Kinetics of thermo-induced micelle-to-vesicle transitions in a catanionic surfactant system investigated by stopped-flow temperature jump

Jingyan Zhang\textsuperscript{ab} and Shiyong Liu\textsuperscript{a}*

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The kinetics of thermo-induced micelle-to-vesicle transitions in a catanionic surfactant system consisting of sodium dodecyl sulfate (SDS) and dodecyltriethylammonium bromide (DEAB) were investigated by the stopped-flow temperature jump technique, which can achieve T-jumps within ~2–3 ms. SDS/DEAB aqueous mixtures ([SDS]/[DEAB] = 2/1, 10 mM) undergo microstructural transitions from cylindrical micelles to vesicles when heated above 33 °C. Upon T-jumps from 20 °C to final temperatures in the range of 25–31 °C, relaxation processes associated with negative amplitudes can be ascribed to the dilution-induced structural rearrangement of cylindrical micelles and to the dissolution of non-equilibrium mixed aggregates. In the final temperature range of 33–43 °C the obtained dynamic traces can be fitted by single exponential functions, revealing one relaxation time (τ) in the range of 82–440 s, which decreases with increasing temperature. This may be ascribed to the transformation of floppy bilayer structures into precursor vesicles followed by further growth into final equilibrium vesicles via the exchange and insertion/expulsion of surfactant monomers. In the final temperature range of 45–55 °C, vesicles are predominant. Here T-jump relaxations revealed a distinctly different kinetic behavior. All dynamic traces can only be fitted with double exponential functions, yielding two relaxation times (τ\textsubscript{1} and τ\textsubscript{2}), exhibiting a considerable decrease with increasing final temperatures. The fast process (τ\textsubscript{1} ~ 5.2–28.5 s) should be assigned to the formation of non-equilibrium precursor vesicles, and the slow process (τ\textsubscript{2} ~ 188–694 s) should be ascribed to their further growth into final equilibrium vesicles via the fusion/fission of precursor vesicles. In contrast, the reverse vesicle-to-micelle transition process induced by a negative T-jump from elevated temperatures to 20 °C occurs quite fast and almost completes within the stopped-flow dead time (~2–3 ms).

Introduction

Amphiphilic molecules possess the ability of self-assembling into a large variety of morphologically different aggregate microstructures, such as spherical and rod-like micelles, vesicles, and lamellar phases.\textsuperscript{1–4} In the past decades, considerable attention has been devoted to theoretical and experimental investigations of the phase behavior, equilibrium microstructures, and morphological transitions of these aggregates.\textsuperscript{5–12} In the context of small molecule surfactant systems, microstructural transformation between micelles and vesicles can be induced via the variation of surfactant concentrations, temperature, pH, and the presence of external additives (salts, surfactants).\textsuperscript{13–24}

Thermo-induced micelle-vesicle transitions are particularly interesting due to the non-invasiveness and facile manipulation of temperature variations.\textsuperscript{25–34} Though vesicle-to-micelle transitions upon heating are quite typical in some ionic surfactant systems,\textsuperscript{33–35} the reverse process, i.e., heating-induced micelle-to-vesicle transition in surfactant systems is quite rare.\textsuperscript{27,30,31} Early examples have been reported by Hoffmann et al.\textsuperscript{26,37} and Blume et al.\textsuperscript{38} concerning zwitterionic surfactant/cosurfactant mixtures and lipid/ionic surfactant mixtures, respectively. Later on, Huang et al.\textsuperscript{30,31} reported several examples of thermo-induced sequential micelle-to-vesicle transitions in catanionic surfactant systems. In one particular example, they observed that the cationic-anionic surfactant system of sodium dodecyl sulfate (SDS)/dodecyltriethylammonium bromide (DEAB) exhibits transitions from cylindrical micelles to vesicles upon heating above ~30 °C.\textsuperscript{30} At a temperature range of 30–50 °C, the number density of the vesicles continuously grows at the expense of the cylindrical micelles. At even higher temperatures, vesicle aggregates can be observed. Recently, Holyst, Hao, and coworkers\textsuperscript{32} reported that the microstructural evolution between micelles and vesicles in salt-free catanionic...
surfactant mixtures of tetradecltrimethylammonium hydroxide (TTAOH) and fatty acid are mainly dependent on surfactant concentrations.

