

Cite this: *RSC Adv.*, 2019, 9, 6556

Aggregation behavior of cetyldimethylethylammonium bromide under the influence of bovine serum albumin in aqueous/electrolyte solutions at various temperatures and compositions: conductivity and molecular dynamics study†

Md. Farid Ahmed,^{ab} Mohammad Robel Molla,^{id ab} Mousumi Saha,^c Imrul Shahriar,^{id c} Mohammad Saidur Rahman,^{id ad} Mohammad A. Halim,^{id c} Malik Abdul Rub,^{id ef} Md. Anamul Hoque^{id *a} and Abdullah M. Asiri^{id ef}

Herein, we have investigated the interaction of bovine serum albumin (BSA), the most abundant globular protein, with a conventional cationic surfactant, cetyldimethylethylammonium bromide (CDMEAB), through a conductivity technique in the absence/presence of electrolyte solutions at various temperatures (298.15–323.15 K). The interaction of the protein with drugs/surfactants and other additives plays a crucial role in the body. Hence, the main concern of the study is to extract the impact of BSA on surfactant molecules and *vice versa*. From the specific conductivity *versus* concentration of surfactant plots, three different noticeable critical micelle concentration (c^*) values were obtained for pure CDMEAB and its mixture with protein/protein + salts. The presence of BSA and electrolytes altered the c^* values of CDMEAB revealing interactions among the studied constituents where the salt solutions reduced the c^* values and created a convenient environment for favorable micellization. The negative magnitudes achieved for standard free energy changes (ΔG_m^0) suggest spontaneity of micellization while the values of ΔH_m^0 and ΔS_m^0 signified the existence of some electrostatic and hydrophobic interactions. The values of molar heat capacity (ΔC_m^0) were positive as well as small which was an indication of less structural deformation. Molecular Dynamics (MD) simulation for all atoms revealed that the salt ions promoted non-covalent interaction between BSA and CDMEAB, and such interactions were not observed in the absence of the salt. Protein structure remained nearly same in spite of strong interaction with CDMEAB as evident from the overall RMSD (root-mean-square deviation) values of the alpha carbons and backbone of the protein and RMSF (root-mean-square fluctuation) values of the amino acid residues present in BSA. In this work thermodynamic parameters of transfer (such as $\Delta G_{m,tr}^0$, $\Delta H_{m,tr}^0$, and $\Delta C_{p,m,tr}^0$) were also evaluated and the results are discussed in detail. Besides, contributions of enthalpy and entropy to free energy changes were also analyzed.

Received 4th January 2019
Accepted 15th February 2019

DOI: 10.1039/c9ra00070d

rsc.li/rsc-advances

^aDepartment of Chemistry, Jahangirnagar University, Savar, Dhaka-1342, Bangladesh.
E-mail: ahoque_ju@yahoo.com; ahoque_ju@juniv.edu; Fax: +880-2-7791052; Tel: PABX: +880-2-7791045-51 extn 1437

^bBangladesh Council of Scientific and Industrial Research (BCSIR), Dhaka, Bangladesh
^cDivision of Quantum Chemistry, The Red-Green Research Centre, BICCB, Dhaka, Bangladesh

^dDepartment of Chemical Engineering, Faculty of Engineering, University of Malaya, 50603, Kuala Lumpur, Malaysia

^eChemistry Department, Faculty of Science, King Abdulaziz University, Jeddah-21589, Saudi Arabia

^fCenter of Excellence for Advanced Materials Research, King Abdulaziz University, Jeddah 21589, Saudi Arabia

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c9ra00070d

1. Introduction

Proteins are associated with various functions including transport, signaling, energy conversion, cell communication, catalysis, *etc.* in biological systems. Bovine serum albumin (BSA) along with other globular proteins has been stimulating a plethora of scientific investigations over the last few decades due to its typical architecture and better performance for carrier properties.¹ This protein is one of the most familiar carrier proteins, which play a crucial role in transport as well as the release of various endogenous & exogenous materials in blood.^{1,2} The biological activities and benefits of proteins can be realized by means of insight from their performance to act as



a hormone as well as transmitters in the central nervous system of living beings.³ BSA has the capability to bind with a diversified array of biologically active molecules (such as drugs, fatty acids, metabolites, amino acids, metals and various organic compounds). This affinity for binding of proteins (BSA) with drugs as well as amphiphilic substances opened a new area of research called “protein–surfactant interaction”, subsequently providing a wide range of applications in the field of drug delivery and drug formulations, biotechnology, detergents, foods and the cosmetics industry.^{4–7} Strong binding lessens the abundance of free drug in plasma while weak binding indicates short lifetime or less distribution. Spectroscopic elucidation of strong binding between BSA and drug such as ceftriaxone has been studied in the absence and presence of Zn by Yue *et al.*⁸ The protein–drug complex directs a model for studying drug–protein interactions which would later facilitate the research of metabolism and transportation. Li *et al.*⁹ studied the effect of chitosan on the microenvironment and the secondary structure of BSA which provides theoretical guidance for the design of new ingredients for the food industry.

Assembly of amphiphilic molecules including surfactants/drugs *etc.* has a spontaneous propensity to form aggregates known as micelles due to the presence of weak non-covalent forces (*i.e.* van der Waals force) among the individual molecules. Micelles are formed at a particular concentration of that amphiphile known as “critical micelle concentration”.^{10–16} It is a unique feature of surfactant which helps to extract information about the interaction of drug and other biomolecules such as protein in the body. In the case of drug delivery, surfactants are important molecules to act as recipients/excipients. Hydrophobic nucleus/core of micelles is significant as it has the capability to enhance the solubility or bioavailability of hydrophobic drugs. Cationic surfactants are such surfactants that are utilized like anti-microbial substances in many types of antiseptics, disinfectants, perfumes as well as in cosmetics.¹¹ Previous studies revealed that the presence of various additional substances exert their effects on the physiological feature of micelles including the extent of departure of opposite-ion binding, catalytic property, and reaction rate which are related to the phase-separation model.^{11,17}

According to Deep and Ahluwalia, native globular protein's stability is noticeably influenced by pH, temperature, and insertion of small molecules like surfactants, co-enzymes, inhibitors and activators that bind specifically to the original state.¹⁸ The interaction between surfactant & globular proteins can help towards an understanding of the action of surfactant as a solubilizing agent for membrane proteins and lipids. Some excellent literatures are available on the study of the interaction between drugs/salts/polymers/polyols with surfactants; however, to the best of our knowledge, no study yet reported the interaction between BSA and CDMEAB.^{19–23} CDMEAB is a cationic surfactant which is used as disinfectant/laboratory purpose.²⁴ Understanding the interaction between BSA and CDMEAB could be a key interest in the meadow study of protein-membrane chemistry model.

Therefore, in our present work, we have employed conductivity technique in order to investigate the interaction between

BSA (Scheme 1(I)) and CDMEAB (Scheme 1(II)). Numerous studies revealed that when the aggregates are exposed to a variety of additives, the physiological properties (aggregate's stability, reaction rate, degree of ionization, and clouding phenomenon) could be affected due to the presence of additives.^{25–27} In a biological system, sodium ion plays a vital role to control the sodium pump and transmitting nerve systems.²⁸ Salt ions have the unique property to decrease the critical micelle concentration (c^*) by reducing the existing electrostatic repulsion force among the polar head groups of the surfactant, therefore, increasing the aggregation number (N_{agg}).²⁹ Therefore, NaCl and Na₂SO₄ were utilized in this work to understand the effect of additives on different physicochemical parameters corresponding to BSA–CDMEAB interaction.

In recent years, MD simulation techniques are being implemented to understand the effect of surfactant on protein structure and on the protein folding phenomenon.^{30–32} Hoque *et al.* have studied the interaction of a cationic surfactant, cetylpyridinium chloride on BSA in presence and absence of electrolytes using MD simulation technique.³³ MD simulation provided detailed information including non-covalent interactions between protein and surfactant, root-mean-square deviation of alpha carbon and backbone of the protein and root-mean-square fluctuation of each amino acid present in the protein. Delgado-Magnero and coworkers have utilized MD simulation method to understand the binding mechanism of nonionic surfactants with BSA and the results obtained suggest the potential use of polysorbates as excipients for minimizing the undesirable effects of protein adsorption and aggregation.³⁴ In the present study, we investigate the nature and type of interactions present among the surfactant molecules and BSA employing MD simulation to support the experimental observations.

