I : intensity of light emerging from a cell of path length L_c (generally of the order of 10^{-2}m). Theoretically, the turbidity can be expressed as

$$\tau_{\text{theo}} = \sum_{k=1}^{\infty} n_k \sigma_k \quad (11)$$

where σ_k : optical cross section of an aggregate of size k . Thus, according to Eqs. 10 and 11, the turbidity of a dispersion measures the amount of light that is lost by the scattering of the aggregates existing in a liquid.

If the time of measurement is short enough and the dispersion is sufficiently dilute, the initial slope of the turbidity as a function of time can be directly related to the rate of doublet formation k_{11} (Lips *et al.*, 1971; Lips and Willis, 1973):

$$\left(\frac{d\tau}{dt} \right)_0 = \left[\frac{\ln 10}{L_c} \right] \left(\frac{d\text{Abs}}{dt} \right)_0 = \left(\frac{1}{2} \sigma_2 - \sigma_1 \right) k_{11} n_0^2 \quad (12)$$

where $\text{Abs} = \log(I_0/I)$: absorbance of the dispersion, $n_0 = n(t=0)$, and σ_1 and σ_2 are the optical cross sections of a spherical drop and a doublet. According to the Rayleigh, Gans and Debye theory (RGD; Kerker, 1969), the cross sections of singlets and doublets can be computed using Eq. 12 whenever

$$C_{\text{RGD}} = (4\pi a/\lambda)(m-a) \ll 1 \quad (13)$$

being λ : wavelength of light in the medium ($\lambda = \lambda_o/n_w$ (where n_w : index of refraction of water, and λ_o : wavelength of light in vacuum)), and m : relative refractive index between the particle and the surrounding medium (n_o/n_w). Whenever Eq. 13 holds,

$$\sigma_k = \frac{4}{9} \pi R_k^2 \alpha_k^4 (m-1)^2 \int_0^{\pi} P_k(\vartheta) (1 + \cos^2 \vartheta) \sin(\vartheta) d\vartheta \quad (14)$$

where ϑ : angle of observation ($\alpha_k = 2\pi R_k/\lambda$), and $P_k(\vartheta)$ is the form factor of an aggregate of size k , deduced by Puertas *et al.* (Puertas *et al.*, 1997, 1998; Maroto y de las Nieves, 1997). In the case of a sphere (singlet),

$$P_1(\vartheta) = \left(3 \frac{\sin u - u \cos u}{u^3} \right)^2 \quad (15)$$

with $u = 2R\sin(\vartheta/2)$. In the case of a doublet,

$$P_2(\vartheta) = \left(2 + \frac{\sin 2u}{u} \right) \left(3 \frac{\sin u - u \cos u}{u^3} \right)^2 \quad (16)$$

In this work, the stability of four different nanoemulsions is

TABLE I
COMPOSITION OF THE EMULSIONS

System / Composition	A	B	C	D
SDS (%wt)	1.5	1.5	1.5	1.5
NaCl (%wt)	0.94	0.94	0.94	0.94
Isopentanol (%wt)	1.04	1.04	1.04	1.04
Weight fraction of water	0.93	0.93	0.93	0.93
Weight fraction of oil	0.07	0.07	0.07	0.07
Weight % of dodecane	100	93	70	56
Weight % of squalene	—	7	—	23
Weight % of tetrachloro-ethylene	—	—	30	21

studied. These emulsions are composed of: a) dodecane (Nano A), b) dodecane (C₁₂) + squalene (SQ) (Nano B), c) C₁₂ + tetrachloroethylene (TCE) (Nano C), and finally d) C₁₂ + SQ + TCE (Nano D). Four types of stability measurements are used:

1. The variation of R³ vs t

$$V_{\text{exp}} = dR_a^3/dt \quad (17)$$

2. The profile of the emulsion transmittance as a function of the container height at different times.

3. The creaming velocity (Eq. 7).

4. The rate of doublet formation k₁₁ (Eq. 12).

According to Kamogawa *et al.* (1999) the addition of an insoluble component to the oil (like squalene) decreases its aqueous solubility. It is also recognized that mixtures of dodecane with a denser substance (like TCE) can be used to produce a neutrally buoyant combination of oils. Hence, it is expected that Nano A will experience all destabilization phenomena occurring in a typical emulsion, while in the emulsions B and C the occurrence of ripening and buoyancy, respectively, will be restricted. Since Nano D contains both TCE and SQ, hypothetically only flocculation and coalescence are expected.

