


Cite this: *RSC Adv.*, 2020, 10, 34161

# Nanoparticle induced formation of self-assembled zwitterionic surfactant microdomains which mimic microemulsions for the *in situ* fabrication and dispersion of silver nanoparticles†

Taichi Nakagawa,<sup>a</sup> Akihito Hibara,<sup>b</sup> Willie L. Hinze<sup>c</sup> and Yoshitaka Takagai<sup>id</sup>\*<sup>a</sup>

The effective synthesis of highly concentrated and stable spherical silver nanoparticles (Ag-NPs) enclosed within microdomains formed by the Ag-NP induced self-assembly of zwitterionic surfactants is described. In this paper, the rapid and efficient synthesis of dispersed and stable Ag-NPs in zwitterionic surfactants, such as 3-(nonyldimethylammonio)- or 3-(decyldimethyl-ammonio)-propyl sulfate (C<sub>9</sub>- or C<sub>10</sub>-APSO<sub>4</sub>) that self-assemble in the presence of the Ag-NPs to form microdomains akin to microemulsions (ME) without the need of any organic solvent, co-surfactant or polymer is presented. Essentially all (>99.9%) of the synthesized Ag-NPs (TEM diameter 16.8 ± 4.3 nm) were incorporated and dispersed within the ME microdomains (213.5 ± 48.0 nm). A mechanism for formation of ME and its enclosing of the Ag-NPs in the ME is proposed.

Received 7th August 2020

Accepted 10th September 2020

DOI: 10.1039/d0ra06824a

rsc.li/rsc-advances

## 1. Introduction

Self-assembling nanoparticles (NPs) on/in confined microdomains, such as microparticles, microdroplets and emulsions have been intensely researched<sup>1</sup> owing to their attractive chemical properties which improve the effectiveness of the resultant materials to function as catalysts,<sup>2</sup> sensors,<sup>3</sup> and medical agents.<sup>4</sup> The synthesis of NPs on surfaces or inside microdomains can have a positive influence on properties such as the reaction speed,<sup>2</sup> efficiency,<sup>5</sup> sensitivity,<sup>6</sup> and enhance phenomenon such as emission.<sup>7</sup> In particular, numerous reports focus on applications of silver (Ag)-NPs synthesized using microemulsion technology in various fields<sup>8,9</sup> such as catalysis, nanoreactors, biology, medicine, optics, and electronics among others. Although it is difficult to control the size and homogeneity of Ag-NPs prepared by chemical reduction methods,<sup>10</sup> utilization of surfactant oil-in-water (o/w) or water-in-oil (w/o) emulsion systems have proven

beneficial in the synthesis of small and uniform Ag-NPs which were formed within the emulsion microdroplets.<sup>8,9</sup> Bulavchenko *et al.* reported that the fabrication efficiency of Ag-NPs was enhanced by utilizing microemulsions relative to larger emulsions.<sup>11</sup> Thus far, the microemulsion systems employed have been composed of water, water-immiscible organic solvent (oil), surfactant and in some cases, a co-surfactant or polymer.<sup>8,9</sup>

Herein, we report the use of simple aqueous solutions of zwitterionic surfactants, such as 3-(nonyldimethylammonio)- or 3-(decyldimethyl-ammonio)-propyl sulfate (C<sub>9</sub>- or C<sub>10</sub>-APSO<sub>4</sub>), for the efficient synthesis of Ag-NPs. At concentrations greater than their critical micelle concentration (CMC), surfactants of this class are thermo-responsive and exist as a homogeneous aqueous micellar solution at temperatures above their upper critical solution temperature (UCST), whereas they phase separate at temperatures below the UCST to form a small surfactant-rich phase and a larger aqueous surfactant-poor phase.<sup>12–14</sup> In this study, the surfactant concentrations were below the CMC values and at temperatures such that homogeneous solutions were formed in which the surfactant was initially present in its monomeric (molecular) form. During the course of the synthetic reaction at elevated temperature, these surfactants undergo induced self-assembly to form microdomains (ME) in which the Ag-NPs are sequestered and dispersed in a manner akin to that observed in o/w microemulsions or swollen micelles except that no added organic solvent, co-surfactant or polymer is required.