Although the physicochemical properties and evolution of equilibrium microstructures of these thermo-induced transitions between micellar and vesicular phases have been well-characterized, much less is known concerning the kinetics of thermo-induced micelle-to-vesicle transition and the microstructural reorganization under various conditions. The kinetics of micelle-to-vesicle transition and the reverse process is not only of fundamental scientific interest, but also plays a vital role in the context of their technological applications. On the one hand, it provides fresh insights into their microstructural stability and offers new approaches for controlling their final microstructures. Moreover, many vesicle systems are not in the thermodynamic equilibrium state and exist as long-lived metastable structures; thus, exploring the kinetic pathways of their formation is crucial. On the other hand, catanionic vesicles have been the subject of extensive experimental and theoretical investigations due to their long-term stability and spontaneous formation from mixed surfactant systems. The versatile physicochemical properties of catanionic vesicles have allowed for several application-related studies, such as the preparation of magnetic nanoparticles and hollow spheres and the formation of polymer-vesicle gels and networks. Furthermore, they have been employed for the encapsulation of probe molecules and pharmaceutical drugs, or as gene delivery nanocarriers. It is worth noting that the stability, and formation/breakdown kinetics of catanionic vesicles are directly correlated with their functional applications.

Previously, vesicle formation kinetics of catanionic systems consisting of sodium octyl sulfate (SOS)/cetyltrimethylammonium bromide (CTAB), SDS/dodecyltrimethylammonium bromide (DTAB), or lecinthin/bile salt mixtures were investigated by the stopped-flow technique in combination with time-resolved laser light scattering (LLS) measurements. It has been tentatively proposed that the vesicle formation process consists of a series of events: the fastest is associated with the formation of non-equilibrium mixed aggregates, the following two relaxation steps are the formation of loose or “floppy” bilayer structures through the rearrangement of non-equilibrium mixed aggregates and the formation of vesicle precursors, and the rate-determining step is attributed to the formation of final equilibrium vesicles. Most recent developments in elucidating the vesicle formation kinetics have come from instrumental progress, and the combination of small angle neutron scattering (SANS) or small angle X-ray scattering (SAXS) with the stopped-flow technique allows the accurate probing of microstructural evolution processes, especially at the early stage of vesicle formation. In this context, Gradzielski et al. and Narayanan et al. investigated the kinetics of micelles-to-vesicle transitions upon stopped-flow equimolar mixing of anionic surfactants with cationic or zwitterionic surfactants detected by SAXS. They proposed that the vesicle formation is a complex multistep process: starting with the quick formation of non-equilibrium mixed globular micelles and the subsequent dissolution with a time constant of 0.5–1 s; after or accompanied with the micelle dissolution process, bilayers then quickly form and slowly transform into unilamellar vesicles; the subsequent process is the growth of vesicles into larger and more monodisperse ones.

Another important issue in understanding the vesicle formation kinetics is to elucidate the underlying mechanisms and pathways. The aggregation dynamics of small molecule surfactant near the association equilibrium can be relatively well described by the Aniansson and Wall (A–W) theory with an important assumption that all changes are due to an elementary process of insertion/expulsion of individual chains (“unimers”) into/out of the micelle. Kahlweit and coworkers later proposed that at higher surfactant concentrations, micelle fusion/fission process is favored instead of the unimer insertion/expulsion pathway. Experimental studies in this aspect revealed that both mechanisms might take effect during the vesicle formation process, which is dependent on specific systems.

To the best of our knowledge, the kinetics of thermo-induced micelle-to-vesicle transition has never been investigated in literature reports possibly due to the following two reasons: (1) the lack of appropriate surfactant systems; (2) the lack of suitable instrumental techniques. Existing techniques of temperature jump are typically induced by laser flash or electric discharge, and the temperature elevation can be only maintained up to 10–100 ms, which is surely not long enough for the monitoring of vesicle formation kinetics. Recent technical progress has allowed the design of stopped-flow devices coupled with the millisecond temperature jump (mT-jump) accessory. Thus, temperature jumps can be conveniently achieved with a typical dead time of 2–3 ms. Most importantly, the measurement time windows can be extended up to several hours. Herein, we report the first kinetic investigation of thermo-induced micelle-to-vesicle transition and the reverse process of SDS/DEAB catanionic surfactant system by employing stopped-flow T-jump coupled with light scattering detectors. By controlling the final temperatures, the transformation kinetics from cylindrical micelles to the micelle-vesicle coexisting phase and to the pure vesicular phase has been probed. In combination with the experimental results, we proposed that the thermo-induced vesicle formation pathway strongly depends on the final temperatures. The thermo-induced vesicle-to-micelle transition kinetics and relevant mechanisms were also explored.