Experimental Procedure

Materials

Dodecane (Aldrich, 99%, 0.75g/cc) was eluted twice through an aluminum column in order to improve its purity. Sodium chloride (Sigma, 99.5%), iso-pentanol (Scharlau Chemie, 99%, 0.81g/cc), tetrachloroethylene (J.T. Ba-

ker, 100%, 1.614g/cc), sodium dodecyl sulfate (Sigma, 99%) and squalene (Aldrich, 99%, 0.809g/cc) were used as received. The water used was distilled and deionized (1.1μS·cm⁻¹ at 25°C) using a Simplicity purifier from Millipore (USA).

Nanoemulsion synthesis and characterization

Nanoemulsions were prepared using the phase inversion composition method (Solè *et al.*, 2006; Wang *et al.*, 2008). In order to guarantee the occurrence of minimum tension during the mixture of the components, the phase diagram of a system composed of dodecane, NaCl, iso-pentanol, SDS and water was previously built (Rahn-Chique *et al.*, 2012a, b). For this study, 0.95g of an 11.5% wt/wt aqueous solution of NaCl were mixed with 0.846g of oil and 0.2g of SDS. Then, enough iso-pentanol was added to reach 6.5% wt/wt in the total mixture. The mixture was then stirred with a mechanical stirrer at a velocity of 14500rpm while additional water was added at a rate of 8-10cm³·s⁻¹. During this time (1min), the agitator was moved conveniently to guarantee the correct homogenization of the mixture. Following this procedure, O/W nano-

emulsions with average diameters of 401-440nm were obtained.

The composition of the oil was systematically varied in order to produce systems with small ripening rate and/or neutral buoyancy. To these aims, previous experiments with non-ionic surfactants (Cruz, 2012) and hexadecane (García-Valero, 2011) showed that mixtures of dodecane with at least 7% wt/wt of squalene (SQ), or 30% wt/wt of tetrachloroethylene (TCE), respectively, were required. Since the addition of TCE promotes a considerable degree of ripening (see below), and the addition of squalene lowers the density of the mixture considerably, it was necessary to make a compromise in order to obtain a system of maximum stability: C₁₂ 56% wt/wt, TCE 23% wt/wt, SQ 21% wt/wt (Nano D). The nominal compositions of the mother emulsions used and their physical properties are given in Tables I and II.

Due to the detection limits of the instruments, two different dilutions were necessary in order to follow the evolution of the systems (Table III). In all dilutions, the final concentrations of NaCl and SDS were adjusted to 5 and 8mM, respectively. The evolution of the average radius as a function of time was measured using a Brookhaven Goniometer at n₀=4×10⁹ particles/cc. Electrophoretic measurements (Delsa 440SX, Beckman-Coulter) and turbidity evaluations as a function of height (Quickscan, Formulacion) were carried out at n₀=4×10¹⁰ particles/cc. The

TABLE II
PROPERTIES OF THE MOTHER EMULSIONS

System / Composition	A	B	C	D
Density of oil mixture (g·ml ⁻¹)	0.750 ±0.001	0.756 ±0.001*	0.991 ±0.001*	0.955 ±0.001*
Refractive index of oils	1.425 ±0.002	1.430 ±0.002	1.449 ±0.002	1.450 ±0.002
Average drop radius (nm)	400 ±100	420 ±90	410 ±70	400 ±100
N ₀ (particles/ml)	2.2×10 ¹²	2.2×10 ¹²	1.7×10 ¹²	8.3×10 ¹²
Zeta potential of drops (mV)	-84 ±14	-89 ±14	-73 ±17	-94 ±14
C _{RGD}	0.313	0.330	0.320	0.396
α	2.19	2.19	2.19	2.19

*These errors were formerly estimated by weighting five 1ml samples of the mixtures. However, such a procedure only validates the precision of the measurements. Since the provider (Aldrich) can only assess the exactness of the density of pure dodecane up to three significant figures, this estimation of the error (0.001) was employed in all cases.