<sup>a</sup>Faculty of Symbiotic Systems Science, Cluster of Science and Technology, Fukushima University, 1 Kanayagawa, Fukushima 960-1296, Japan. E-mail: s015@ipc.fukushima-u.ac.jp

<sup>b</sup>Institute of Multidisciplinary Research for Advanced Materials, Tohoku University, 2-1-1 Katahira, Aoba-ku, Sendai 980-8577, Japan

<sup>c</sup>Department of Chemistry, Wake Forest University, P. O. Box 7486, Winston-Salem, North Carolina 27109, USA. E-mail: hinze@wfu.edu

† Electronic supplementary information (ESI) available: The elemental mapping; the impact of citric acid concentration; absorption spectra; the relationship between the surfactant concentration and the volume of second phase; relationship between the conductivity and the concentration of C<sub>10</sub>-APSO<sub>4</sub> surfactant; the relationship between the C<sub>10</sub>-APSO<sub>4</sub> surfactant and conductivity. See DOI: 10.1039/d0ra06824a



## 2. Experimental

### 2.1. Instrumentation

Ultraviolet-visible spectra were obtained using a JASCO V-650 double beam spectrophotometer equipped with quartz cuvette (1 cm × 1 cm). The sample solution was measured without dilution. Dynamic light-scattering (DLS) measurements were performed using a HORIBA SZ-100 nanoparticle analyzer with a fluorescence analytical grade quartz cuvette (1 cm × 1 cm) at room temperature (25.0 °C) and green laser output at 532 nm and the measurement scattering angle set to 90°. DLS measurement of particle size is strongly influenced by the viscosity of sample solution. Thus, a sine-wave vibro viscometer SV-A (A&D Co. Ltd, Tokyo, Japan) was used to measure the viscosity of the sample solutions prior to the DLS measurement. The viscosity was then imputed as the parameter of DLS measurement. The hydrodynamic diameters reported are the mean diameters. Transmission electron microscopy (TEM) images were obtained using a JEOL JEM-1400plus transmission electron microscope (TEM; Tokyo, Japan) operating at 120 kV. The HD-2700 (Hitachi, Ltd) microscope was employed for the scanning TEM (STEM) analysis. The acceleration voltage was 200 kV at magnification between 30 000 and 100 000. The Octane T Ultra (AMETEK Co., Ltd) energy-dispersive X-ray (EDX) analyzer was used, which was connected to the STEM apparatus. TEM samples were prepared by placing 5.0 µL aliquots of the aqueous sample onto a carbon-filmed 200-mesh Cu grid (Nishin EM Co. Ltd, Tokyo, Japan). After sample loading, the grids were dried at room temperature prior to the TEM measurements. The microemulsion size distribution was typically determined by measuring 20 such grids. Nanoparticles size distribution and concentration measurements were made using a ViewSizer3000 (HORIBA Ltd, Japan). The dimension of the cell was 10 × 10 × 45 mm with four-plane transmission (Hella Analytics, Germany). The measurement conditions were set temperature 22 °C, 3 lasers (450, 520, 650 nm) were used with the scattering angle set to 90°. The zeta potential measurements were made employing a Nano Z ZEN3600 Zetasizer (Malvern Instruments Ltd, Worcestershire, UK). The pH measurements were made using a HORIBA handy D-50 pH meter equipped with a long ToupH combination 9680-10D electrode. This electrode probe was employed to directly measure solution pH in the conical-shaped centrifuge tube. The conductivity measurements were made using a HORIBA handy D-74T water quality meter with a Conductivity Cell (Submersible Type) 9382-10D electrode. This electrode was employed to directly measure solution conductivity in the centrifuge tube.

### 2.2. Materials

All chemicals and reagents were used without any further purification, unless otherwise noted. *N,N*-Dimethyldecylamine (>95.0%) was purchased from Tokyo Chemical Industry Co., Ltd (Japan). Trimethylene sulphate (98%) was obtained from Combi-Blocks, Inc. (USA). Acetone (≥99.5%), trisodium citrate dihydrate (≥99.0%), sodium hydroxide (≥97.0%) and diethyl ether (≥99.5%) were acquired from FULIFILM Wako Chemicals

Co. (Japan). Silver nitrate (≥99.0%) was obtained from Sigma-Aldrich (USA). Distilled water was obtained using a water purification apparatus RFD240NA (ADVANTEC, Japan).

The zwitterionic surfactants, C<sub>10</sub>-APSO<sub>4</sub>, C<sub>9</sub>-APSO<sub>4</sub> and C<sub>8</sub>-APSO<sub>4</sub> were synthesized using published literature procedures.<sup>12,14</sup>

### 2.3. Procedures

**2.3.1. Determination of the zeta potential.** The zeta-potential of the surfactants was determined by an extrapolation method. Namely, the surfactants were diluted with a water-methanol mixed solvent system containing different amounts of methanol and for each, the zeta potential was measured. A plot of the apparent zeta potential *versus* the methanol content was constructed and found to be linear. The zeta potential of the surfactant was taken to be its extrapolated value for 0% methanol.

**2.3.2. Determination of the CMC.** The CMC of the surfactant was determined by measuring the electrical conductivity of a series of aqueous surfactant solutions which contained different concentrations (*C*) of the surfactant. A graph of molar conductivity *versus*  $\sqrt{C}$  was made and the CMC was determined from the inflection point of the curve.