Experimental section

Materials

Sodium dodecyl sulfate (SDS) (analytical grade, Shanghai Chemical Reagent Co.) was recrystallized from anhydrous ethanol. Dodecyltrimethylammonium bromide (DEAB) was synthesized from dodecyl bromide and triethylamine, and the crude product was recrystallized three times from acetone/ethanol mixtures. Stock solutions of SDS/DEAB mixtures were prepared in deionized water. For stopped-flow studies, all solutions prior to mixing were filtered through 0.45 μm nitrocellulose filters (Acrodisc) to remove dusts.
Transmittance measurements

The optical transmittance of the aqueous solution at a wavelength of 600 nm was recorded on a Unico UV/vis 2802PC spectrophotometer equipped with a thermostatically controlled cuvette.

Stopped-flow temperature jump with light-scattering detection

Stopped-flow studies were carried out using a Bio-Logic SFM300/S stopped-flow instrument. It is equipped with three 10 mL step-motor-driven syringes (S1, S2, and S3), which can be operated independently to carry out single- or double-mixing. The stopped-flow device is attached to a MOS-250 spectrometer; kinetic data are fitted using the Biokine program provided by Bio-Logic. For the light scattering detection at a scattering angle of 90°, both the excitation and emission wavelengths were adjusted to 335 nm with 10 nm slits. Using FC-08 or FC-15 flow cells, typical dead times are 1.1 ms and 2.6 ms, respectively. All kinetic traces were averaged from at least five consecutive T-jump kinetic measurements.

The millisecond temperature jump (mT-jump) accessory is equipped with a standard Bio-Logic stopped-flow observation cell, which achieves temperature changes by mixing two solutions of different initial temperatures ($T_1$ and $T_2$), and the final temperature of the mixture ($T_{\text{final}}$) is determined by the initial temperature ($T_1$ and $T_2$) and the mixing ratio. Three thermoelectric Peltier elements are used to control the initial temperatures of the two solutions and that of the observation cell after mixing. The temperature of the mixed solution was stabilized to be the same as the observation cell (Peltier controlled) with the aid of a thermosensitive fluorescent dye, N-acetyl-L-tryptophanamide (NATA). The precision of the temperature jump is within ±0.1 °C, and the temperature stability in the observation cell after the temperature jump is <1% in 30 s. For the experimental setup in this work, the T-jump of the SDS/DEAB aqueous mixture ([SDS]/[DEAB] = 2/1, 20 mM) from low to high temperatures (to induce micelle-to-vesicle transition) at an initial temperature of 20 °C ($T_1$) was performed by 1:1 v/v mixing with water at different temperatures ($T_2$, 20-90 °C) to target varying final temperatures ($T_{\text{final}}$, 20-55 °C) and a final concentration of 10 mM. On the other hand, the vesicle-to-micelle transition was induced by T-jump from high to low temperatures for the surfactant mixture ([SDS]/[DEAB] = 2/1, 30 mM) at varying initial temperatures (38-50 °C) by 1:2 v/v mixing with cold water (10-22 °C) to target a final temperature of 20 °C and a final concentration of 10 mM.

Results and discussion

Temperature-dependent optical transmittance measurements

Previously, Huang et al.30 systematically investigated thermo-induced micelle-to-vesicle transitions in SDS/DEAB catanionic surfactant systems by employing a combination of dynamic LLS, freeze fracture electron microscopy (FF-EM), and rheological measurements. At 20 °C, they observed that cylindrical micelles are the major aggregates coexisting with a few small spherical vesicles, and the intensity-average hydrodynamic radius, ($R_h$), is ~25 nm. As the temperature increased to ~30 °C, more vesicles are observed ($R_h$ ~ 100 nm). Further temperature increase is associated with an increase in the number density of vesicles at the expense of cylindrical micelles. At ~40 °C, vesicles are the dominating aggregates, whereas at 50 °C, cylindrical micelles almost completely disappear and a few vesicle aggregates also form. They ascribed the observed thermo-induced micelle-to-vesicle transition to the weakened hydration of surfactant head groups at elevated temperatures.