2. Experimental procedure

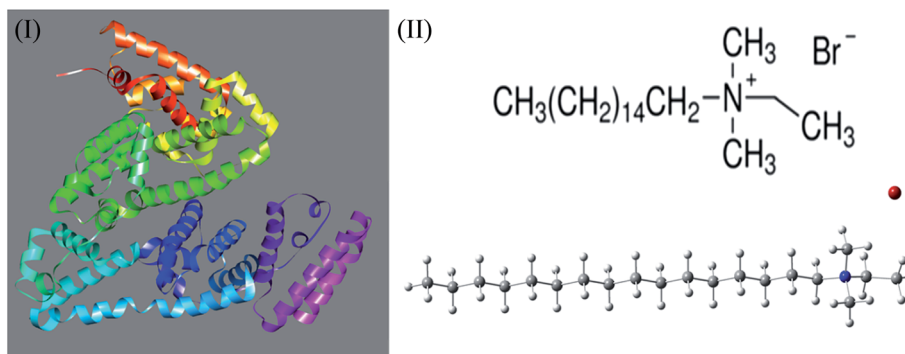
2.1. Materials and solutions

All the chemicals used in the study were of analytical grade and collected to use without further purification. CDMEAB (mass fraction purity, 0.99), BSA (mass fraction purity, >0.98), NaCl (mass fraction purity, 0.99), Na₂SO₄ (mass fraction purity, 0.99) were purchased from Across Organics (USA), Merck (Germany), Merck (India) and Merck (India) respectively.

2.2. Conductivity measurement technique

CDMEAB and CDMEAB + BSA solutions were made in molal concentration (with/without a particular concentration of salts solution) by using distilled-deionized water having specific conductivity 1.7–2.0 $\mu\text{S cm}^{-1}$ (over the temperature range 298.15–323.15 K). Weights of samples during solution preparation were taken by using an electric balance (Metler Toledo). In order to evaluate the magnitudes of specific conductivity of pure surfactant/(BSA + surfactant) system, we have used 4510 conductivity meter (manufacturer: Jenway, UK) including a glass cell electrode having cell constant 0.97 cm^{-1} . Standard KCl solution having appropriate concentration was used to





Scheme 1 Bovine serum albumin (BSA) (I) and molecular structure of CDMEAB (II).

perform the calibration of the conductivity meter. Alternative current (AC) having frequency 60 Hz was employed to study the conductivity. At first, 25 mmol kg⁻¹ solution of CDMEAB in water was prepared by utilizing an aqueous solution of BSA (at room temperature) and the surfactant solution was inserted into 20 mL BSA solution (with/without a fixed concentration of salts). These solutions were prepared by using salts (NaCl and Na₂SO₄) for examining the influence of salts in which the same concentration of salts was considered. Conductivity was measured after every addition of prepared solutions with thoroughly mixing and time was provided for the attainment of temperature equilibration; therefore, this process was repeated after every addition. The results obtained provide good matching with literature results.^{25–28,35} During the whole temperature range, RM6 Lauda circulating water bath was used to control the temperature and temperature error ± 2 K was considered. Three different (c^*) magnitudes were extracted from the abrupt alteration of specific conductivity with concentration of CDMEAB solution and (CDMEAB + BSA) mixtures.

2.3. Molecular dynamics simulations

The crystal structure of serum albumin 4F5S from *Bos taurus* (Bovine) was collected from protein data bank (shown in Scheme 1(I)). This structure was chosen based on the resolved amino acid residues (25–607) and relatively high resolution of 2.47 Å compared to the other available structures in UniProt (<http://www.uniprot.org/uniprot/P02769>). The molecular dynamics simulation of one BSA and 40 CDMEAB molecules with water was carried out for 20 ns in the absence and presence of salt (NaCl) using YASARA Dynamics program.^{36,37} The Amber14 force field was used for all the simulations.³⁸ A cut off radius of 8 Å was used for short-range van der Waals and Coulomb interactions. The long-range electrostatic interactions were calculated using the particle-mesh Ewald method.³⁹ Periodic boundary condition (cell dimension of 105 Å \times 105 Å \times 105 Å) and temperature 298 K were considered for all simulations. Time step 1.25 fs was used and simulation snapshots were saved at every 100 ps. The solvent density of 0.998 g ml⁻¹ for both systems was maintained by adding 34 699 molecules of water. Macro programmed protocol was used for steepest decent minimization and equilibration for salt and no salt

environment. Molecular dynamics trajectory, root mean square deviation (RMSD) of alpha carbon and backbone, solvent accessible surface area (SASA), radius of gyration as well as the residue-specific root mean square fluctuations (RMSF) of protein were calculated by specific macros written in Yanaconda language. Collision cross section measurements of the protein structures were performed using the IMPACT program.⁴⁰

3. Results and discussion

3.1. Determination of critical micelle concentration (c^*) and counter-ion binding parameter (β) of pure and mixed systems

The concentration above which surfactant micellization takes place is called critical micelle concentration (c^*). The c^* values can be evaluated from the abrupt change of physico-chemical properties with the change of concentration of the surfactant solutions. In the current study, the specific conductivity (κ) of CDMEAB in H₂O/H₂O + BSA solutions having different concentrations is measured and the κ values are found to vary with the concentration of surfactant. Fig. 1 depicts the variation of specific conductivity associated with different concentration of CDMEAB in aqueous solution for pure CDMEAB and CDMEAB + BSA mixture. Every intersection points of the plots demonstrate the certain concentration of amphiphiles which is regarded as critical micelle concentration (c^*). Here we have obtained three different noticeable break points for pure CDMEAB and (BSA + CDMEAB) mixtures which are designated as c_1^* , c_2^* , and c_3^* . The primary kink reflects the formation of BSA-surfactant complex (convey of the concentration of critical aggregation), second breakpoint indicates the CDMEAB micellization in presence of BSA whereas the tertiary breakpoint is acknowledged as the structural modification of micelles such as spherical shape to rod shape.⁴¹ More than one critical micelle concentration (c^*) for various amphiphiles has been observed by many researchers as well as in our previous study.^{42,43}

The obtained magnitudes of c_1^* , c_2^* , and c_3^* of CDMEAB are found 0.74, 1.49, and 6.57 mmol kg⁻¹ respectively at 303.15 K. On the other hand, the values of c_1^* , c_2^* , and c_3^* in aqueous BSA (0.005–0.100 mmol kg⁻¹) + CDMEAB mixed systems are found (0.75–1.05 mmol kg⁻¹), (2.22–2.49 mmol kg⁻¹), and (6.35–7.67 mmol kg⁻¹) respectively (Table 1). The values of CDMEAB are in good agreement with literature values.⁴⁴ In presence of



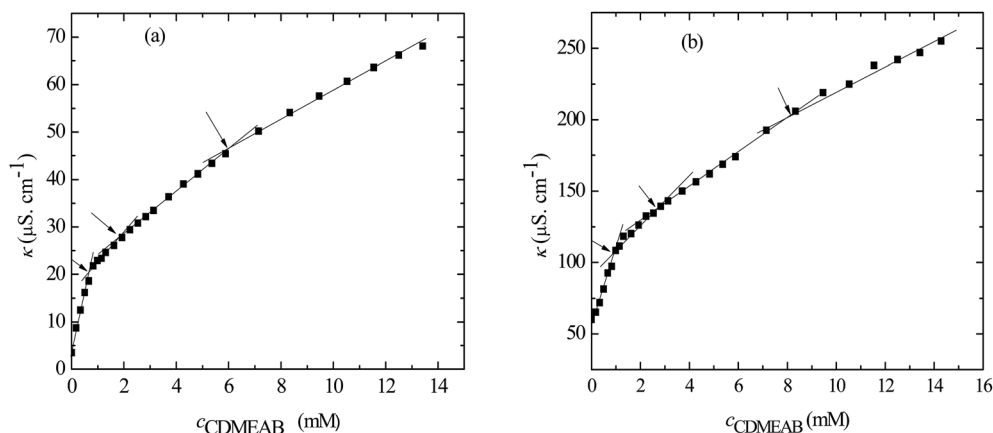


Fig. 1 Specific conductivity (κ) versus concentration of CDMEAB for (a) pure CDMEAB and (b) (BSA + CDMEAB) mixed system containing $0.03 \text{ mmol kg}^{-1}$ BSA in water at 303.15 K .

Table 1 Values of c^* for pure CDMEAB and (BSA + CDMEAB) systems in an aqueous solution containing different concentrations of BSA at 303.15 K^a

c_{BSA} (mmol kg^{-1})	c_1^* (mmol kg^{-1})	c_2^* (mmol kg^{-1})	c_3^* (mmol kg^{-1})
0.000	0.74	1.49	6.57
0.005	0.75	2.27	6.79
0.010	0.77	2.32	7.67
0.030	0.91	2.49	6.84
0.050	0.93	2.31	6.52
0.100	1.05	2.22	6.35

^a Relative standard uncertainties (u_r) is $u_r(c_1^*/c_2^*/c_3^*) = 0.03$.