**2.3.3. Ag-NP synthesis.** The recipe for synthesis of Ag-NPs in MEs was as follows: typically, 1.00 mL of an aqueous 0.50 w/w% zwitterionic surfactant C<sub>9</sub>- or C<sub>10</sub>-APSO<sub>4</sub>, 0.50 mL of 5.0 mM trisodium citrate aqueous solution were mixed together in a conical centrifuge tube (tube) and heated in an oil bath (108 °C) for 2 min to prepare a homogeneous solution. The resultant solution was adjusted to pH 11 by the addition of 0.25 mL of 0.10 M NaOH solution with continued heating in an oil bath (108 °C) for 2 min. Next, 0.50 mL of 0.50 mM AgNO<sub>3</sub> solution was added to the tube and the resultant solution kept static at 108 °C for 8 min (for synthesis of Ag-NPs). This solution was then cooled for 5 min down to room temperature (25 °C). The solution was characterized by UV-visible absorption spectrophotometer, DLS, TEM, and STEM with EDX. To count and measure the NPs, a NP tracking analyzer (NTA) was employed.

## 3. Results and discussion

### 3.1. Synthesis and characterization

The Ag-NPs prepared in the ME appeared as a yellow homogeneous solution (see inset to Fig. 1A) and exhibited a surface plasmon resonance (SPR) absorption band at 401.5 nm (Fig. 1A solid red line) with the peak width at half maximum (PWHM) of 95 nm. The TEM images indicate that the Ag-NPs are monodisperse and roughly spherical-shaped in the ME, which is composed of the surfactant C<sub>10</sub>-APSO<sub>4</sub> (Fig. 1B). Furthermore, STEM-EDX mapping (Fig. S1 in ESI†) confirmed that the Ag-NPs were dispersed in the C<sub>10</sub>-APSO<sub>4</sub> surfactant ME microdomains. The mean diameter of the Ag-NPs in the ME was 16.8 ± 4.3 nm (*n* = 50) as determined from the TEM images, whereas the average hydrodynamic diameter obtained from DLS was 55.3 ± 46.3 nm (Fig. 1C). Essentially all (>99.9%) of the synthesized Ag-NPs were incorporated and dispersed within the ME. The size of