We measured the temperature dependence of optical transmittance at a wavelength of 600 nm for SDS/DEAB aqueous mixtures at [SDS]/[DEAB] = 2/1 and a total surfactant concentration of 10 mM (Fig. 1). In reasonable agreement with the results reported by Huang et al.,30 we found that heating the aqueous mixture from 15 °C to 33 °C leads to almost no changes in optical transmittance, apparently indicating the lack of any major morphological transformations. Above 33 °C, the optical transmittance abruptly decreases to ~70% at 60 °C, which stabilizes out at even higher temperatures. As reported by Huang et al.,30 this can be ascribed to the micelle-to-vesicle transition. Higher temperatures are advantageous to the formation of vesicles with larger number densities. At a temperature range of 15–75 °C, no macroscopic phase separation can be observed. Fig. 1 also shows temperature-dependent changes of optical transmittance in the cooling cycle, indicating that the micelle-to-vesicle transition is quite reversible in nature.

Kinetics of thermo-induced micelle-to-vesicle transition: T-jump from 20 °C to final temperatures in the range of 25–31 °C

Fig. 2 depicts time-dependent scattered light intensities recorded for an SDS/DEAB aqueous mixture ([SDS]/[DEAB] = 2/1) upon a stopped-flow temperature jump from 20 °C to final temperatures in the range of 25–31 °C. All dynamic traces exhibit an initial decrease and then stabilize out. We can apparently observe that the higher the final temperatures, the slower the kinetic processes. Note that for stopped-flow experiments, the observed kinetics upon T-jump should be ascribed to dynamic processes occurring for the surfactant mixture at 10 mM (final concentration after stopped-flow 1:1 v/v mixing) as the uniform physical mixing process (associated with dilution) completes within a millisecond. We also plotted
of cylindrical micelles upon dilution. Previously, we investigated
associated with changes in micellar sizes and the size distributions
(PTHC).53,54 Initial decrease of scattering intensities upon
oxybenzene sulfonate (MOBS) and
micelles consisting of sodium 4-(8-methacryloyloxyoctyl)-
the kinetics of dilution-induced disintegration of worm-like
20
is reasonable considering that within the investigated final
aggregate structures due to their poor structural stability. This
dissolution/structural rearrangement of non-equilibrium
decrease in scattering light intensity can be attributed to the
non-equilibrium mixed aggregates, whereas the subsequent
tentatively ascribe the initial fast process to the formation of
In combination with the results reported by Hatton
only observed relaxation processes with negative amplitudes.
stopped-flow mixing) at higher final temperatures implies that
dilution effects. The initial fast process (which completes
more kinetic events might have occurred in addition to the
dilution effects. The initial fast process (which completes
3–4 s. At 20 °C, cylindrical micelles are the major form of aggregates for SDS/DEAB
aqueous mixtures with the total concentration being 10 or 20 mM.
Thus, the initial decrease of scattered intensities occurring at
20 °C (i.e., no T-jump) can be ascribed to the dilution effect
associated with changes in micellar sizes and the size distributions
of cylindrical micelles upon dilution. Previously, we investigated
the kinetics of dilution-induced disintegration of worm-like
micelles consisting of sodium 4-(8-methacryloyloxyoctyl)-
oxylbenzene sulfonate (MOBS) and p-toluidine hydrochloride
(PTHC).53,54 Initial decrease of scattering intensities upon
stopped-flow dilution is also observed at final concentrations
much higher than the critical aggregation concentrations
(CACs).

Although the relaxation process at other final temperatures
associated with negative amplitudes are similar to that occurring
at 20 °C, the values of the initial and final equilibrium
scattering intensities apparently increase with the final
temperatures. The larger scattering intensity at higher final
temperatures should be ascribed to the slight increase,
albeit very low, of the number density of spherical vesicles,
as suggested by Huang and coworkers30,31 through dynamic
LLS and rheology measurements. The more prominent
growth of the initial scattering intensity (i.e., ~2–3 ms after
stopped-flow mixing) at higher final temperatures implies that
more kinetic events might have occurred in addition to the
dilution effects. The initial fast process (which completes
within the stopped-flow dead time, ~2–3 ms) associated with
the increase in light scattering intensity occurs although we
only observed relaxation processes with negative amplitudes.
In combination with the results reported by Hatton et al.,45 we
tentatively ascribe the initial fast process to the formation of
non-equilibrium mixed aggregates, whereas the subsequent
decrease in scattering light intensity can be attributed to the
dissolution/structural rearrangement of non-equilibrium
aggregate structures due to their poor structural stability. This
is reasonable considering that within the investigated final
temperature range (25–31 °C), cylindrical micelles are the
predominant form. We thus established that dynamic traces
are a combination of the results of the dilution effects and the
initial formation of non-equilibrium mixed aggregates, and
their subsequent structural rearrangement.