BSA, c_1^* values are increased with increasing the concentration of BSA. The c_2^* and c_3^* values are first increased, reach a maximum and then decreased. These variations in c^* values signify the interaction between BSA and surfactant. The decrease in c_2^* and c_3^* values at higher concentration of BSA possibly due to the electrostatic interaction between BSA and the positively charged surfactant. However, interactions sustained through the nonpolar interactions of the hydrophobic portion of BSA and CDMEAB. The values of the degree of micelle ionization (α) were extracted by means of taking the corresponding ratios of the slopes relating to the closest straight lines before & after the c^* .^{21,22,45,46} Then α_1 , α_2 , and α_3 can be obtained from the ratios of S_2/S_1 , S_3/S_1 and S_4/S_1 respectively, where S_1 , S_2 , and S_3 are the first, second and third straight lines. The fraction of counter ion binding (β) can be obtained from the relation: $\beta = 1 - \alpha$.

At a particular concentration of BSA ($0.03 \text{ mmol kg}^{-1}$) in the c^* values for BSA + CDMEAB mixed systems (ESI Fig. 1†) are decreased with temperature up to a fixed limit and then increased as well as provided a U-contoured architecture. The ESI Fig. 1† demonstrates that micellization became favored with increasing temperature due to the dehydration of hydrophobic moiety which was predominated over the dehydration of hydrophilic moiety; therefore, decline the propensity of hydrophilic hydration, which is highly significant than the propensity

of hydrophobic dehydration at higher temperatures and lessen the micellization process.

3.2. Impact of salt solutions (NaCl and Na_2SO_4) on c^* and β of (BSA + CDMEAB) mixed system at various temperatures

The obtained values of c^* and β of pure CDMEAB & (BSA + CDMEAB) systems in the presence of two different sodium salts are viewed in Table 2. In cell membranes, the presence as well as the concentration of electrolytes is an important consideration as they may affect the interaction of surfactant. So, drug-surfactant interaction is an important approach where the coagulation process of surfactant molecules should be brought under the study of the impact of electrolytes, temperatures, pH, and the concentration of various additives. The values of c_1^* for (BSA + CDMEAB) in salt medium is noticeably decreased than pure CDMEAB and (BSA + CDMEAB) mixed system while the second c_2^* values show the opposite pattern. On the other hand, the third c_3^* values of (BSA + CDMEAB) in presence of salts are comparatively lower than pure (BSA + CDMEAB) missed system. Overall, the electric double layer is compressed due to the presence of an aqueous solution of salts, and subsequently boosted the lessening of repulsion affinity among the head groups of CDMEAB and BSA. At a particular concentration of BSA and salts, c_1^* and c_2^* are first decreased and then increased. However, the c_3^* values did not exhibit any regular trend. In our study, we have chosen such salts that play a vital role in human body. Our principal interest is to understand the way how they interact with each other as well as providing additional information about thermodynamic conditions. These findings provoke better information about surfactant-protein interaction, drug-surfactant interaction in the presence of salts. Salts give a convenient environment for micellization. Table 2 demonstrate that the presence of salts decreases the c^* values and enhances the aggregation process. The obtained values of c^* are comparatively lower in case of Na_2SO_4 than NaCl which reveals that aggregation process is highly suitable with NaCl for (BSA + CDMEAB) mixture.



Table 2 Values of c^* and β for (BSA + CDMEAB) mixed system (with/without salts) at different temperature^a

Medium	$c_{\text{BSA}}/\text{mmol kg}^{-1}$	$I_{\text{Salt}}/\text{mmol kg}^{-1}$	T/K	$c_1^*/\text{mmol kg}^{-1}$	$c_2^*/\text{mmol kg}^{-1}$	$c_3^*/\text{mmol kg}^{-1}$	β_1	β_2	β_3
CDMEAB									
H ₂ O	0.0	0.00	298.15	0.84	1.94	6.08	0.70	0.76	0.83
			303.15	0.74	1.49	5.83	0.75	0.80	0.87
			308.15	0.64	2.28	5.46	0.71	0.81	0.89
			313.15	0.77	2.36	6.06	0.72	0.83	0.90
			318.15	0.83	2.55	6.16	0.70	0.83	0.89
			323.15	0.97	2.67	6.26	0.70	0.82	0.89
BSA + CDMEAB									
H ₂ O	0.03	0.00	298.15	0.99	2.44	8.46	0.69	0.88	0.88
			303.15	0.91	2.49	7.97	0.66	0.75	0.84
			308.15	0.83	2.61	7.67	0.68	0.79	0.85
			313.15	1.03	2.71	7.58	0.64	0.76	0.85
			318.15	1.08	2.89	6.71	0.69	0.69	0.88
BSA + CDMEAB									
H ₂ O + NaCl	0.03	0.50	303.15	0.88	3.26	7.24	0.28	0.79	0.84
			303.15	0.78	3.08	7.31	0.69	0.76	0.82
			303.15	0.71	2.86	7.52	0.54	0.67	0.77
			303.15	0.85	2.67	7.41	0.58	0.69	0.84
			303.15	0.89	2.63	—	0.58	0.69	—
BSA + CDMEAB									
H ₂ O + Na ₂ SO ₄	0.03	0.50	303.15	0.79	1.96	5.87	0.57	0.72	0.81
			303.15	0.71	2.67	6.56	0.57	0.67	0.77
			303.15	0.65	3.09	6.68	0.60	0.69	0.77
			303.15	0.67	3.78	7.91	0.60	0.62	0.72
			303.15	0.79	4.82	—	0.57	0.56	—
BSA + CDMEAB									
H ₂ O + NaCl	0.03	1.50	298.15	0.87	3.24	8.31	0.65	0.63	0.67
			303.15	0.71	2.86	7.52	0.70	0.80	0.85
			308.15	0.75	2.24	7.04	0.68	0.77	0.83
			313.15	0.81	2.13	6.99	0.61	0.77	0.84
			318.15	0.85	2.43	7.22	0.65	0.78	0.85
			323.15	0.93	2.66	7.49	0.70	0.78	0.85
BSA + CDMEAB									
H ₂ O + Na ₂ SO ₄	0.03	1.50	298.15	0.89	2.79	7.59	0.58	0.61	0.67
			303.15	0.65	3.09	6.68	0.56	0.68	0.82
			308.15	0.69	3.09	6.87	0.56	0.68	0.82
			313.15	0.72	3.56	7.47	0.55	0.66	0.77
			318.15	0.76	3.76	7.61	0.53	0.65	0.76
			323.15	0.86	4.07	7.75	0.56	0.68	0.77

^a Relative standard uncertainties (u_r) are $u_r(c_1^*/c_2^*/c_3^*) = 0.03$ and $u_r(\beta_1/\beta_2/\beta_3) = 0.04$.

According to some previous study, chloride ion shows chaotropic nature along with reduction of the stability of hydrophobic moiety or associates of surfactants. Another anion is sulfate ion which carries high charge density and acts as water structure maker (a powerful kosmotrope). Therefore, raises the stability of hydrophobic aggregates of surfactants.^{47,48} Table 2 also indicates that Na₂SO₄ have better salting out effect than NaCl. The β magnitudes of CDMEAB & (BSA + CDMEAB) mixed systems in presence of electrolytes did not follow any hemolytic orientation while the aggregates stability is proved due to the high values of β (almost in all cases above 70% (± 2), except c_1^*).

3.3. Temperature effects on the micellization of CDMEAB and (BSA + CDMEAB) mixtures

van der Waals interaction among the hydrophobic portion of surfactants are responsible for the aggregation phenomena. Effect of the variation of temperature of experimental system is one of the major considerable factors which affect the micellization process. Due to the propensity of hydration of polar portions, micellization may be delayed. Table 2 shows the values of c^* at a particular temperature in the presence of electrolytes. In the case of pure CDMEAB, first and third c^* values are initially decreased, attain the minimum position and



then increased with rise of temperature whereas the second c^* values are gradually increased with temperature. For the (BSA + CDMEAB) mixed system, the second and third c^* values are gradually increased. After that, they decreased while first c^* values are decreased, powered a nearest flat, and then increased with an accretion of temperatures. At a certain temperature (303.15 K), in salt medium, first c^* values are increased initially and then decreased. On the other hand, the second & third c^* values are gradually increased in Na_2SO_4 solution and gradually decreased in NaCl solution. Besides, with varying temperature in NaCl/ Na_2SO_4 solution, c^* values are first decreased and then increased (except c_2^* values in Na_2SO_4 solution).