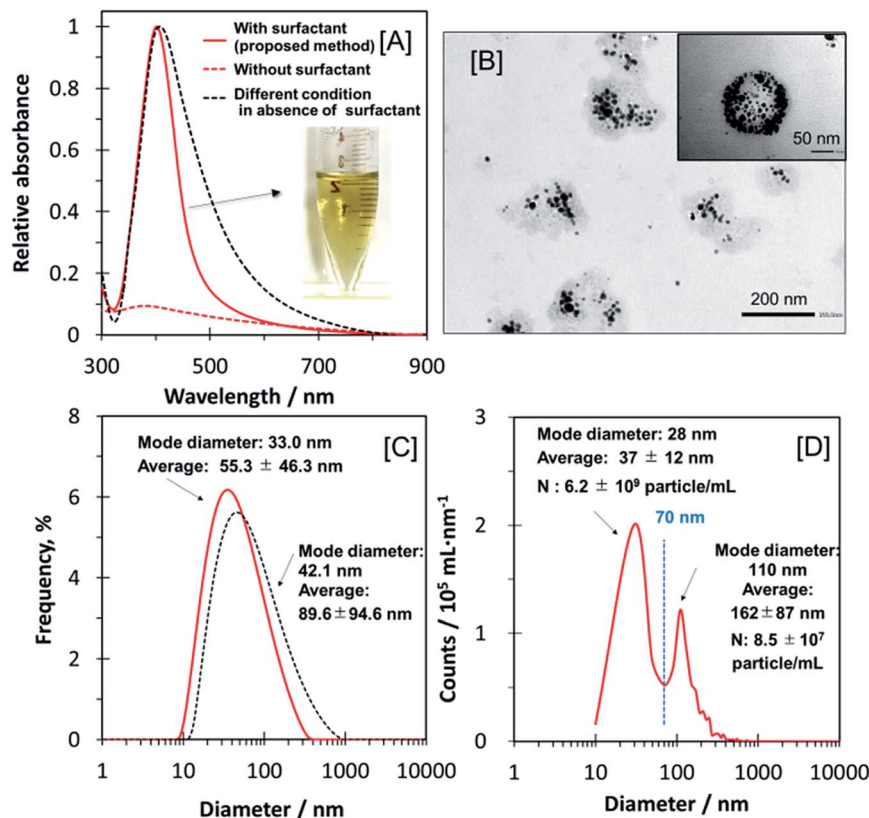


Fig. 1 Characterization of the Ag-NPs in ME. Panel [A] shows a photograph of the solution containing Ag-NPs and the UV-visible absorption spectra of Ag-NPs prepared by this method in the presence (solid red line) and absence of the surfactant (red dotted line). The black dotted line is spectra of Ag-NPs obtained using a typical citric acid reductant method.<sup>17</sup> Panel [B] shows transmission electron microscopy (TEM) image of Ag-NPs in the (ME). Panel [C] shows the dynamic light scattering (DLS) profile of the Ag-NPs prepared by this method (red line) and a typical citric acid reductant method<sup>17</sup> (black dotted line). Panel [D] shows the NP tracking analyzer (NTA) profile which indicates the presence of both the Ag-NPs (mode diameter 28 nm) and the ME (mode diameter 110 nm). Experimental conditions:  $[C_{10}\text{-APSO}_4]_T = 0.20\%$  (w/w);  $[AgNO_3]_T = 0.11$  mM;  $[trisodium\ citrate]_T = 1.10$  mM;  $[NaOH]_T = 0.010$  mM; total volume = 2.25 mL; pH = 11; reaction time: 8 min at 108 °C. TEM voltage 120 kV (30k to 50k magnification).

the ME microdomains was determined as  $213.5 \pm 48.0$  nm ( $n = 25$ ; dry state) from the TEM image. On conducting the synthetic protocol in the absence of  $C_{10}\text{-APSO}_4$ , flocculation (aggregation) of the Ag-NPs occurred (Fig. 1A dotted red line). We propose that the surfactant adsorbs onto the surface of Ag-NPs providing a capping layer, which stabilizes the Ag-NPs and mediates their growth.<sup>8</sup> In the absence of  $C_n\text{-APSO}_4$  surfactant, no such confinement of growth occurs and the resultant NPs are not uniform and unstable. A typical Ag-NP synthesis by the citric acid method (without surfactant) was reported as yielding a range of Ag-NP sizes (10–200 nm) with their SPR bands ( $\lambda_{\max}$ ) ranging from 397 to 427 nm.<sup>12</sup> For comparison, the previously PWHM (near 130 nm) and size distribution ( $89.6 \pm 94.6$  nm)<sup>15</sup> which are larger than those of the Ag-NPs synthesized in this work (Fig. 1A and C dotted black lines). The smaller number of  $\lambda_{\max}$  and PWHM represents a smaller size and more uniform size distribution for the Ag-NPs.<sup>15,16</sup>

The NTA profile (Fig. 1D) indicates the presence of two major size distributions ( $37 \pm 12$  and  $162 \pm 87$  nm). The two peaks correspond to the values obtained from DLS (33 nm, modal Ag-NP diameter) and TEM ( $213.5 \pm 48.0$  nm as dried ME). As ME

liquid droplets were dried for TEM observation, the apparent diameter (after drying) in the TEM image is larger than that of the dispersive state in NTA. This method of synthesizing Ag-NPs in an ME serves to regulate the size (57.7 to 72.1 nm; described in the Section 3.2) and shape (spherical) of the NPs. NTA indicated that the number of Ag-NPs and ME clusters were  $6.2 \pm 10^9$  and  $8.5 \pm 10^7$  particle per mL, respectively. The resultant concentration for Ag-NPs in ME was 73 particle/ME, whereas the mean apparent concentration of the Ag-NPs in ME was  $15 \pm 10$  particle/ME ( $n = 50$ ) as determined from TEM images (max. frequency).

### 3.2. Effect of the concentration of $Ag^+$ ion

Fig. 2 shows the relationships between the concentration of  $Ag^+$  and the diameter of the Ag-NPs and their concentration in the ME. Although the formed Ag-NPs (average size) only slightly decreased (from 72.1 to 57.7 nm) with increases in the silver ion concentration (Fig. 2 blue line), the uncertainties associated with these values were ca. 25%. Thus, there was no appreciable difference in the size of Ag-NPs with variations in the concentration of  $Ag^+$  ion. Thus, this method can synthesize a specific





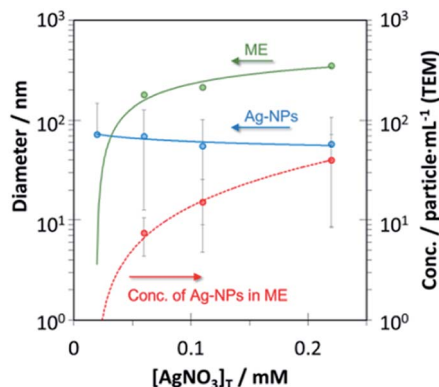


Fig. 2 Influence of the variation of  $\text{Ag}^+$  concentration upon the Ag-NP particle size (blue line) and surfactant ME diameter (green line) as well as the Ag-NP concentration in the ME (red line). Experimental conditions:  $[\text{C}_{10}\text{-APSO}_4]_{\text{T}} = 0.20\%$  (w/w);  $[\text{trisodium citrate}]_{\text{T}} = 1.10$  mM;  $[\text{NaOH}]_{\text{T}} = 0.010$  mM; total volume = 2.25 mL; pH = 11; reaction time: 8 min at  $108^\circ\text{C}$ . Diameters and concentrations determined from replicate ( $n = 10$ ) DLS and TEM measurements, respectively.