TABLE III
COMPOSITION AND PROPERTIES OF DILUTED NANO EMULSIONS

System / Composition	A	B	C	D
SDS (M)	8×10 ⁻³	8×10 ⁻³	8×10 ⁻³	8×10 ⁻³
NaCl (M)	5×10 ⁻³	5×10 ⁻³	5 x 10 ⁻³	5×10 ⁻³
N ₀ (particles/ml) for turbidity measurements	4×10 ⁹	4×10 ⁹	4×10 ⁹	4×10 ⁹
N ₀ (particles/ml) for creaming velocity evaluation	4×10 ¹⁰	4×10 ¹⁰	4×10 ¹⁰	4×10 ¹⁰
Creaming Velocity (cm·s ⁻¹)	(3.6 ±0.5)×10 ⁻⁶	(2.1 ±0.5)×10 ⁻⁶	~0	(-3.4 ± 0.5)×10 ⁻⁷
dR ³ /dt (m ³ ·s ⁻¹)	(2.7 ±0.9)×10 ⁻²⁶	(1.6 ±0.9)×10 ⁻²⁸	(4.7 ±0.2)×10 ⁻²⁶	(-3.1 ±0.8)×10 ⁻²⁷

latter measurements were undertaken at 200 acquisitions/h during eight days. The creaming rate was calculated with the Migration software (Formulation, 2002) employing

$$V_g = \frac{2 R_i^2 \Delta \rho g}{9 \eta} \left[1 - \phi \right] \left[1 + 4.6 \phi / (1 - \phi)^3 \right] \quad (18)$$

Evaluation of the flocculation rate for doublet formation (k_{11})

To select a convenient wavelength for turbidity measurements, the absorption spectra of each emulsion and each emulsion component were measured separately between 200 and 1100nm using a UV-Visible spectrophotometer (Turner SP 890). The optimum wavelength for scattering studies is that which warrants negligible adsorption of all components, and shows a significant variation of the absorbance as a function of time. For the present studies a value of $\lambda = 800\text{nm}$ was selected.

The appropriate particle concentration for aggregation studies was established from plots of $(dAbs/dt)_0$ vs n_0 . For this purpose, the mother emulsions were diluted to produce initial particle concentrations (n_0) between 8×10^9 and 6×10^{10} drops/cc (in the case of system A the higher limit was extended to 2×10^{11} drops/cc). The absorbance of each emulsion (t_{max}) was measured at least thrice during 60s after the addition of 600mM NaCl, following the procedure previously described by Rahn-Chique (2012a, b). From these curves, the initial slope of the absorbance $(dAbs/dt)_0$ was computed. An approximately linear region of $(dAbs/dt)_0$ vs n_0 was found to be below 6×10^{10} particles/cc.

Using a fixed particle concentration of $n_0 = 4.0 \times 10^{10}$ particles/cc, the absorbance of each emulsion (A, B, C and D) was evaluated as a function of time, at different electrolyte concentrations (350-600mM NaCl). One mother emulsion of each type (A, B, C, D) was synthesized for each salt concentration. The average radius of the drops and the polydispersity of each type of emulsion were quite reproducible: A (407-430nm, 19-22%), B (407-414nm, 20-22%), C (401-443nm, 19-21%) and D (400-450nm, 25-27%). The appropriate amount of salt was introduced directly into the sample cell of

the spectrophotometer (Turner, SP 890) by injecting 0.6cc of brine to 2.4cc of each diluted emulsion.

Results and Discussion

According to LSW the rate of Ostwald ripening is given by a linear slope of R_c^3 vs t (V_{exp} in Eq. 17). However our simulations (Urbina-Villalba *et al.*, 2009; 2012a, b) suggest that flocculation and coalescence contribute significantly to V_{exp} . Thus, $V_{exp} = V_{FCOB}$ (where F, C, O and B stand for flocculation, coalescence, Ostwald ripening, and buoyancy). If a repulsive potential exists between the drops, the calculations predict a concave downward curve whose slope decreases with time approaching the LSW limit. The curve oscillates above and below its average slope because it results from two opposing trends: 1) the increase of the average radius of the emulsion produced by the elimination of drops either by coalescence and/or complete dissolution, and 2)

the decrease of R_a by the molecular exchange between the drops, which only lowers the average radius. Consequently, the curve of R^3 vs t shows a saw-tooth variation in which R_a grows when the number of drops diminishes, but it decreases at a constant number of drops.

Due to their small size, nanoemulsion drops are expected to be non-deformable. In the present case the drops are also highly charged, due to the adsorption of SDS. The electrostatic surface potential of the drops increases in the order $C < A < B < D$ (Table II) evidencing that the composition of the oil affects the adsorption process. Consequently, their flocculation rates are expected to decrease in the order $C > A > B > D$. Besides, since aggregation is a necessary step prior to coalescence, both flocculation and coalescence are restricted for these systems in the absence of salt. Incidentally, the highest value of V_{exp} corresponds to system C, and its lowest variation to system D (Figure 1).