The effect of temperature on micellization of surfactant depends on the nature of surfactants whether it is ionic or non-ionic. Usually ionic amphiphiles exhibit U-shaped behavior,^{49,50} while non-ionic amphiphiles show regular trend;^{51–53} though it has been noted that non-ionic amphiphiles also provide the U-shaped curve.⁵⁴ In our case, ESI Fig. 1† provides good harmony with literature. The alteration of critical micelle concentration due to the variation of temperatures may be analyzed by means of mode of hydration of the single surfactant and protein-surfactant mixtures. In the saturated solution of surfactant and its mixtures, both hydrophobic and hydrophilic hydration is possible while the aggregates of surfactant faced by hydrophilic hydration. However, reduction of both types of hydration is possible with an elevation of temperature and the desolvation of polar head groups causes the favorable micellization. Many studies showed that elevation of temperature causes the disruption of H_2O entities and rupturing of H-bonds around the hydrocarbon moiety of CDMEAB/BSA, stimulates the lessening of hydrophobic de-hydration, therefore, to decrease the micellization rate.^{11,17,50–52} Again, the β values of CDMEAB and (BSA + CDMEAB) mixed systems did not follow any regular bias with regard to the temperature in the presence and absence of sodium salts (Table 2).

3.4. Investigation of micellization process insight from thermodynamic states

The theoretical & practical investigation of drug-surfactant interaction, protein-surfactant interaction *etc.* significantly relies on the thermodynamic parameters. Collection of information of thermodynamic specifications helps to understand the consequences of a mode of collaboration among the studied constituents. Herein, we have evaluated different thermodynamic parameters of pure CDMEAB and (BSA + CDMEAB) systems in the presence as well as absence of salts through the following equations.^{32,55–58}

$$\Delta G_m^0 = (1 + \beta)RT \ln(X_{c^*}) \quad (1)$$

$$\Delta H_m^0 = -(1 + \beta)RT^2(\partial \ln X_{c^*}/\partial T) \quad (2)$$

$$\Delta S_m^0 = (\Delta H_m^0 - \Delta G_m^0)/T \quad (3)$$

Eqn (1) and (2) contains X_{c^*} which denotes mole fraction of c^* . ESI Fig. 1† depicts non-linear curve of $\ln(X_{c^*})$ versus T plot and these type of plots are used to calculate the magnitudes of

ΔH_m^0 . The values of tangents were measured as per temperature and then slopes are taken from the obtained tangents as the tantamount of $\ln(X_{c^*})$.^{59,60} The obtained magnitudes of different thermodynamic specifications for pure CDMEAB and (BSA + CDMEAB) mixed systems with and without salts (NaCl & Na_2SO_4) are summarized in Table 3.

The obtained magnitudes of ΔG_m^0 (free energy changes) are conceived to be negative which provides strong evidence that the aggregation process is spontaneous in nature. The negative values of free energy changes are elevated with an elevation of temperatures. Such negative values of ΔG_m^0 for the micellization of surfactants were also reported.^{19–22,55–58} These values are comparatively lower in presence of salts suggesting the slight reduction of dynamic forces for perfect interaction.

In the case of pure CDMEAB and (BSA + CDMEAB), the $\Delta H_{1,m}^0$, and $\Delta H_{3,m}^0$ values are negative at the initial temperatures which suggest the process is exothermic in nature. These negative values are decreased with rising temperatures and revealed the endothermic process by altering the negative values to positive sign. The $\Delta H_{2,m}^0$ values are positive (*i.e.* endothermic process) which are raised by means of rising temperatures. Again, all the values of entropy changes ($\Delta S_{1,m}^0$, $\Delta S_{2,m}^0$, and $\Delta S_{3,m}^0$) of pure CDMEAB and (BSA + CDMEAB) are positive and these values are raised by means of rising temperatures. Changes of entropy values point out that the aggregation process is entirely entropy controlled, especially at elevated temperatures. However, at the initial temperatures, the systems are both entropy and enthalpy controlled. These values of enthalpy and entropy also indicate the existence of hydrophobic (at the higher temperatures) and electrostatic (at the lower temperatures) interactions.

In presence of salts, the values of enthalpy ($\Delta H_{1,m}^0$, $\Delta H_{2,m}^0$, and $\Delta H_{3,m}^0$) for pure CDMEAB and (BSA + CDMEAB) are negative at the initial temperatures (except $\Delta H_{2,m}^0$ values in presence of Na_2SO_4), which suggests the process is exothermic in nature. These negative values are decreased with rising temperatures and revealed the endothermic process by altering the negative values to positive sign. On the other hand, the values of entropy changes are positive except $\Delta S_{1,m}^0$ and $\Delta S_{2,m}^0$ (in presence of NaCl) & $\Delta S_{1,m}^0$ (in presence of Na_2SO_4) values at 298.15 K and 303.15 K temperatures.

The negative values of ΔH_m^0 demonstrate the existence of electrostatic interaction between the head groups, while the positive ΔH_m^0 magnitudes demonstrate the disruption of water structure as well as the presence of hydrophobic interaction. Such negative and positive values of ΔH_m^0 also reported by some groups previously.^{61–64}

The positive magnitude of entropy changes (ΔS_m^0) of pure CDMEAB and (BSA + CDMEAB) mixed system can be analyzed by means of taking two considerable factors. Firstly, the disruption of iceberg structures as well as shifting of the hydrophobic moiety from H_2O environment to hydrophobic environment of micelle cores. Secondly, the enhancement of free rotation of hydrophobic moiety inside the micelle core compared to H_2O environment.⁶⁵ On the other hand, a negative magnitude of entropy change (ΔS_m^0) may be possible only when





Table 3 Values of thermodynamic parameters for (BSA + CDMEAB) mixed system in water and in an aqueous solution of NaCl and Na₂SO₄^b

System	Medium	I_{salt}^a mmol kg ⁻¹	T/K	$\Delta G_{1,m}^0$ kJ mol ⁻¹	$\Delta G_{2,m}^0$ kJ mol ⁻¹	$\Delta G_{3,m}^0$ kJ mol ⁻¹	$\Delta H_{1,m}^0$ kJ mol ⁻¹	$\Delta H_{2,m}^0$ kJ mol ⁻¹	$\Delta H_{3,m}^0$ kJ mol ⁻¹	$\Delta S_{1,m}^0$ J mol ⁻¹ K ⁻¹	$\Delta S_{2,m}^0$ J mol ⁻¹ K ⁻¹	$\Delta S_{3,m}^0$ J mol ⁻¹ K ⁻¹	$\Delta C_{1,m}^0$ kJ mol ⁻¹ K ⁻¹	$\Delta C_{2,m}^0$ kJ mol ⁻¹ K ⁻¹	$\Delta C_{3,m}^0$ kJ mol ⁻¹ K ⁻¹
CDMEAB	H ₂ O	0.00	298.15	-46.77	-44.75	-41.46	-41.67	20.50	-11.56	17.11	218.83	100.28	3.75	0.52	1.12
			303.15	-49.60	-47.66	-43.13	-23.21	23.11	-6.09	87.05	233.44	122.18	4.07	0.53	1.21
			308.15	-49.80	-46.76	-44.75	-0.89	25.65	0.34	158.72	234.98	146.33	4.39	0.53	1.30
			313.15	-50.02	-47.95	-45.04	21.48	28.50	7.08	228.33	244.15	166.45	4.72	0.53	1.38
			318.15	-50.31	-48.14	-45.54	45.74	31.03	14.36	301.91	248.84	188.29	5.04	0.53	1.47
BSA + CDMEAB	H ₂ O	0.00	298.15	-51.22	-48.76	-46.07	71.66	33.73	21.60	380.25	255.27	209.41	5.34	0.54	1.55
			303.15	-45.85	-46.86	-40.91	-16.65	7.46	-20.15	97.94	182.19	69.64	2.15	0.36	0.74
			308.15	-46.12	-44.20	-40.93	-5.73	9.09	-16.22	133.20	175.77	81.52	2.20	0.40	0.78
			313.15	-47.70	-45.74	-42.06	4.81	11.71	-12.19	170.40	186.42	96.93	2.26	0.45	0.81
			318.15	-46.48	-45.55	-42.97	16.32	14.01	-8.27	200.40	190.20	110.81	2.32	0.49	0.85
BSA + CDMEAB	H ₂ O–NaCl	1.50	298.15	-48.36	-43.97	-44.81	29.69	15.83	-3.84	245.30	187.95	128.78	2.38	0.53	0.89
			303.15	-45.92	-46.95	-45.41	40.01	19.35	0.78	265.90	205.16	142.92	2.44	0.58	0.92
			308.15	-43.71	-39.32	-36.52	-78.09	-65.10	-27.83	-115.29	-86.48	29.17	5.16	3.59	1.52
			313.15	-48.40	-44.68	-41.48	-52.95	-49.20	-21.50	-14.99	-14.91	65.91	5.59	4.13	1.76
			318.15	-48.25	-45.84	-42.06	-22.62	-25.84	-10.86	83.17	64.91	101.25	6.05	4.69	2.00
BSA + CDMEAB	H ₂ O–Na ₂ SO ₄	1.50	298.15	-46.82	-46.95	-42.93	9.56	1.41	0.28	180.05	154.43	137.98	6.52	5.24	2.26
			303.15	-48.31	-47.30	-43.82	42.28	28.05	11.93	284.72	236.83	175.23	6.98	5.80	2.50
			313.15	-50.17	-47.69	-44.33	78.35	57.03	24.38	397.70	324.06	212.62	7.41	6.35	2.74
			318.15	-41.42	-39.62	-36.76	-92.92	15.58	-13.08	-172.71	185.16	79.42	5.81	0.60	1.14
			303.15	-43.80	-41.38	-39.94	-64.48	18.77	-7.69	-68.21	198.40	106.39	6.25	0.63	1.26
	H ₂ O		308.15	-45.38	-42.13	-41.98	-31.23	22.21	-0.80	45.93	208.81	133.64	6.68	0.66	1.37
			313.15	-45.53	-41.72	-41.03	4.40	25.22	6.66	159.45	213.74	152.30	7.14	0.69	1.52
			318.15	-45.46	-41.84	-41.52	39.90	28.52	14.42	268.32	221.16	175.81	7.58	0.72	1.62
	323.15	-46.57	-43.01	-42.30	79.05	32.66	22.48	388.75	234.17	200.47	8.00	0.75	1.73		

