Thermodynamic Properties of Potassium Lauryl Sulfate, Sodium Lauryl Sulfate and Sodium Lauryl Benzene Sulfonate (KDS, NaDS, and NaBDS) in Alcohol-Water System in Presence of Urea at Various Temperatures

Yogyta Singh*, Purva Saxena and Pratima Jain
Department of Chemistry, Government KRGPG (Autonomous) College, Gwalior, M.P

*Corresponding author: Yogyta Singh, Department of Chemistry, Government KRGPG (Autonomous) College, Gwalior M.P, Tel: +918004684253; E-mail: Yogitapatel1505@gmail.com

Received: June 14, 2017; Accepted: July 31, 2017; Published: August 04, 2017

Abstract

Surface Physico-chemical and thermodynamic studies of some aqueous surfactant KDS, NaDS, NaDBS, solution were carried out as a function of molality and temperature. Thermodynamic properties of micellization behavior of anionic surfactants KDS, NaDS, NaDBS, in presence of different concentration of urea and butanol-water were studied at 303 K, 308 K, 313 K, 318 K. CMC and degree of ionization of the micelle, β, were estimated from the dependence of the specific conductivity on molality. The temperature dependence of lnXcmc was studied by the curve lnXcmc/temperature by keeping in mind Muller equation.

Keywords: KDS; NaDS; NaDBS; Enthalpy; Entropy; Gibbs free energy; Conductivity

Introduction

The chain reaction of cleaner on the micellization bearing of ionic surfactants in aqueous solutions has been highly studied long using different techniques [1]. The CMC of surfactants in recent literature has been defined as the concentration of the surfactants solution at which the molecule self-aggregate to from spherically shaped micelles. The CMC indicates commonlinear order of absorption separation breaking point, at down which highest of the monomeric case and beyond which basically all further surfactants access the micelle case. The reviews are being focused on the significance of extreme organic solvents on the stock behavior of surfactants and the structure properties of aggregates formed [2-4]. The micelle design of a fluid solution knows to be changed by inorganic supplement and this has been various surveys as regards the effect of organic additives on the CMC of anionic surfactants. Kabir-ud-Din [5] announced that CMC of anionic and non-anionic surfactants greatly increase with adding of urea in fluid solution.

© 2017 Trade Science Inc.
The effect of alcohol-water solvent system with different composition of additive urea and varying temperature on the collection behavior of aqueous solution of KDS, NaDS, NaDBS has been study through conductivity measurements. The plots of conductivity vs. temperature were used to determine the CMC and degree of ionization, \( \beta \), of different salts KDS, NaDS, NaDBS micelles at the different molality and temperature. The degree of micelle ionization was inversely proportional to mixture composition but directly propositional to surfactant concentration temperature, there trends are indicative of the interaction of SDS ionic head group and the association of SDS molecule with the binary mixed solvents [6].

The free energy, entropy, and enthalpy, of micellization additionally the variation in heat capacity (\( \Delta C_{p,m}^{*} \)) of surfactant are the monomeric micelle structures on the CMC, are given by using equations:

\[
\Delta H_{m}^{*} = -RT \frac{d \ln(CMC)}{dT} \quad (1)
\]

Free energy of micellization is obtained using relation [7]:

\[
\Delta G_{m}^{*} = RT \ln(CMC) \quad (2)
\]

Where \( R \) is the universal gas constant, \( T \) is the absolute temperature and CMC is the critical micelle concentration.

Entropy of micellization was calculated using this equation:

\[
\Delta S^{*} = \frac{\Delta H^{*} - \Delta G^{*}}{T} \quad (3)
\]

Heat capacity of micellization is obtained by using relation [8]:

\[
\Delta C_{p,m}^{*} = \left[ \frac{d \Delta H_{m}^{*}}{dT} \right] = T \left[ \frac{d \Delta S_{m}^{*}}{dT} \right] \quad (4)
\]

These analyzing the temperature dependent on CMC in aqueous solution of KDS, NaDS, NaDBS were taken thermodynamic parameters derived from these equation were found out.

**Materials and Methods**

The recrystallized KDS and urea were used to prepare KDS solution by mixing required quantity of alcohol in water. After recrystallization it was used for experiment. Triple distal water was used to prepare aqueous solution all throughout studies [9].