According to Eq. 2 the ripening rate of dodecane in the stationary regime should be equal to $1 \times 10^{-26} \text{m}^3 \cdot \text{s}^{-1}$. Figure 1 illustrates the long-time behavior of the systems. It can be observed that the value of R^3 oscillates as a function of time. If the average slope of each curve is subtracted from $R^3(t)$, a deviations histogram can be calculated. The frequency of the deviations can then be adjusted to a Gaussian distribution. The standard deviation of the distribution quantifies the amplitude of the oscillations, which is found to decrease in the order $A > C > D \geq B$ (Figure 2). Thus, the addition of SQ to systems A and C (systems B and D, respectively), decreases the amplitude of the oscillations.

Moreover, the average slope of systems A and C also decrease (Table III). These results can be justified in terms of the lower solubility of the oil mixture, which decreases with the addition of SQ. In fact, increasing the solubility of C_{12} (system A) by the addition of TCE (system C) should increase V_{exp} , as it was experimentally found (Table III). This does not occur when we move from system B to system D, because in this case the solubility increases with the addition of TCE, but also decreases with the increase of SQ (from 7 to 23%; Table I).

Thus, the following question can be asked: Is the variation of R_a^3 vs t a consequence of the degree of floccula-

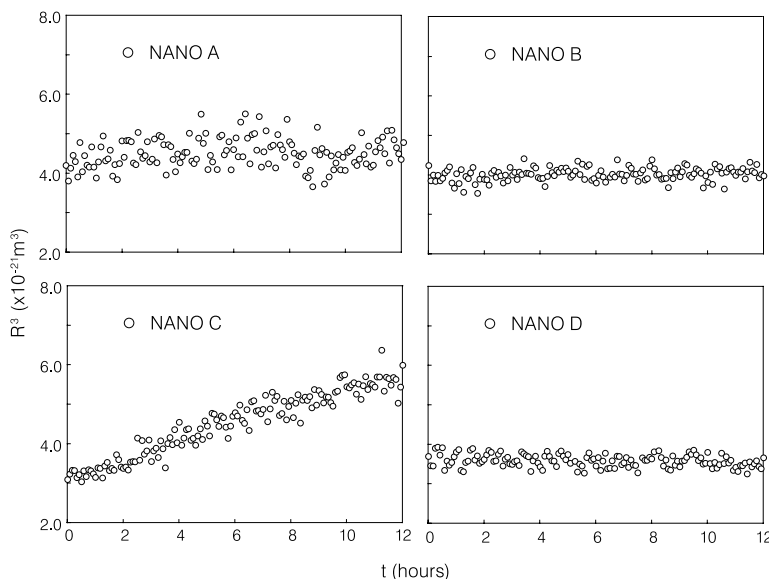


Figure 1. Change of R_a^3 vs t for each nanoemulsion.

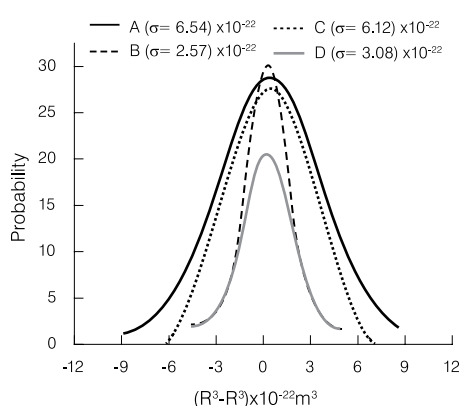


Figure 2. Frequency of the deviations of R_a^3 vs t with respect to its average slope.

tion of the emulsions, or is it the result of the Ostwald ripening phenomenon? The truth is that when several destabilization processes are combined, V_{exp} is neither a sound measure of flocculation (see below) nor a good measure of the ripening rate. In fact, system D shows a negative slope ($-3.1 \times 10^{-27} \text{m}^3 \cdot \text{s}^{-1}$) that cannot be explained by LSW theory, but is perfectly consistent with our previous simulations: if the rate of elimination of the drops is substantially decreased by lowering both the coalescence rate and the dissolution of the particles, only molecular exchange survives. In this case the simulations predict a decrease of the average radius as a function of time that LSW is unable to justify. In our view, this information, along with the fact that the zeta potential of systems A and B is very similar (Table II) but their behavior is distinct, indicates that the Ostwald ripening phenomenon predominates: $V_{exp} \approx V_{OB}$.