All the dynamic traces in Fig. 2 can be fitted with single
exponential functions. The final temperature dependence
of relaxation times is plotted in Fig. 3, which is in the range of
~1.2–2.8 s. We can clearly observe an increase in relaxation
times with increasing final temperatures. At elevated temperatures,
the higher structural stability of the initially formed
non-equilibrium mixed aggregates might lead to longer time
scales required for aggregate disintegration and structural
rearrangement.

**T-jump from 20 °C to final temperatures in the range of
33–43 °C**

Upon a stopped-flow temperature jump from 20 °C to final
temperatures in the range of 33–43 °C, the time dependence
of scattered light intensity recorded for SDS/DEAB aqueous
mixture is shown in Fig. 4a. Closer examination of the
dynamic trace recorded upon T-jump from 20 °C to 33 °C
(Fig. 4b) reveals that the scattered light intensity shows an
abrupt decrease within the first ~20 s, and then gradually
increases with time even after ~3000 s. As discussed in the
previous section, the initial decrease in scattered light intensity
should be mainly ascribed to the disintegration of quickly
formed non-equilibrium mixed aggregates. Since vesicles starts
to appear at final temperatures ≥ 33 °C, it is quite expected
that the observed increase in scattered light intensity at later
stages should be attributed to the formation of floppy bilayer
structures, the subsequent closure into precursor vesicles, and
then the vesicle growth.

At these final temperatures, only one relaxation process
associated with positive amplitude can be typically observed,
and we do not observe any relaxation process associated with
negative amplitude. It can be inferred from previous examples
that during vesicle formation, the formation of intermediate
states (e.g., floppy bilayer structures) is a very fast process
(within ~1 s).46,55 The observed time-dependence of scattered
light intensity is drastically different compared to that at a
final temperature of 33 °C, which exhibits dramatic and abrupt

![Fig. 2](image_url)  
**Fig. 2** Time dependence of scattered light intensities recorded for SDS/DEAB aqueous mixture ([SDS]/[DEAB] = 2/1) upon stopped-flow temperature jump from 20 °C to varying final temperatures in the range of 25–31 °C. All temperature jump experiments involve 1:1 v/v dilution. Also shown (grey line) is the dynamic trace obtained upon stopped-flow 1:1 v/v dilution at 20 °C. The final total surfactant concentration is 10 mM.

![Fig. 3](image_url)  
**Fig. 3** Final temperature dependence of relaxation times obtained from the single exponential fitting of stopped-flow dynamic traces shown in Fig. 2.
changes in scattered light intensity. Moreover, the final scattered intensities increase with final temperatures, suggesting an increase in the number density of the aggregates (cylindrical micelles and vesicles). It should be noted that at a final temperature range of 33–43 °C, vesicles are not the dominating morphology and most of the aggregates exist in the form of cylindrical micelles.

The time dependence of scattered light intensity $I_t$ can be converted to a normalized function, namely, $(I_{\infty} - I_t)/I_{\infty}$ versus $t$, where $I_{\infty}$ is the value of $I_t$ at an infinitely long time. All dynamic traces at the final temperature range of 35–43 °C in Fig. 4a can be fitted with a single exponential function:

$$\frac{I_{\infty} - I_t}{I_{\infty}} = ce^{-t/\tau}$$

where $c$ is the normalized amplitude, $\tau$ is the relaxation time associated with a specific kinetic process. Fig. 5a shows a typical fit of the dynamic trace upon stopped-flow T-jump from 20 °C to 37 °C, resulting in a relaxation time, $\tau$, of 315 s. The quality of the fit is assessed from the reduced $\chi^2$ error values, which is defined by

$$\chi^2 = \frac{1}{N} \sum_{i=1}^{N} \left(x_i - \bar{x}_i\right)^2$$

where $N$, $x_i$, and $\bar{x}_i$ are the number of data points, and values of the experimental data and fitting data, respectively. For a temperature jump from 20 °C to 37 °C, the $\chi^2$ error value for the single-exponential fitting is 0.49, and a double-exponential fitting was also tested. The fitting was not significantly improved via the introduction of the second exponential expression as $\chi^2$ decreased only by 15%. The single exponential fitting results of dynamic traces recorded at final temperatures in the range of 35–43 °C are plotted in Fig. 5b. Apparently, the formation of final vesicles is much faster at higher final temperatures. The obtained relaxation times vary in the range of 82–440 s, which decrease dramatically with increasing temperatures.