Conductivity of the solution was measured in an Elico conductivity meter at frequentness of 1000 Hz applying platinized electrode called common cell habitual. Applicable changing was studied for the changes of conductivity of clear water with temperature. The critical micelle concentration character was determined on the break points of around two linear lines portions in the conductivity vs. concentration point. The critical micelle concentration of surfactant in residence of urea, monohydric alcohols in water at various temperatures limits (303 K, 308 K, 313 K, and 318 K).

**Results and Discussion**

These points of particular conductivity vs. temperature are declared in the FIG. 1, for aqueous solution of KDS of molalities within 0.003 m and 0.023 m. The pre-micellar absorption limits on the specific conductivity of KDS increment in straight line upon temperature. Found in molalities larger than the critical micelle concentration, the degree of increments of specific conductivity become for temperature that can be assumed with regular increase in the degree of micelles ionization.
The degree of micelles ionization increased with increasing in the chain length, but it decreases with increasing surfactants concentration.

Every unit of the either conductivity calculated at the given temperature. The standard isotherm for KDS solution is shown in FIG. 2. The molalities and conductivity increases with increase in temperature. The CMC standards are calculated by the interaction of the straight-line fragmentation of isotherm. The proportion of slopes of the straight-line fragment of the isotherm alike to the post and pre-micelle molality range is appropriate like the degree of micelle ionization $\beta$. The degree of micelle ionization of KDS was determined by the ratio of slope of specific conductance vs. molality curves above and below the variation point at different temperature.

Degree of micelles ionization of a rises according to the temperature indication that counter ions all being desorbed from the micelle during dilution and same effect observed with increasing temperature [10].
FIG. 3. Temperature depends on the degree of micelles ionization $\beta$, of potassium lauryl sulfate solution.

The degree of micelle ionization or apparent degree of counter ion dissociation $\beta$ is defined by asserting that the aggregation number is dependent only on the concentration, $C$, of counter ions in the aqueous solution [11]. Temperature dependence of the calculated $\beta$ value of the analyzing salts FIG. 3, shows $\beta$ rises of temperature. The degree of micelle ionization increased with increasing in the chain length. Let us assume furthermore that the typical changes in the heat capacity vary vertically by means of temperature following Muller [12] equation is found.

$$
\ln X_{\text{cmc}} = \ln X_{\text{cmc}} \left[ 1 + \frac{\alpha}{\beta T} \left( \frac{T - 2 - \alpha}{\beta T} \right) + \frac{1}{2 - \alpha} \left( 1 + \frac{\alpha}{2 - \beta T} \right) \right] + \frac{\Delta C_{\text{pm}}}{(2 - \alpha) R} \left[ 1 - \frac{T}{T_c} - \frac{\alpha}{2 - \beta T} \left( \frac{T^* - 2 - \alpha}{T^* - 2 + \alpha} \right) \right] (5)
$$

Where $\beta$ is the degree of ionization; $\Delta C_{\text{p,m}}$ is the heat capacity.

FIG. 4. Temperature depends on $\ln X_{\text{cmc}}$ (1) KDS, (2) NaDS, (3) NaDBS.
In the FIG. 4, lnX_{cmc} is designed next to temp of the three salt compounds beneath assessment. lnX_{cmc} increase with increasing temperature, but the rate of increase of lnX_{cmc} with temperature is different in different salts. This increase is almost constant in case of KDS. In case of NaDS the rate is first constant but higher after certain temperature, while in case of NaDBS, the increase of rate is having anomalous behavior these trends are against Muller equation (TABLES 1-3).