sized Ag-NP in the  $\text{C}_{10}\text{-APSO}_4$  ME. This is in contrast to a literature report in which the Ag-NPs size typically increased with an increase in  $[\text{Ag}^+]$ .<sup>8</sup> The concentration of Ag-NPs in the ME increases with concentration of  $\text{Ag}^+$  (Fig. 2 red line). At concentrations  $> 0.22$  mM  $\text{AgNO}_3$ , formation of agglomerated or aggregated Ag-NPs resulted owing to saturation. The ME size is also impacted by the silver ion concentration (Fig. 2, green line). Over the concentration range examined, the size of ME increases from *ca.* 100 to 200 nm with increases in  $[\text{Ag}^+]$ . This result suggests that the presence of  $\text{Ag}^+$  is an important factor in the formation of the ME. No ME formation was observed in a mixture of  $\text{Ag}^+$  and  $\text{C}_{10}\text{-APSO}_4$  surfactant without citric acid (reducing agent). This indicates that the formation of the surfactant ME microdomains is related to the presence of Ag-NP precursor. The precursor functions to induce assembly of surfactant monomers initially present in solution to form aggregates with microdomains which mimic the behavior of a microemulsion (ME). It is important to note that no such ME formed if pre-prepared citrate coated Ag-NPs were used instead of  $\text{Ag}^+$  (*i.e.*  $\text{AgNO}_3$ ) in the described procedure as the Ag-NPs existed in an as-dispersed state in solution. It is considered that rapid exchange of  $\text{C}_n\text{-APSO}_4$  surfactant with the pre-coated citric acid Ag-NPs is difficult resulting in the absence of a surfactant capping layer on the Ag-NPs.

### 3.3. Impact of the concentration of citrate ions and pH

The effect of the citrate ion concentration upon the size and concentration of the Ag-NPs in the ME was briefly examined. The results (Fig. S2 in ESI<sup>†</sup>) indicate that at citrate concentrations  $> ca.$  1.0 mM, the concentration of Ag-NPs in the ME noticeably decreased and their size increased (due to aggregation). The results suggest that citrate functions as a reductant; however, at higher concentrations, it is not only a reductant but also a dispersing agent. The pH impact on the fabrication of both Ag-NP and ME is shown in Fig. S3 in ESI<sup>†</sup>. The formation of

Ag-NPs in the ME was only observed at pH 11. Dong *et al.* reported that the fabrication rate of Ag-NPs at pH 11 was faster than other pH values (citric acid reduction method) owing to the reduction activity.<sup>17</sup> Perhaps the rate of the surfactant binding onto the precursor of Ag-NPs is faster than that of citric acid (as capping agent). At pH values  $> 11$  Ag-NP, synthesis is difficult due to formation of  $\text{AgOH}$  ( $K_{\text{sp}} = 2 \times 10^{-8}$ ).<sup>18</sup>

### 3.4. Effect of surfactant and impact on ME production

The preparation of dispersed Ag-NPs in the ME was successful using both  $\text{C}_9\text{-}$  and  $\text{C}_{10}\text{-APSO}_4$  as surfactants (Fig. 3). The number of Ag-NPs in the ME was essentially the same ( $15 \pm 4$  for  $\text{C}_9\text{-APSO}_4$  and  $15 \pm 10$  for  $\text{C}_{10}\text{-APSO}_4$ ). The diameter of the Ag-NPs in the  $\text{C}_9\text{-APSO}_4$  and  $\text{C}_{10}\text{-APSO}_4$  MEs were also very similar. The ME size was  $100 \pm 31$  nm (from TEM image) for  $\text{C}_9\text{-APSO}_4$  which is smaller than that of  $\text{C}_{10}\text{-APSO}_4$ . However, use of the shorter carbon chain surfactant, ( $\text{C}_8\text{-APSO}_4$ ), only resulted in the formation of Ag-NP aggregates (Fig. 3C). It should be noted that the diameters of the  $\text{C}_9\text{-}$  and  $\text{C}_{10}\text{-APSO}_4$  ME aggregates containing the dispersed Ag-NPs are much greater relative to normal micelles formed from similar zwitterionic sulfobetaine surfactants.<sup>19,20</sup> For instance, the surfactant SB3-10 (same formula as  $\text{C}_{10}\text{-APSO}_4$  except the terminal anionic group is sulfonate,  $-\text{SO}_3^-$ , rather than sulfate) has a diameter of only 2–4 nm.<sup>19,20</sup>