**T-jump from 20 °C to final temperatures in the range of 45–55 °C**

Upon jumping from 20 °C to final temperatures in the range of 45–55 °C, relaxation processes with quite large positive amplitudes are observed. Fig. 6 shows the time dependence of scattered light intensity obtained for SDS/DEAB aqueous mixture upon stopped-flow temperature jump from 20 °C to the final temperature range of 45–55 °C. It seems that the final scattered light intensities remain nearly constant upon further heating. Fig. 7 shows the final temperature dependence of the scattered light intensity recorded upon a stopped-flow temperature jump from 20 °C to varying final temperatures in the range of 33–55 °C. The final scattered light intensity continuously increases up to 45 °C and then stabilizes out. Above 45 °C, cylindrical micelles completely transform into vesicles, which is the dominant morphology.

It is worth noting that in Fig. 7, the final scattering intensity upon T-jump remains almost constant above ~45 °C, whereas in Fig. 1 the optical transmittance is still decreasing above
Fig. 6 Time dependence of scattered light intensities obtained for SDS/DEAB mixed aqueous solutions ([SDS]/[DEAB] = 2/1) upon a stopped-flow temperature jump from 20 °C to varying final temperatures in the range of 45–55 °C. All temperature jump experiments involve 1:1 v/v dilution. The final total surfactant concentration is 10 mM.

45 °C. As for the observed apparent discrepancy, we tentatively interpret it as follows: (1) all kinetic data shown in Fig. 7 results from a T-jump from 20 °C to varying final temperatures (45–55 °C), the fast T-jump (within a millisecond) results in different kinetic pathways towards the formation of vesicles as compared to the slow heating rate; whereas in Fig. 1, the temperature is gradually increasing and at every temperature point ~20 min is required for the thermal equilibration. Thus, the different thermal history during the micelle-to-vesicle transition should contribute to the observed discrepancy in final scattering intensities; (2) for the kinetic data shown in Fig. 6, the measuring time window is 0–1500 s; note that the scattering intensity is still further increasing at extended time periods during the process forming final equilibrium vesicles, and sometimes, this process might take a few days or weeks.

Single- and double-exponential fitting results of dynamic traces upon stopped-flow T-jump from 20 °C to 55 °C are shown in Fig. 8. It is found that the single exponential function cannot fit the relaxation curve very well (Fig. 8a), especially for the first 10 s; the associated \( \chi^2 \) error is \( \approx 17.6 \). However, the dynamic trace can be fitted by a double exponential function (Fig. 8b, eqn (2)), resulting in a prominent improvement of \( \chi^2 \) of 0.61:

\[
\frac{(I_{\infty} - I_d)}{I_{\infty}} = c_1 e^{-t/t_1} + c_2 e^{-t/t_2}
\]

where \( c_1 \) and \( c_2 \) are the normalized amplitudes \( (c_2 = 1 - c_1) \), \( t_1 \) and \( t_2 \) are the relaxation times for two processes, \( t_1 < t_2 \). The mean relaxation time for the overall micelle formation, \( t_f \), can be calculated as

\[
t_f = c_1 t_1 + c_2 t_2
\]

Both processes associated with \( t_1 \) and \( t_2 \) possess positive amplitudes. For a stopped-flow temperature jump experiment from 20 °C to 55 °C, \( t_1 \) and \( t_2 \) are approximately 5.2 s and 188 s, respectively. The calculated \( t_f \) based on eqn (3) is \( \approx 73 \) s. The fitting results are summarized in Table S1 in the ESI.

Fig. 9 shows the double exponential fitting results of dynamic traces in Fig. 6. The relaxation time, \( t_1 \), is in the range of 5.2–28.5 s, and decreases with increasing temperatures; \( t_2 \) ranges from 188 to 694 s, and also decreases with increasing temperatures. The calculated \( t_f \) based on eqn (3) is in the range of 73–162 s. We can apparently judge that the higher the final temperatures, the faster the growth rate of the aggregates. On the other hand, the amplitudes associated with the two processes, \( c_1 \) and \( c_2 \), exhibit distinctly different final temperature dependences, with the former \((c_1, 0.58–0.88)\) decreasing with increasing temperatures, and the latter \( c_2 \) (0.12–0.42) increasing with an increase in the final temperatures.