^a I = ionic strength of salts used. ^b Relative standard uncertainties (u_r) limits are $u_r(\Delta G_m^0)$, $u_r(\Delta H_m^0)$, $u_r(\Delta S_m^0)$, and $(\Delta_m C_p^0)$ are 0.03, 0.03, 0.03, 0.04 and 0.04 respectively.

the molecules of BSA and CDMEAB are affected due to the construction of iceberg structure around them.

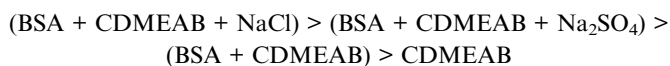
ESI Fig. 2† shows the results of free energy changes (ΔG_m^0) which is form the enthalpic contribution ($H_{1,m}^0$) together with entropic augmentation ($-T\Delta S_{1,m}^0$). ESI Fig. 2† also depicts that enthalpy augmentation is gradually enhanced with rising temperatures while entropy augmentation showed opposite style in case of (BSA + CDMEAB) system in water. On the other hand, the supplementation of enthalpy along with entropy showed an interesting pattern in the salt medium. With an elevation of temperature, enthalpy contribution is increased and entropy contribution is decreased & showed a cross point. It is also clear from ESI Fig. 2† that both contributions are comparatively higher in salts medium than aqueous medium and provided the following pattern:



Molar heat capacity is an important thermodynamic aspect through which we may noticeably understand the way by which drug-surfactant or protein-surfactant interaction get motion restriction along with structural rearrangement/response to binding of ligands. The value of changes of molar heat capacity (ΔC_m^0) for the micellization of pure CDMEAB and (BSA + CDMEAB) mixed system are summarized in Table 3. These values were achieved by taking the corresponding slopes from the plots of enthalpy changes (ΔH_m^0) vs. temperature (T) by utilizing the below-mentioned relation (4).^{66,67}

$$\Delta C_m^0 = ((\partial H_m^0)/\partial T)_p \quad (4)$$

The achieved magnitudes of ΔC_m^0 are positive in all cases and these values are clearly higher in the presence of salts than aqueous medium & followed the following trend (Table 3);



The alteration of molar heat capacity in case of BSA & CDMEAB mixtures can be considered to associate with motion restriction event. It is also nearly related to the molecular surface area as well as staying in the similar horizon of change in the solvent accessible surface area.⁶⁸ The little positive molar heat capacity values, as well as entropy values, indicate the less structural changes in studied surfactant's micelle over the period of binding interaction.⁴³

3.5. Transfer attributes of thermodynamics for (BSA + CDMEAB) mixed systems

The following eqn (5)–(7) were utilized in order to manipulate the transfer attributes of thermodynamic (such as free energy of transfer = $\Delta G_{m,\text{tr}}^0$, enthalpy of transfer = $\Delta H_{m,\text{tr}}^0$, & molar heat capacity of transfer = $\Delta C_{p,m,\text{tr}}^0$) for (BSA + CDMEAB) mixed systems.^{26,69,70}

$$\Delta G_{m,\text{tr}}^0 = \Delta C_m^0(\text{aq. additive}) - \Delta G_m^0(\text{aq.}) \quad (5)$$

$$\Delta H_{m,\text{tr}}^0 = \Delta H_m^0(\text{aq. additive}) - \Delta H_m^0(\text{aq.}) \quad (6)$$

and

$$\Delta C_{p,m,\text{tr}}^0 = \Delta C_{p,m}^0(\text{aq. additive}) - \Delta C_{p,m}^0(\text{aq.}) \quad (7)$$

The evaluated values of free energy of transfer ($\Delta G_{m,\text{tr}}^0$), enthalpy of transfer ($\Delta H_{m,\text{tr}}^0$), & molar heat capacity of transfer ($\Delta C_{p,m,\text{tr}}^0$) for (BSA + CDMEAB) mixed systems in pure water and in salts solution are shown in ESI Table 1†. The calculated values of free energy of transfer ($\Delta G_{m,\text{tr}}^0$) are positive in the cases of (BSA + CDMEAB) mixed system in water and salts (except $\Delta G_{2,m,\text{tr}}^0$ at 298.15 K in water). The first transfer enthalpy values for (BSA + CDMEAB) system in water are positives at (298.15–308.15) K, while other values including second & third are negative. On the other hand, in the presence of electrolytes, the obtained values of transfer enthalpy are negative and the negative values are gradually lessened with rising temperatures and at the high temperature, it changed to positive magnitude. Including our previous study, some researchers also reported the negative as well as positive values of transfer enthalpy.^{3,26,71} Jha *et al.*⁷² reported that negative values of transfer enthalpy are possible about the driving of organic (amino acids) and inorganic (various salts) molecules to urea solution from the water. The transfer of amphiphilic molecules from water to (BSA + CDMEAB) and (BSA + CDMEAB + salts) medium creates exothermic as well as endothermic environment respectively. The molar heat capacity of transfer ($\Delta C_{p,m,\text{tr}}^0$) is negative (except at 318.15 & 323.15) of (BSA + CDMEAB) mixed system in water while the others values of (BSA + CDMEAB) mixed system in salts are positive. These positive values of $\Delta C_{p,m,\text{tr}}^0$ signify that the micelles faced high hydration tendency which is determinable to achieve high H-bonding. On the contrary, negative magnitudes of $\Delta C_{p,m,\text{tr}}^0$ signify that the hydration of the polar head group is more significant than structured water molecules about a non-polar group of monomeric surfactants.⁷³ Apparently, the $\Delta C_{2,m,\text{tr}}^0$ value at 318.15 K is 0.00 $\text{kJ K}^{-1} \text{mol}^{-1}$ which is not actually 0.00 $\text{kJ K}^{-1} \text{mol}^{-1}$ except very low value. All the molar heat capacity of transfer ($\Delta C_{p,m,\text{tr}}^0$) are approximately constant which relates the noticeable structural changeover.

3.6. Investigation of enthalpy-entropy compensation for (BSA + CDMEAB) mixed systems

The following eqn (8) was utilized to evaluate the enthalpy-entropy compensation phenomena among the studied components. A linear relationship was also studied between ΔH_m^0 and ΔS_m^0 (ESI Fig. 3†). The straight lines between ΔH_m^0 and ΔS_m^0 provided excellent R^2 values with a range of 0.950–0.999 which are more or less approximate to unity.^{26,74,75}

$$\Delta H_m^0 = \Delta H_m^{0,*} + T_c \Delta S_m^0 \quad (8)$$

In the eqn (8), $\Delta H_m^{0,*}$ and T_c signs are denoted as intrinsic enthalpy gain and compensation temperature respectively. The values of $\Delta H_m^{0,*}$ and T_c for (BSA + CDMEAB) mixtures in water



Table 4 Enthalpy–entropy compensation parameters for (BSA + CDMEAB) systems containing 0.03 mM BSA in water and in aqueous salts solution^b

Medium	$I^a/\text{mmol kg}^{-1}$	$\Delta H_{1,m}^{0,*}/\text{kJ mol}^{-1}$	$\Delta H_{2,m}^{0,*}/\text{kJ mol}^{-1}$	$\Delta H_{3,m}^{0,*}/\text{kJ mol}^{-1}$	$T_{c,1}/\text{K}$	$T_{c,2}/\text{K}$	$T_{c,3}/\text{K}$
H ₂ O	0.00	−50.03	−61.88	−39.27	331.13	397.92	278.67
H ₂ O + NaCl	1.50	−45.91	−42.96	−38.92	309.03	300.65	290.67
H ₂ O + Na ₂ SO ₄	1.50	−42.74	−52.63	−38.85	307.64	363.6	300.82

^a I is the ionic strength. ^b Relative standard uncertainties (u_r) limit is $u_r(\Delta H_m^{0,*}) = 0.04$.

and salts solution are summarized in Table 4. The intrinsic enthalpy ($\Delta H_m^{0,*}$) values are found negative, suggesting the convenient and stable micelle formation even at $\Delta S_m^0 = 0^{20}$, therefore stable micelle formation in water compared to the salt solution is also noticed by means of reduction of $\Delta H_m^{0,*}$ values in salts medium.