Although a  $\text{C}_n\text{-APSO}_4$  surfactant is neutral overall, the zeta-potential of the solution including this surfactant at pH 3 (without Ag-NPs) was determined to be +23.2, +10.2 and  $-20.0$  mV for  $\text{C}_8\text{-}$ ,  $\text{C}_9\text{-}$ , and  $\text{C}_{10}\text{-APSO}_4$  respectively. These values are in the range of reported values for other zwitterionic surfactant systems which can range from slightly positive to  $-70$  mV depending upon the type and concentration of added salts and pH.<sup>20–23</sup> The bipolar ionic moiety of the zwitterionic surfactants can interact with the uncapped or citrate capped Ag-NPs and the values become more negative, *i.e.*,  $-89 \pm 12$  mV for  $\text{C}_{10}\text{-APSO}_4$  at pH 11. A similar phenomenon was reported for the potential of a NP-surfactant bilayer composite, in which the potential was gradually decreased by the surfactant-coating on metallic NPs from 0 mV to  $-57.9$  mV.<sup>24,25</sup> Taking into account the fact that the zeta potentials for zwitterionic surfactant systems (as well as citrate coated Ag-NPs themselves) decrease with increases in solution pH, the zeta potential obtained for the  $\text{C}_{10}\text{-APSO}_4/\text{Ag-NP}$  ME in this work is reasonable and in the general range reported for Au- and Ag-NPs dispersed or capped with other zwitterionic species.<sup>13,26,27</sup>

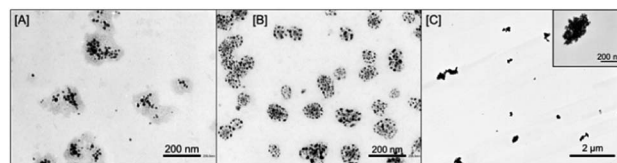


Fig. 3 TEM images of the Ag-NPs formed using three different zwitterionic surfactants in the synthetic procedure: [A]  $\text{C}_{10}\text{-APSO}_4$ . [B]  $\text{C}_9\text{-APSO}_4$ . [C]  $\text{C}_8\text{-APSO}_4$ .



The  $C_{10}$ -APSO<sub>4</sub> concentration range that allows formation of the ME for the synthesis of Ag-NPs as reported is rather limited. The upper concentration is dictated by the phase separation behavior (UCST) exhibited by this surfactant. As shown (Fig. S4 in ESI†), at concentrations  $\geq 0.6$  w/w% (18.5 mM), a heterogeneous solution results which phase separates (the volume of the surfactant-rich phase formed is indicated in the figure in the presence and absence of Ag-NPs). In addition, the concentration of the surfactant needs to be smaller than the CMC at the final temperature in the procedure (25 °C). The CMC was 0.24 w/w% (7.4 mM) at 25 °C (Fig. S5; green line in ESI†). In addition, the CMC value modestly increased with the formation of Ag-NPs as shown in Fig. 4, whereas the presence of Ag<sup>+</sup> (at any concentration) did not affect the CMC values (Fig. 4, supported by Fig. S6 in ESI†). The CMCs of a homologous series of zwitterionic surfactants in water are temperature dependent; namely, as the temperature is increased, an initial slight decrease in the CMC value is observed until a minimum value is reached after which the CMCs start to increase with temperature.<sup>28–30</sup> The CMC of  $C_{10}$ -APSO<sub>4</sub> increased with temperature (*i.e.* 0.24 w/w%, 0.38 w/w%, and 0.44 w/w% for 25 °C, 35 °C, and 42 °C, respectively; Fig. S5 in ESI†).

### 3.5. Mechanism

Fig. 5 illustrates a potential mechanism for fabrication of the Ag-NPs in the  $C_9$ - or  $C_{10}$ -APSO<sub>4</sub>. Initially, the surfactants exist in solution in monomeric form along with the Ag<sup>+</sup> ion reactant (stage I). As citrate ion is added, silver ion is converted to Ag-NPs to which the bipolar zwitterionic surfactant head group bind at a faster rate relative to the citrate ions (stage II). Several Ag-NPs can then interact with each other *via* hydrophobic interactions; *i.e.*, the alkyl chain of one surfactant coated Ag-NP interacts with that of another (or several) other such coated Ag-NPs (stage III). Lastly, the exterior of such self-assembled zwitterionic surfactant microdomains or clusters (*i.e.*, the alkyl chain of the coated Ag-NPs) can interact with the alkyl chain of the monomeric surfactants such that the self-assembled microdomains then has the bipolar zwitterionic group in contact with the bulk water solvent (stage IV). The formation of such zwitterionic surfactant