In regard to buoyancy, Figure 2 confirms that despite the small size of their drops, the systems with the lower densities (A and B) are subject to a substantial degree of creaming. The base of the container clarifies as a function of time, showing higher transmittances, and the top is progressively obscured. Instead, the systems with TCE (systems C and D) do not show a significant amount of creaming. Notice the very different behaviors of these two systems in regard to V_{exp} (Figure 1), despite their similarities with respect to V_g (Figure 3 and Table III). Figures 1 and 3 demonstrate that V_{exp} and V_g show different trends in regard to emulsion stability. In particular, the creaming rate is not a convenient measurement of emulsion stability, because creaming is not the main destabilization phenomenon in the present systems. In our understanding, the most unstable system is the one which presents the quickest departure from its initial state (and vice-versa). This is not necessarily equivalent to the fastest variation of the macroscopic properties. Moreover, such change can be caused by a predominant destabilization phenomenon, or a combination of several contributing phenomena. The system with the highest V_{exp} rate (C) will only show the highest creaming rate if $\Delta\rho$ is appreciable, and this is not the case.

Figure 4 shows the variation in the initial slope of the absorbance as a function of the ionic strength. This slope is proportional to the initial aggregation rate (k_{11} in Eq. 12), but also depends on the optical cross sections of singlets and doublets (at constant n_0). With respect to $(dAbs/dt)_0$ each set of data can be divided into two linear regions which in the case of solid particles correspond to diffusion limited cluster aggregation (DLCA) and reaction limited cluster aggregation (RLCA),

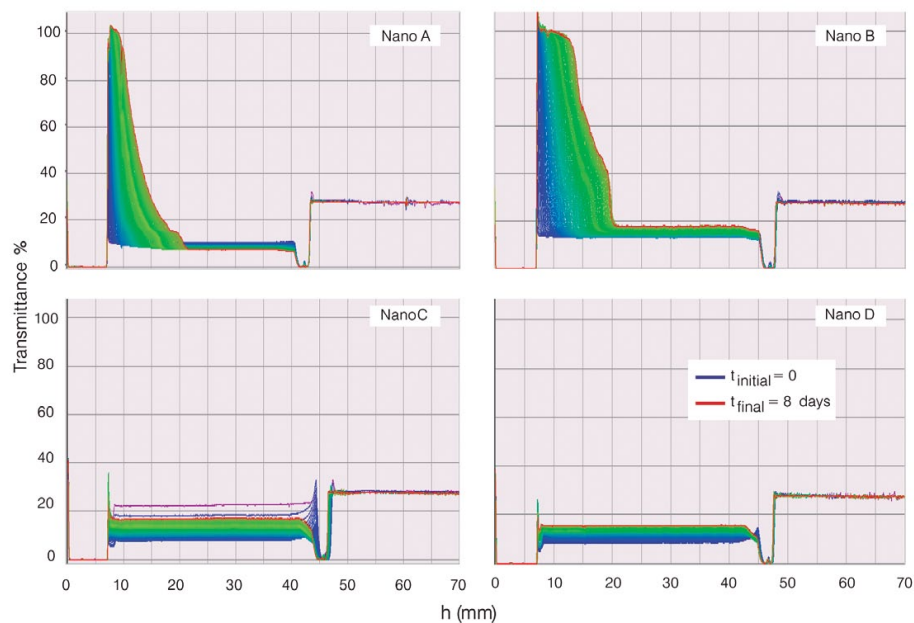


Figure 3. Change in transmittance of the mother emulsions as a function height and time.

respectively. The inflexion point is located around 400mM NaCl, which determines the critical coagulation concentration (CCC). Above this salt concentration, the surface charge of the drops is screened by the electrolyte, and a maximum aggregation rate is achieved. Below 400mM NaCl electrostatic interactions dominate.

As it is evident from Figure 5, σ_1 and σ_2 are sensitive functions of the average radius of the emulsions (Table II). Within the DLCA regime, systems A and C show maxima for minimum values of R_a and vice-versa. Since the average radius of the emulsions was measured in the absence of salt, it appears to be coincidental that the emulsions of these two systems show similar dependences of k_{11} with respect to the salt concentration. Instead, a smooth variation of k_{11} vs $[\text{NaCl}]$ is exhibited by the systems containing squalene (B and D) in the same range of salt concentrations. It is also remarkable that the curves of k_{11} vs $[\text{NaCl}]$ are not straight lines within the RLCA regime. All systems show a change of slope around 370-385mM (see also Figure 6 and related discussion in Rahn-Chique *et al.*, 2012a).