The $\Delta H_m^{0,*}$ values demonstrate the solute–solute interaction along with an increase in hydrophobicity during micelle growth. The compensation temperature (T_c) values lie in the range of 278–363 K in the current study which shows good agreement with the literature values.^{10,20,48} Lumry *et al.*⁷⁶ reported compensation temperature (T_c) from 270 to 300 K and suggested that it is the indication of H₂O contributions in the protein solution as well as comparable to a biological fluid. Their finding supports our result with a good harmony. Ionic surfactants in water solution also showed same phenomena about compensation temperature.⁷⁷

3.7. Insights from molecular dynamics

Molecular dynamics (MD) simulation can provide microscopic details about the interactions present between a protein and drug or surfactant molecules. Simulation results can provide information about the stability of a system and the structural changes that a protein may undergo in presence of other small molecules. To gain atomistic level understanding of the aggregation behavior of CDMEAB surfactant molecules under the influence of BSA, we

have performed 20 ns all-atom simulation in the presence and absence of NaCl (Fig. 2). Salt ions are found to have significant contribution in stabilizing the protein–surfactant system as the positively charged sodium ions bind to the negatively charged amino acid residues of BSA. For instance, one sodium ion interacts with four amino acid residues Asp 13, Asp 254, Asp 255 and Asp 258 through three electrostatic interactions with the COO[−] moiety of the corresponding amino acids and a metal–acceptor bond with the C=O group of the peptide linkage between Asp 254 and Asp 255 (ESI Fig. 4†). BSA contains 99 negatively charged residues including aspartic acid and glutamic acid. An interesting fact is that BSA being a plasma protein has significant ability to bind metal ions including Na⁺. Fig. 3(A) shows the change in overall Gibbs free energy of the system with simulation timescale and it is observed that the presence of 5% NaCl causes a reduction in the overall Gibbs free energy of the system. Therefore, it may be inferred that the presence of salt ion causes the protein–surfactant system to stabilize to a greater extent. The 20 ns simulation snapshot in ESI Fig. 5† depicts that in presence of NaCl, the surfactant molecules form larger aggregates and come in close contact with BSA compared to when salt ions are not present. BSA has three distinct albumin domains: 19–209; 210–402; 403–600. When salt ions are present, the surfactant molecules are found to be equally distributed in these three domains but in absence of salt, no surfactant molecules interact with the 210–402 domain. The surfactant molecules are also found to be strongly interacting with the unstructured regions of BSA.

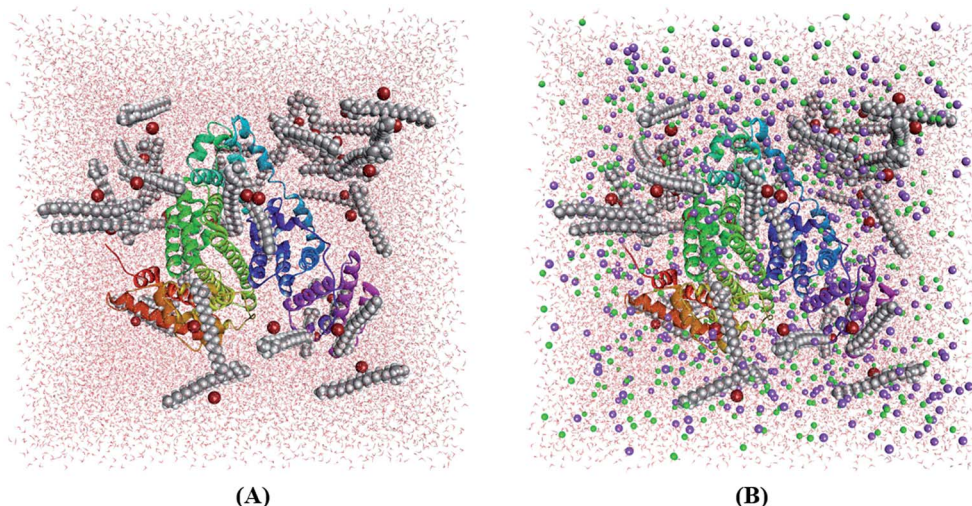


Fig. 2 Simulation box containing BSA + 40 CDMEAB in (A) no-salt and (B) 5% NaCl environments (sodium in purple and chloride in green).



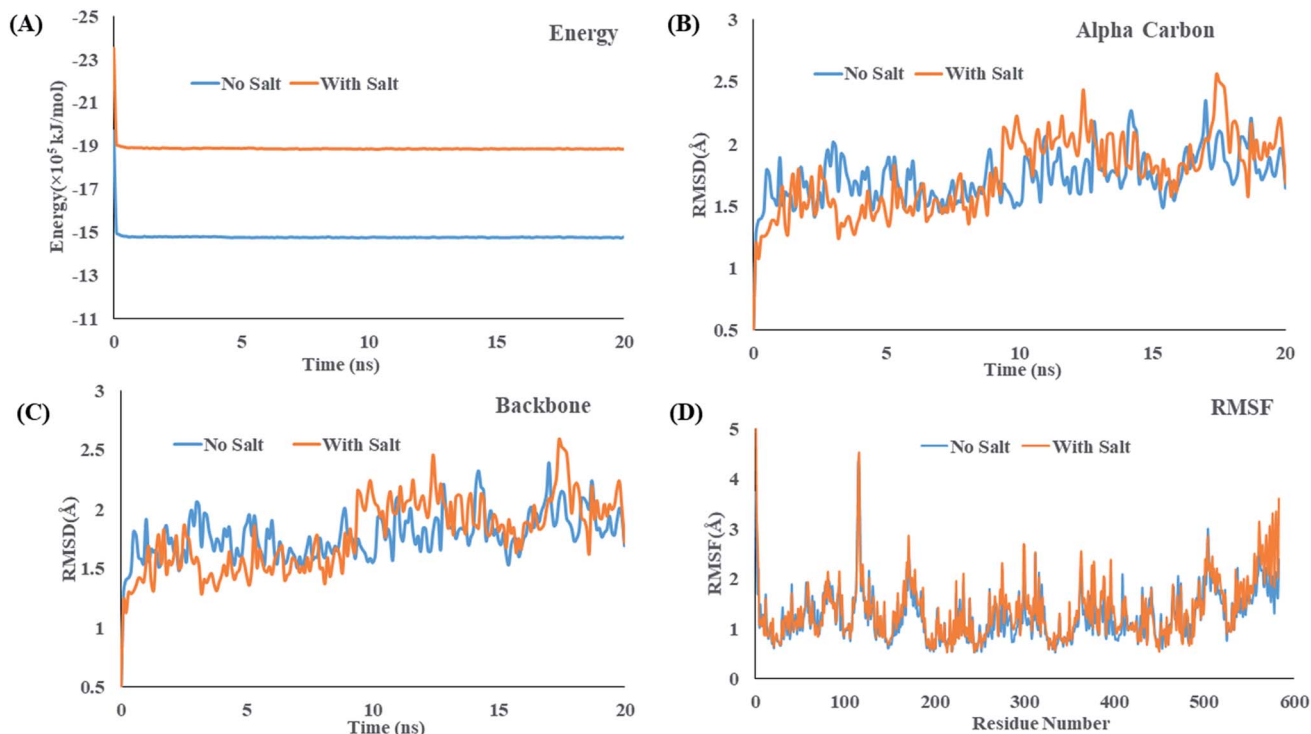


Fig. 3 (A) Gibbs free energy change over 5 ns molecular dynamics, comparative RMSD trajectory of (B) alpha carbon, (C) backbone of BSA, (D) RMSF trajectory of BSA.