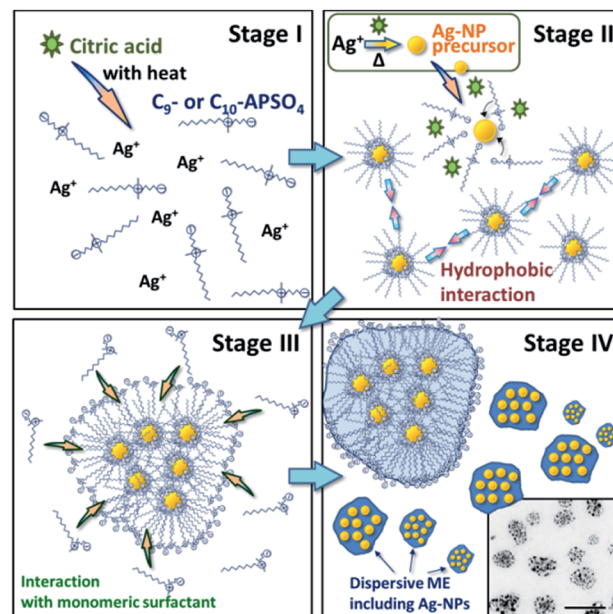


Fig. 5 Illustration of the possible mechanism of fabrication of dispersive ME enclosed Ag-NPs, prepared by following the synthetic procedure (see Section 2.3.3).

coated (monolayer or bilayer) individual metal NPs or citrate coated NPs has literature precedence.<sup>21–23,31</sup> In this work, it appears that the entire exterior of the  $C_{10}$ -APSO<sub>4</sub>-Ag-NP microdomains (clusters) are thus coated as well, mimicking an oil-in-water microemulsion system that houses the dispersed Ag-NPs.

## 4. Conclusions

In summary, the rapid and efficient synthesis of dispersed and stable Ag-NPs in zwitterionic  $C_n$ -APSO<sub>4</sub> surfactants that self-assemble in the presence of the Ag-NPs to form microdomains akin to microemulsions without the need of any organic solvent or co-surfactant was developed. Essentially all the Ag-NPs were present within the ME. The effects of silver ion concentration and pH upon the size and concentration of the Ag-NPs were assessed. In addition, the concentration of the surfactant, its zeta-potential, and the variation of CMC was investigated. The mechanism for the formation of ME and the enclosing of Ag-NPs in the ME was presented. It is thought that the Ag-NPs in the surfactant ME could potentially be employed as a suitable surfactant-based dispersion solution for different applications.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

The authors gratefully acknowledge funding from the Japan Atomic Energy Agency (JAEA) Nuclear Energy S&T and Human

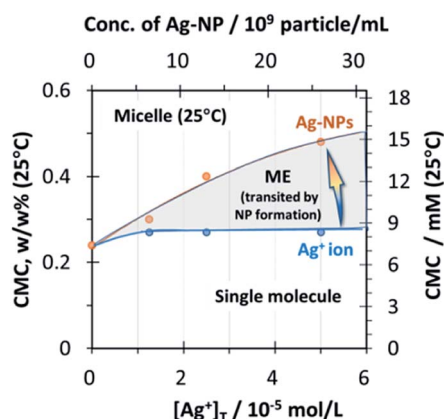


Fig. 4 The transition of critical micelle concentration (CMC) values in the presence of Ag<sup>+</sup> (blue line) and Ag nanoparticles (orange line).

Resource Development Project through concentrating wisdom Grant Number JPJA19H19210081 and Japan Society for the Promotion of Science (JSPS) Grant-in-Aid for Scientific Research(B) JP20H04352.