Theoretically, the most unstable system with regard to aggregation is the one with highest k_{11} , lowest CCC, and highest $dk_{11}/d[\text{NaCl}]$ (for $[\text{NaCl}] < \text{CCC}$). All systems studied show a CCC of 400mM. However, for the DLCA regime the absolute values of k_{11} decrease in the order: $C \gg D \approx B \geq A$, while for the RLCA regime they vary as $B > A > D > C$. In regard to the change of k_{11} with respect to the ionic strength ($dk_{11}/d[\text{NaCl}]$), $B < A < D < C$. In our view, the absolute value of k_{11} is a determining factor in regard to

aggregation, and therefore, the relative stability of these systems depends on the salt concentration.

In any event, since the change of R_a^3 vs t is the result of all destabi-

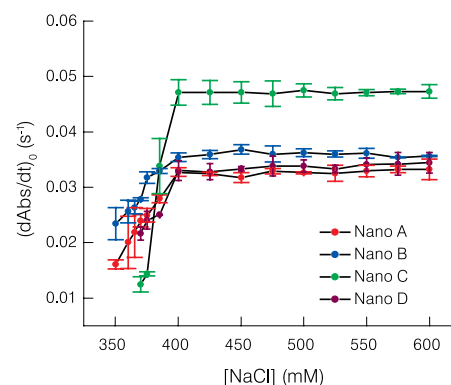


Figure 4. Initial slope of the absorbance as a function of ionic strength for systems A, B, C and D.

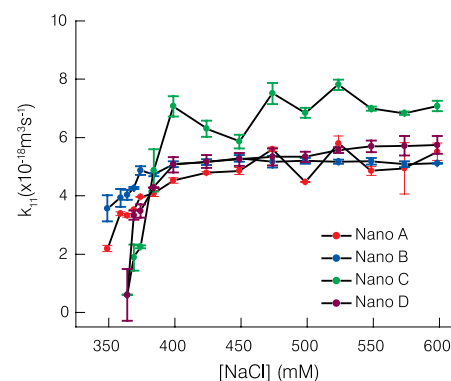


Figure 5. Dependence of k_{11} on the salt concentration for systems A, B, C and D.

bilization phenomena, it appears to be the most reliable measure of stability in these nanoemulsions.

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INFLUENCIA DE LA MADURACIÓN Y FORMACIÓN DE CREMA EN LA TASA DE AGREGACIÓN DE NANOEMULSIONES DE DODECANO-AGUA. ¿ES LA TASA DE FORMACIÓN DE CREMA UNA MEDIDA APROPIADA DE LA ESTABILIDAD DE LA EMULSION?

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RESUMEN

Se estudia el comportamiento de cuatro nano-emulsiones iónicas de aceite en agua (O/W) compuestas por dodecano puro y mezclas de dodecano con esqualeno y tetracloroetileno. Las nanoemulsiones fueron estabilizadas con dodecil-sulfato de so-

dio (DSS). La variación del radio promedio y la turbidez fue estudiado en función del tiempo. Los resultados ilustran las limitaciones de caracterizar la estabilidad de emulsiones a través de su tasa de formación de crema.

INFLUÊNCIA DO AMADURECIMENTO E FORMAÇÃO DE CREME NA TAXA DE AGREGAÇÃO DE NANOEMULSÕES DE DODECANO/ÁGUA. É A TAXA DE FORMAÇÃO DE CREME UMA MEDIDA APROPRIADA DA ESTABILIDADE DA EMULSÃO?

Yorlis Mendoza, Kareem Rahn-Chique, Antonio M. Puertas, Manuel S. Romero-Cano e German Urbina-Villalba

RESUMO

Estuda-se o comportamento de quatro nano-emulsões iônicas de óleo em água (O/W) compostas por dodecano puro e misturas de dodecano com esqualeno e tetracloroetileno. As nanoemulsões foram estabilizadas com dodecil-sulfato de sódio (DSS).

O comportamento do raio médio e a turbidez foi estudada em função do tempo. Os resultados ilustram as limitações de caracterizar a estabilidade de emulsões a través de sua taxa de formação de creme.