**TABLE 1. CMC value for various surfactant salts at different temperature.**

<table>
<thead>
<tr>
<th>T (K)</th>
<th>CMC (mol/litre)</th>
<th>KDS</th>
<th>NaDS</th>
<th>NaDBS</th>
</tr>
</thead>
<tbody>
<tr>
<td>303</td>
<td>0.0375</td>
<td>0.0466</td>
<td>0.00679</td>
<td></td>
</tr>
<tr>
<td>308</td>
<td>0.0410</td>
<td>0.0505</td>
<td>0.00705</td>
<td></td>
</tr>
<tr>
<td>313</td>
<td>0.0450</td>
<td>0.0552</td>
<td>0.00853</td>
<td></td>
</tr>
<tr>
<td>318</td>
<td>0.0500</td>
<td>0.0640</td>
<td>0.00872</td>
<td></td>
</tr>
</tbody>
</table>

**TABLE 2. Parameters resulting from the ΔC_{p,m} at various temperature.**

<table>
<thead>
<tr>
<th>T (K)</th>
<th>ΔC_{p,m} (heat capacities)</th>
<th>KDS</th>
<th>NaDS</th>
<th>NaDBS</th>
</tr>
</thead>
<tbody>
<tr>
<td>303</td>
<td>-4.4</td>
<td>-3.9</td>
<td>-1.8</td>
<td></td>
</tr>
<tr>
<td>308</td>
<td>-5.3</td>
<td>-5.1</td>
<td>-11.0</td>
<td></td>
</tr>
<tr>
<td>313</td>
<td>-6.9</td>
<td>-9.8</td>
<td>-1.4</td>
<td></td>
</tr>
<tr>
<td>318</td>
<td>-7.1</td>
<td>-7.8</td>
<td>-6.1</td>
<td></td>
</tr>
</tbody>
</table>

**TABLE 3. lnX_{cmc} for various salts at various temperature T.**

<table>
<thead>
<tr>
<th>T (K)</th>
<th>lnX_{cmc}</th>
<th>KDS</th>
<th>NaDS</th>
<th>NaDBS</th>
</tr>
</thead>
<tbody>
<tr>
<td>303</td>
<td>-9.58</td>
<td>-9.36</td>
<td>-8.98</td>
<td></td>
</tr>
<tr>
<td>308</td>
<td>-9.49</td>
<td>-9.28</td>
<td>-8.95</td>
<td></td>
</tr>
<tr>
<td>318</td>
<td>-9.29</td>
<td>-9.04</td>
<td>-8.73</td>
<td></td>
</tr>
</tbody>
</table>

The parameter enable us to estimate the standards of free energy of the micelle, The surfactants like a role of temperature over and above the enthalpy and entropy contribution toward ΔG_{m} illustrates the tempt dependence of ΔG_{m} for the three surfactants. The FIG. 5 explains that KDS and NaDS show almost equal results. And increase in temperature there are regular decrease in free energy and trend follows in NaDBS but is more drastic change than that of KDS and NaDS. The plots clarify that the value of the free energy of formation of KDS, NaDS, NaDBS micelles are negative at different temperature. They become more negative with increasing temperature, showing increasing stability.
While studying effect of enthalpy or entropy with increasing temperature KDS and NaDS are showing same behavior is decreasing and also increasing enthalpy or entropy with increasing temperature (FIG. 6 and 7). However breaks are seen in both the cases that may be assigned to CMC. Probably benzene ring in the long chain does permit electrons to more freely
and create hindrance; hence it is showing abnormal behavior when compared to KDS or NaDS. NaDBS is showing abnormal behavior first enthalpy or entropy both are decreasing sharply with increasing temperature, but after 308 K sharp increase is seen up to 313 K and again decreases after 318 K, this is due to the presence of benzene ring.

If we caste a glance at structures of KDS, NaDS and NaDBS we find that KDS and NaDS are having similar structures except in cation. KDS has a better cleansing action indicated by the presence of the potassium ion. NaDBS again due to the presence of benzene ring does not easily undergo oxidation and show abnormal behavior.

**Conclusion**

From the results obtained, it was shown that degree of micelle ionization $\beta$, molality of salts KDS, NaDS and NaDBS, may be projected from the plot of definite conductivity vs. molality. The $\beta$ morals were originated to be significantly superior of the KDS than the NaDBS. The CMC standards reveal linearly reliance on temperature and chain length. It was also shown that temperature dependence of $\ln X_{\text{cmc}}$ can be fitted to Muller equation partly. We calculated the temperature dependencies of $\beta$, and $\Delta C_p^{m}$ for the micellization. From these parameters are enthalpy, entropy and Gibbs free energy of micelles are expected as a purpose of temperature.

**REFERENCES**