## Notes and references

- 1 N. Yan, Y. Zhu and W. Jiang, *Chem. Commun.*, 2018, **54**, 13183–13195.
- 2 Y. Zhao, X. Ni, S. Ye, Z.-G. Gu, Y. Li and T. Ngai, *Langmuir*, 2020, **36**, 2037–2043.
- 3 Y. Liu, A. Teitelboim, A. Fernandez-Bravo, K. Yao, M. V. P. Altoe, S. Aloni, C. Zhang, B. E. Cohen, P. J. Schuck and E. M. Chan, *ACS Nano*, 2020, **14**, 1508–1519.
- 4 F. Huang, Y. Gao, Y. Zhang, T. Cheng, H. Ou, L. Yang, J. Liu, L. Shi and J. Liu, *ACS Appl. Mater. Interfaces*, 2017, **9**, 16880–16889.
- 5 M. B. Schütz, K. Lê, S. Ilyas and S. Mathur, *Langmuir*, 2020, **36**, 1552–1558.
- 6 A. Del Tedesco, V. Piotto, G. Sponchia, K. Hossain, L. Litti, D. Peddis, A. Scarso, M. Meneghetti, A. Benedetti and P. Riello, *ACS Appl. Nano Mater.*, 2020, **3**, 1232–1241.
- 7 L. Bao, H. You, L. Wang, L. Li, R. Qiao, Y. Zhang, Y. Zhong, Y. Xiong and Z. Li, *J. Mater. Chem. C*, 2014, **2**, 8949–8955.
- 8 J. N. Solanki and Z. V. P. Murthy, *Ind. Eng. Chem. Res.*, 2011, **50**, 12311–12323.
- 9 M. Sanchez-Dominguez, K. Pemartin and M. Boutonnet, *Curr. Opin. Colloid Interface Sci.*, 2012, **17**, 297–305.
- 10 H. D. Beyene, A. A. Werkneh, H. K. Bezabh and T. G. Ambaye, *Sustain. Mater. Technol.*, 2017, **13**, 18–23.
- 11 A. I. Bulavchenko, A. T. Arymbaeva, M. G. Demidova, P. S. Popovetskiy, P. E. Plyusnin and O. A. Bulavchenko, *Langmuir*, 2018, **34**, 2815–2822.
- 12 T. Saitoh and W. L. Hinze, *Anal. Chem.*, 1991, **63**, 2520–2525.
- 13 Y. Takagai, R. Miura, A. Endo and W. L. Hinze, *Chem. Commun.*, 2016, **52**, 10000–10003.
- 14 P. G. Nilsson, B. Lindman and R. G. Laughlin, *J. Phys. Chem.*, 1984, **88**, 6357–6362.
- 15 N. G. Bastús, F. Merkoçi, J. Piella and V. Puentes, *Chem. Mater.*, 2014, **26**, 2836–2846.
- 16 L. Mulfinger, S. D. Solomon, M. Bahadory, A. V. Jeyarajasingam, S. A. Rutkowsky and C. Boritz, *J. Chem. Educ.*, 2007, **84**, 322.
- 17 X. Dong, X. Ji, H. Wu, L. Zhao, J. Li and W. Yang, *J. Phys. Chem. C*, 2009, **113**, 6573–6576.
- 18 The Chemical Society of Japan, *Handbook of Chemistry: Pure Chemistry*, Maruzen Publication, Tokyo, 5th edn, 2004.
- 19 J. Wang, O. Morales-Collazo and A. Wei, *ACS Omega*, 2017, **2**, 1287–1294.
- 20 D. P. Santos and R. L. Longo, *J. Phys. Chem. B*, 2016, **120**, 2771–2780.
- 21 F. D. Souza, B. S. Souza, D. W. Tondo, E. C. Leopoldino, H. D. Fiedler and F. Nome, *Langmuir*, 2015, **31**, 3587–3595.
- 22 J. P. Priebe, F. D. Souza, M. Silva, D. W. Tondo, J. M. Priebe, G. A. Micke, A. C. O. Costa, C. A. Bunton, F. H. Quina, H. D. Fiedler and F. Nome, *Langmuir*, 2012, **28**, 1758–1764.
- 23 C. Yao, P. Wang, L. Zhou, R. Wang, X. Li, D. Zhao and F. Zhang, *Anal. Chem.*, 2014, **86**, 9749–9757.
- 24 A. Sambasivam, A. V. Sangwai and R. Sureshkumar, *Langmuir*, 2016, **32**, 1214–1219.
- 25 E. K. Perttu and F. C. Szoka, *Chem. Commun.*, 2011, **47**, 12613.
- 26 Z. Qiao, Y. Yao, S. Song, M. Yin and J. Luo, *J. Mater. Chem. B*, 2019, **7**, 830–840.
- 27 T. C. Prathna, N. Chandrasekaran and A. Mukherjee, *Colloid. Surface. Physicochem. Eng. Aspect.*, 2011, **390**, 216–224.
- 28 N. Muller, *Langmuir*, 1994, **10**, 2202–2205.
- 29 E. Sikorska, D. Wyrzykowski, K. Szutkowski, K. Greber, E. A. Lubecka and I. Zhukov, *J. Therm. Anal. Calorim.*, 2016, **123**, 511–523.
- 30 C. Brinatti, L. B. Mello and W. Loh, *Langmuir*, 2014, **30**, 6002–6010.
- 31 H. D. Fiedler, E. E. Drinkel, B. Orzechowicz, E. C. Leopoldino, F. D. Souza, G. I. Almerindo, C. Perdona and F. Nome, *Anal. Chem.*, 2013, **85**, 10142–10148.