Fig. 7 Final temperature dependence of scattered light intensities for SDS/DEAB mixed aqueous solutions ([SDS]/[DEAB] = 2/1) recorded 1500 s after a stopped-flow temperature jump from 20 °C to varying final temperatures in the range of 33–55 °C. The final total surfactant concentration is 10 mM.

Fig. 8 (a) Single and (b) double exponential fitting results of the dynamic trace recorded for SDS/DEAB aqueous mixture ([SDS]/[DEAB] = 2/1) upon stopped-flow temperature jump from 20 to 55 °C. The final total surfactant concentration is 10 mM.
Thermo-induced micelle-to-vesicle transition mechanisms

The thermo-induced morphological transformation between micelles and vesicles in SDS/DEAB mixed surfactant solutions can be explained by the well-known theory of the packing parameter \( p \), proposed by Mitchell et al. \cite{Mitchell1993} and Israelachvili et al. \cite{Israelachvili1991, Mitchell1993}. For spherical micelles, \( 0 \leq p \leq 1/3 \), and for cylindrical micelles, \( 1/3 \leq p \leq 1/2 \). The packing parameter \( p \) can be expressed as \( v/a_0 l_c \), where \( v \) is the surfactant tail volume, \( l_c \) is the tail length, and \( a_0 \) is the equilibrium area per molecule at the aggregate surface. With the increase of temperatures, it is expected that the hydration of surfactant ionic head groups are progressively weakened, hence, the surface area of the micelle per surfactant ion occupied \( (a_0) \) is reduced. Correspondingly, an increase in the packing parameter \( p \) then leads to the formation of larger aggregates such as cylindrical micelles and vesicles.

Recent theoretical considerations and experimental results concerning the vesicle formation kinetics have established that the micelle-to-vesicle transition typically involves a series of kinetic steps: rapid formation of non-equilibrium mixed aggregates and floppy bilayer intermediates, growth of disk-like micelles and their enclosure into vesicle precursors, and the formation of final equilibrium vesicles. \cite{Cao2006, Huang2004, Huang2005, Huang2006, Huang2007, Huang2008}

Fig. 9 Double exponential fitting results of dynamic traces obtained upon stopped-flow temperature jump from 20 °C to varying final temperatures in the range of 45–55 °C. The experimental conditions are the same as those described in Fig. 6.

A closer examination of kinetic traces shown in Fig. 6 can reveal that all kinetic traces recorded upon T-jump from 20 °C to the final temperature range of 45–55 °C apparently exhibit two-stage characteristics, i.e., the initial fast increase in scattering intensity followed by the subsequent much slower process. All relaxation curves can be fitted with double-exponential functions. This is drastically different to those occurring at lower final temperatures (35–43 °C). We ascribe the fast process \( (\tau_1 \sim 5.2–28.5 \text{ s}) \) to the formation of non-equilibrium precursor vesicles, and the slow process \( (\tau_2 \sim 188–694 \text{ s}) \) to their further growth into final equilibrium vesicles. Both relaxation times, \( \tau_1 \) and \( \tau_2 \), exhibit a considerable decrease with an increase of final temperatures (Fig. 9). As for the relaxation process associated with \( \tau_1 \), we further plotted the temperature-dependence of \( \tau \) and \( \tau_1 \) at a final temperature range of 35–55 °C (Fig. S1†). To our surprise, all the data points constitute into a smooth transition curve, which continuously decreases with increasing final temperatures. This further supports the theory that the kinetic process associated with \( \tau_1 \) in the range of 45–55 °C is of similar origin to that associated with \( \tau \) in the final temperature range of 35–43 °C.

In general, the slow relaxation process \( (\tau_2) \), the rate-determining step, can occur via the vesicle fusion/fission mechanism or unimer insertion/expulsion mechanism. In the final temperature range of 45–55 °C, all cylindrical micelles transform into vesicles, accompanied with a considerable increase in the number density of vesicles and the formation of vesicle aggregates. Thus, we propose that the slow process \( (\tau_2) \) should proceed via vesicle fusion/fission. \cite{Huang2004, Huang2005, Huang2006, Huang2007, Huang2008}