Fig. 3(B and C) shows that the root means square deviation (RMSD) values of the alpha carbons and backbones of BSA remains almost similar throughout the simulation timescale and fluctuates around an average value. Presence of salt causes no significant change in the RMSD values which infers that the salt ions cause no significant structural changes of the protein. On the other hand, Fig. 4(A) depicts that the Solvent Accessible Surface Area (SASA) of the protein is much smaller in presence of NaCl. An explanation for the phenomenon can be inferred for the 20 ns simulation snapshot shown in ESI Fig. 5†. It reveals that in presence of salt, the larger aggregates of surfactant

molecules stay in close contact with the protein surface that makes the surface less accessible to solvent molecules.

Fig. 3(D) shows that the root means square fluctuation (RMSF) of the amino acid residues (1–560) remains almost similar in the both aqueous and salt environment but the RMSF values of the residues 561–583 are significantly larger when salt ions are present. As proteins may undergo significant structural changes while interacting with other molecules, in recent years, the calculation of collision cross section (CCS) has received great attention for providing precise information about the conformation and size of a protein in different environments.

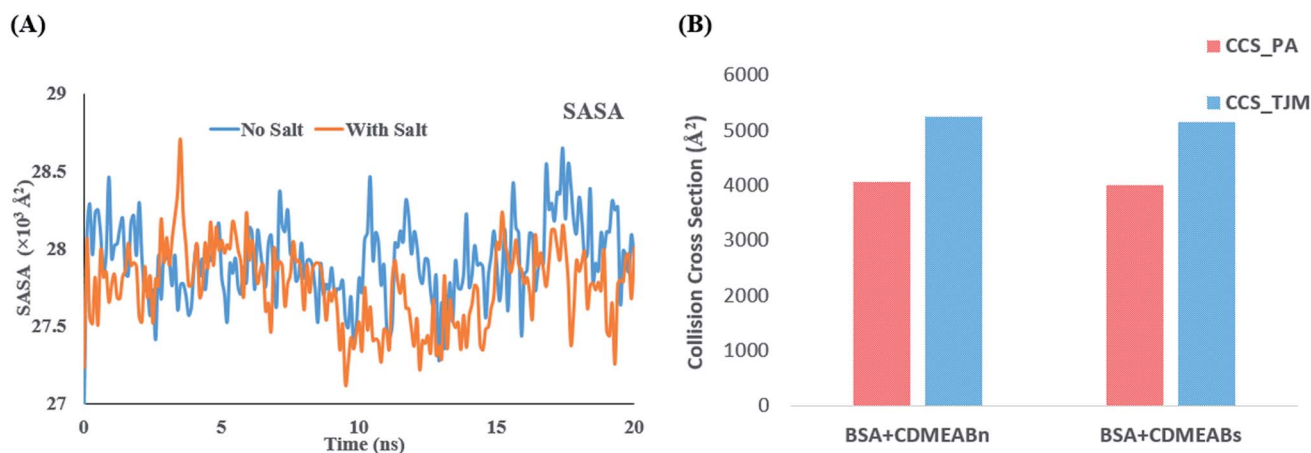


Fig. 4 (A) Change in SASA over the simulation timescale (B) collision cross sections of BSA in salt (CDMEABs) and no salt (CDMEABn) environment.



Both the projection approximation (PA) and trajectory method (TJM) has been utilized to measure perform the CCS analysis of BSA in the presence and absence of salt.^{78,79} The CCS_PA and CCS_TJM values of BSA in presence of salt are 4000.59 Å² and 5157.86 Å² respectively which are slightly lower than when no salt ions are present (Fig. 4(B)). In spite of strong interaction between the protein and surfactant molecules, the overall collision cross section of BSA + CDMEAB system remains almost similar to the CCS value of native BSA.

4. Conclusion

Micellization and thermodynamic properties of cationic surfactant CDMEAB with the globular protein bovine serum albumin (BSA) have been investigated in aqueous solution together with the presence of two different salts at various temperatures through conductometric method. The values of critical micelle concentration (c^*) enhanced due to the presence of BSA while the presence of salts comparatively declined the c^* values. Different parameters of thermodynamics (ΔH_m^0 and ΔS_m^0) demonstrated the existence of hydrophobic, exothermic and electrostatic interactions which are acting as binding forces among the protein and surfactant. Negative free energy changes (ΔG_m^0) are a strong indication of spontaneous micellization. The compensation temperature (T_c) of the studied systems suggest that the behavior of BSA + surfactant system along with additives are much comparable to a biological fluid. The negative $\Delta H_m^{0,*}$ values demonstrate the excellent stability of BSA + surfactant aggregate. Molecular dynamics (MD) simulation results unveiled some interesting features of the molecular interaction between BSA and CDMEAB in the presence and absence of salt environment. Firstly, strong and homogenous interaction was predicated between BSA and CDMEAB in the presence of NaCl. Secondly, RMSD and RMSF values of alpha carbon and amino acid residues and collision cross section (CCS) of the protein remained same in aqueous and salt environments that showed that there is no significant structural change occurred to the protein structure.

Conflicts of interest

There are no conflicts of interest to declare.

Acknowledgements

Dr Mohammad A. Halim likes to acknowledge The World Academy of Science (TWAS) for providing funding (17-479 RG/CHE/AS_I) to purchase high performance computer for performing molecular dynamics simulation.

References

- 1 K. A. Majorek, P. J. Porebski, A. Dayal, M. D. Zimmerman, K. Jablonska, A. J. Stewart, M. Chruszcz and W. Minor, *Mol. Immunol.*, 2012, **52**, 174.
- 2 T. Kosa, T. Maruyama and M. Otagiri, *Pharmaceutical Research*, 1997, **14**, 1607.
- 3 Y. Matsui and A. Okimoto, *Bull. Chem. Soc. Jpn.*, 1978, **51**, 3030.
- 4 D. Bhatt, K. C. Maheria and J. Parikh, *J. Surfactants Deterg.*, 2013, **16**, 547.
- 5 K. S. Sharma, S. R. Patil and A. K. Rakshit, *Colloids Surf., A*, 2003, **219**, 67.
- 6 D. Kumar and M. A. Rub, *J. Mol. Liq.*, 2018, **269**, 1.
- 7 T. Randolph and L. Jones, *Pharm. Biotechnol.*, 2002, **13**, 159.
- 8 Q. Yue, T. Shen, C. Wang, C. Gao and J. Liu, *Int. J. Spectrosc.*, 2012, **2012**, 1678.
- 9 G. Li, J. Huang, T. Chen, X. Wang, H. Zhang and Q. Chen, *Carbohydr. Polym.*, 2017, **176**, 75.
- 10 S. M. A. Ahsan, M. R. Molla, M. S. Rahman, M. A. Hoque, M. A. Rub and M. A. Khan, *J. Mol. Liq.*, 2019, **274**, 484.
- 11 D. Attwood and A. Florence, *Surfactant Systems—Their Chemistry, Pharmacy and Biology*, Chapman and Hall, New York, 1983.
- 12 N. Azum, A. Z. Naqvi, M. A. Rub and A. M. Asiri, *J. Mol. Liq.*, 2017, **240**, 189.
- 13 U. Ashraf, O. A. Chat, M. Maswal, S. Jabeen and A. A. Dar, *RSC Adv.*, 2015, **5**, 83608–83618.
- 14 N. Azum, M. A. Rub and A. M. Asiri, *J. Dispersion Sci. Technol.*, 2017, **38**, 96.
- 15 D. Kumar, K. E. Neo and M. A. Rub, *J. Surfactants Deterg.*, 2016, **19**, 101.
- 16 Y. K. Shrestha and F. Yan, *RSC Adv.*, 2014, **4**, 37274.
- 17 M. J. Rosen, *Surfactants and interfacial phenomena*, Wiley, New York, 1959.
- 18 S. Deep and J. C. Ahluwalia, *Phys. Chem. Chem. Phys.*, 2001, **3**, 4583.
- 19 M. R. Amin, S. Mahbub, S. Hidayathulla, M. M. Alam, M. A. Hoque and M. A. Rub, *J. Mol. Liq.*, 2018, **269**, 417.
- 20 S. Aktar, R. M. Molla, S. Mahbub, A. M. Rub, M. A. Hoque and D. S. Islam, *J. Dispersion Sci. Technol.*, 2018, DOI: 10.1080/01932691.2018.1472005.
- 21 M. A. Hoque, M. M. Alam, M. R. Molla, S. Rana, M. A. Rub, M. A. Halim, M. A. Khan and A. Ahmed, *J. Mol. Liq.*, 2017, **244**, 512.
- 22 M. A. Hoque, M. M. Alam, M. R. Molla, S. Rana, M. A. Rub, M. A. Halim, M. A. Khan and F. Akhtar, *Chin. J. Chem. Eng.*, 2018, **26**, 159.
- 23 H. Hamishehkar, S. Hosseini, A. H. Naseri, A. Safarnejad and F. Rasoulzadeh, *BioImpacts*, 2016, **6**, 125.
- 24 D. Otzen, *Biochim. Biophys. Acta, Proteins Proteomics*, 2011, **1814**, 562.
- 25 D. Kumar and M. A. Rub, *J. Mol. Liq.*, 2018, **250**, 329.
- 26 A. Z. Naqvi, M. A. Rub and Kabir-ud-Din, *Colloid J.*, 2015, **77**, 525.
- 27 M. A. Rub, N. Azum and A. M. Asiri, *Russ. J. Phys. Chem. B*, 2016, **10**, 1007.
- 28 I. Restrepo-Angulo, A. D. Vizcaya-Ruiz and J. Camacho, *J. Appl. Toxicol.*, 2010, **30**, 497.
- 29 W. Caetano and M. Tabak, *J. Colloid Interface Sci.*, 2000, **225**, 69.
- 30 M. Heshmat, V. P. Nicu and E. J. Baerends, *J. Phys. Chem. A*, 2012, **116**, 3454.
- 31 D. Lu, Z. Liu and J. Wu, *J. Chem. Phys.*, 2007, **126**, 064906.



- 32 S. Jana, S. Ghosh, S. Dalapati and N. Guchhait, *Photochem. Photobiol. Sci.*, 2012, **11**, 323.
- 33 M. A. Hoque, F. Ahmed, M. A. Halim, M. R. Molla, S. Rana, M. A. Rahman and M. A. Rub, *J. Mol. Liq.*, 2018, **260**, 121.
- 34 K. H. Delgado-Magnero, P. A. Valiente, M. Ruiz-Peña, A. Pérez-Gramatges and T. Pons, *Colloids Surf., B*, 2014, **116**, 720–726.
- 35 R. Amin, M. R. Molla, S. Rana, M. A. Hoque, M. A. Rub, M. Kabir and A. M. Asiri, *Phys. Chem. Liq.*, 2018, 1–17, DOI: 10.1080/00319104.2018.1519565.
- 36 E. Krieger, T. Darden, S. B. Nabuurs, A. Finkelstein and G. Vriend, *Proteins: Struct., Funct., Bioinf.*, 2004, **57**, 678.
- 37 F. Krieger, B. Fierz, O. Bieri, M. Drewello and T. Kiefhaber, *J. Mol. Biol.*, 2003, **332**, 265.
- 38 J. A. Maier, C. Martinez, K. Kasavajhala, L. Wickstrom, K. E. Hauser and C. Simmerling, *J. Chem. Theory Comput.*, 2015, **11**, 3696.
- 39 T. Darden, D. York and L. Pedersen, *J. Chem. Phys.*, 1993, **98**, 10089.
- 40 E. G. Marklund, M. T. Degiacomi, C. V. Robinson, A. J. Baldwin and J. L. Benesch, *Structure*, 2015, **23**, 791.
- 41 I. García-Mateos, S. Pérez and M. M. Velázquez, *J. Colloid Interface Sci.*, 1997, **194**, 356.
- 42 M. A. Rub, A. M. Asiri, J. M. Khan, F. Khan and R. H. Khan, *Taiwan Inst. Chem. Eng.*, 2014, **45**, 2068.
- 43 M. A. Hoque, M.-O.-F. Patoary, M. R. Molla, M. A. Halim, M. A. Khan and M. A. Rub, *J. Dispersion Sci. Technol.*, 2017, **38**, 1578.
- 44 Sk. M. A. Ahsan, M. D. Hossain, M. A. Hoque and M. A. Khan, *Indian J. Chem., Sect. A: Inorg., Bio-inorg., Phys., Theor. Anal. Chem.*, 2016, **55**, 160.
- 45 M. Rub, A. Asiri, D. Kumar, N. Azum and F. Khan, *Acta Phys.-Chim. Sin.*, 2014, **30**, 699.
- 46 F. Khan, M. S. Sheikh, M. A. Rub, N. Azum and A. M. Asiri, *J. Mol. Liq.*, 2016, **222**, 1020.
- 47 M. A. Hoque, M. D. Hossain and M. A. Khan, *J. Chem. Thermodyn.*, 2013, **63**, 135.
- 48 M. Rahman, M. A. Khan, M. A. Rub and M. A. Hoque, *J. Mol. Liq.*, 2016, **223**, 716.
- 49 M. Schick, *J. Phys. Chem.*, 1963, **67**, 1796.
- 50 G. Nemethy and A. Ray, *J. Phys. Chem.*, 1971, **75**, 809.
- 51 B. Sharma and A. K. Rakshit, *J. Colloid Interface Sci.*, 1989, **129**, 139.
- 52 C. La Mesa, *Phys. Chem. Liq.*, 1990, **94**, 323.
- 53 E. Crook, G. Trebbi and D. Fordyce, *J. Phys. Chem.*, 1964, **68**, 3592.
- 54 S. Chauhan, S. Kumari and K. Singh, *J. Chem. Thermodyn.*, 2017, **105**, 337.
- 55 J. Mata, D. Varade and P. Bahadur, *Thermochim. Acta*, 2005, **428**, 147.
- 56 M. R. Molla, S. Rana, M. A. Rub, A. Ahmed and M. A. Hoque, *J. Surfactants Deterg.*, 2018, **21**, 231.
- 57 S. Mahbub, M. Rahman, S. Rana, M. A. Rub, M. A. Hoque and M. A. Khan, *J. Surfactants Deterg.*, 2019, **22**, 137–152.
- 58 N. Azum, M. A. Rub and A. M. Asiri, *J. Solution Chem.*, 2016, **45**, 791.
- 59 M. A. Rub, N. Azum, F. Khan and A. M. Asiri, *J. Chem. Thermodyn.*, 2018, **121**, 199.
- 60 G. McDonnell and A. D. Russell, *Clin. Microbiol. Rev.*, 1999, **12**, 147.
- 61 F. Khan, M. A. Rub, N. Azum and A. M. Asiri, *J. Phys. Org. Chem.*, 2018, **31**, e3812.
- 62 D. Kumar, S. Hidayathulla and M. A. Rub, *J. Mol. Liq.*, 2018, **271**, 254.
- 63 D. Kumar and M. A. Rub, *J. Mol. Liq.*, 2017, **238**, 389.
- 64 M. K. Al-Muhanna, M. A. Rub, N. Azum, S. B. Khan and A. M. Asiri, *J. Dispersion Sci. Technol.*, 2016, **37**, 450.
- 65 G. Stainsby and A. Alexander, *Trans. Faraday Soc.*, 1950, **46**, 587.
- 66 M. A. Hoque, M. M. Rashid, M. R. Molla and M. A. Rub, *J. Solution Chem.*, 2017, **46**, 682.
- 67 O. Owoyomi, J. Ige and O. Soriyan, *Chem. Sci. J.*, 2011, **25**, 1.
- 68 I. Jelesarov and H. R. Bosshard, *J. Mol. Recognit.*, 1999, **12**, 3.
- 69 K. S. Sharma and A. K. Rakshit, *J. Surfactants Deterg.*, 2004, **7**, 305.
- 70 M. Rahman, M. A. Khan, M. A. Rub, M. A. Hoque and A. M. Asiri, *J. Chem. Eng. Data*, 2017, **62**, 1464.
- 71 A. Rakshit and B. Sharma, *Colloid Polym. Sci.*, 2003, **281**, 45.
- 72 R. Jha and J. C. Ahluwalia, *J. Chem. Soc., Faraday Trans.*, 1993, **89**, 3465.
- 73 H. S. Nalwa, *Handbook of surfaces and interfaces of materials, five-volume set*, Elsevier, 2001.
- 74 L.-J. Chen, S.-Y. Lin and C.-C. Huang, *J. Phys. Chem. B*, 1998, **102**, 4350.
- 75 M. Rahman, M. A. Hoque, M. A. Khan, M. A. Rub and A. M. Asiri, *Chin. J. Chem. Eng.*, 2018, **26**, 1110.
- 76 R. Lumry and S. Rajender, *Biopolymers*, 1970, **9**, 1125.
- 77 G. Sugihara and M. Hisatomi, *J. Colloid Interface Sci.*, 1999, **219**, 31.
- 78 E. Mack Jr, *J. Am. Chem. Soc.*, 1925, **47**, 2468.
- 79 M. Mesleh, J. Hunter, A. Shvartsburg, G. C. Schatz and M. Jarrold, *J. Phys. Chem.*, 1996, **100**, 16082.