This is in agreement with the fact reported by Huang et al., \cite{Huang2004} who revealed that above 50 °C they can clearly observe the presence of vesicle aggregates with dimensions in the range of 200–300 nm. More intriguingly, we found that the amplitudes, \( c_1 \) and \( c_2 \) (associated with \( \tau_1 \) and \( \tau_2 \)), exhibit different temperature dependences, with the former decreasing.
and the latter increasing with final temperatures. The above analysis indicates that the vesicle formation pathways are highly dependant on the final temperatures. Below and above the critical temperature of 45 °C, the vesicle growth transforms from the unimer insertion/expulsion mechanism to the fusion/fission mechanism. Theoretically, the detailed vesicle growth mechanism can be differentiated via the concentration dependences of associated relaxation times \(t_{1}, t_{2}\). However, in the current case, the critical micelle-to-vesicle transition temperature varies dramatically with the total surfactant concentrations.

**Kinetics of thermo-induced vesicle-to-micelle transition**

The temperature dependent optical transmittance measurements shown in Fig. 1 suggest that upon cooling, the SDS/DEAB aqueous mixture exhibits the reverse vesicle-to-micelle transition. We further employed the stopped-flow \(T\)-jump technique to investigate the associated kinetic processes. It should be noted that the samples for the investigation of vesicle-to-micelle transition kinetics were subjected to \(~15 \text{ days storage prior to the measurements, allowing them to reach the final equilibrium. Fig. 10 shows the time dependence of scattered light intensities recorded for the SDS/DEAB aqueous mixture upon stopped-flow \(T\)-jump from varying initial temperatures (50 °C, 45 °C, and 38 °C) to 20 °C. All kinetic studies are associated with 1/2 v/v dilutions with the final total surfactant concentration being 10 mM. The grey line is a dynamic trace obtained upon a stopped-flow 1:2 v/v dilution at 50 °C, which exhibits an almost straight line. This indicates that no apparent microstructural changes can be discerned with the stopped-flow technique within the 1/2 v/v dilution process.

Upon a temperature jump from 50, 45, 38 °C to 20 °C, again, almost straight lines with much lower scattered intensities compared to that at 50 °C are observed. These results reveal that the reverse vesicle-to-micelle transition process is extremely rapid, and that most of the kinetic events complete within the stopped-flow dead time (a few millisecond). However, a closer examination of the dynamic traces (see inset in Fig. 10) still reveals the decrease of scattered light intensities, though quite small, within the time window of 0–800 s. The observed decrease in scattered intensity should be ascribed to the later stage during the process of thermo-induced transition from vesicles to cylindrical micelles. Within the stopped-flow dead time, all vesicles quickly disintegrate into small aggregates, and the subsequent slow process with relaxation time in the order of a few hundred seconds is associated with the structural rearrangement of small aggregates into cylindrical micelles.

**Conclusions**

The stopped-flow temperature jump technique was applied to probe the kinetics of thermo-induced micelle-to-vesicle and vesicle-to-micelle transitions in the SDS/DEAB catanionic surfactant system. The obtained temporal variations in scattered light intensities during microstructural changes were employed to elucidate the underlying transition mechanism. We conclude that kinetic sequences associated with the thermo-induced micelle-to-vesicle transition are highly dependant on the final temperatures. If the final temperature is in the range of 33–43 °C, the obtained dynamic traces can be fitted by single exponential functions, and the obtained relaxation times \(t\) decrease with an increase in the final temperatures. We tentatively ascribe this process to the transformation of floppy bilayer structures (which quickly form upon \(T\)-jumps) into precursor vesicles, accompanied with their further growth into final equilibrium vesicles. Considering the relatively low number density of precursor vesicles, the vesicle growth process should mainly proceed via the exchange and insertion/expulsion of surfactant monomers. In the final temperature range of 45–55 °C, vesicles are the predominant morphology. All dynamic traces can only be fitted with double exponential functions, yielding two relaxation times \(t_{1}\) and \(t_{2}\), which again exhibit a considerable decrease with an increase of the final temperatures. The fast process should be attributed to the formation of non-equilibrium precursor vesicles, and the slow process is associated with their further growth into final equilibrium vesicles via the fusion/fission mechanism. On the other hand, the reverse vesicle-to-micelle transition process induced by stopped-flow \(T\)-jump from elevated temperatures to 20 °C occurs quite fast and most of the kinetic events complete within the stopped-flow dead time (\(~2–3 \text{ ms}\)). We can only discern a slow relaxation process (with the relaxation time on the order of a few hundred seconds) at the later stage of thermo-induced vesicle-to-micelle transition.

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