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# Corrosion and hydrogen evolution rate control for X-65 carbon steel based on chitosan polymeric ionic liquids: experimental and quantum chemical studies†

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The corrosion performance of carbon steel was tested in four polymeric ionic liquids (PILs) that differed only in the fatty acid linked to the chitosan (CS) amine group. The measurements were implemented involved the hydrogen evolution rate (HER), gravimetric measurements, potentiodynamic polarization (PDP), electrochemical impedance spectroscopy (EIS), and quantum chemical estimations. The morphology and the elements arranged on the metal were considered by a scanning electron microscopy (SEM) system attached to an energy dispersive X-ray (EDX) system. The addition of polymeric ionic liquids hindered the rate of hydrogen generation. The order of the inhibitors efficiency was CSPTA-lauric > CSPTA-myristic > CSPTA-palmitic > CSPTA-stearic. The polarization method proved that the percentage inhibition efficiency increases with increasing the inhibitors concentration in 1 M HCl, representing a drop in the corrosion rate of carbon steel. On the other hand, the percentage inhibition decreased with the increase in temperature. Quantum chemical calculations revealed that the tested ionic liquids could react with the iron surface *via* electron transfer from the metal atom to ionic liquid molecule.

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## 1. Introduction

Developing environmentally friendly polymeric ionic liquids (PILs) has attracted great interest due to their unique chemical and physical properties. These properties are the key factors for considering PILs for green innovative applications because ILs can meet green environmental requirements, such as related to their purity, non-toxicity, and commercial accessibility. In addition, they have the ability to inhibit the corrosion of materials used for pipelines, tanks, and technical equipment.<sup>1,2</sup>

Ionic liquids (ILs) are salts that are organic liquids composed of unpacked well ions, permitting free movement, and consequently have flow properties.<sup>3,4</sup> Moreover, there is a great interest in ILs as green corrosion inhibitors because of their large number of advantages, such as low toxicity, high thermal steadiness, low volatility, elevated ionic conductivity, and great action as corrosion inhibitors.<sup>5–8</sup> Among these features of ILs, their very small vapor pressure (will not vaporize), non-polluted environmentally friendly nature are attractive and make them less harmful as metal corrosion inhibitors. The metal corrosion

inhibited by PILs is because of their adsorption on the iron surface.<sup>9–11</sup>

Imidazolium ILs and their derivatives are one of the most effective and the most commonly used ILs as corrosion inhibitors in different corrosive media for carbon steel.<sup>12–17</sup> In addition, it was reported that quaternary ammonium,<sup>18,19</sup> phosphonium,<sup>20</sup> and pyridinium salts ILs<sup>21,22</sup> could be used for the protection of steel. Taghavikish *et al.*<sup>23</sup> reported the improved anticorrosion ability for PILs based on thiol-ene derivatives. In addition, Atta *et al.*<sup>24</sup> investigated the enhanced anticorrosion ability of hyperbranched PILs. The results from the present work are in good agreement with our previous work<sup>25</sup> in which PILs were prepared based on acrylamides followed by quaternization with different aliphatic tertiary amines.

Chitosan (CS) ionic liquid polymers are attractive materials to apply as green corrosion inhibitors.<sup>26–28</sup> CS is a harmless, ecofriendly decomposable, and  $\beta$ -D-glucose(1,4)amine polysaccharide linked with *N*-acetyl glucosamine.<sup>29,30</sup> It is produced from the deacetylation of chitin,<sup>31–33</sup> the most widespread polysaccharide in the environment after cellulose.<sup>34</sup> It is found in shrimp and crab and insects cuticles.<sup>35</sup> CS and its products have widespread potential in the field of medicine, paper, textile, food, cosmetics, and several other industries.<sup>36–38</sup>

Several new materials can be produced from CS because of its high capability of being functionalized. Modifications of CS ionic functional groups (*e.g.*, NH<sub>2</sub> or OH groups) allow it to be applied

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in several manufacturing processes, owing to its improved water solubility over CS itself.<sup>39–42</sup>

The aim of this research was to prepare PILs from chitosan by means of interacting with *p*-toluene sulfonic acid. Then, amidation with some fatty acids, for example, stearic (C<sub>18</sub>), palmitic (C<sub>16</sub>), myristic (C<sub>14</sub>), and lauric (C<sub>12</sub>) acids. Moreover, we aimed to study the control of hydrogen generation by PILs derivatives. In addition, the electrochemical steel anticorrosion behavior in 1 M solution of HCl was inspected by way of different techniques using potentiodynamic polarization and impedance spectroscopy (EIS) (Nyquist and Bode plots). Chemical quantum estimations were approved to elucidate the inhibition process by the prepared PILs at a molecular level. Furthermore, the surface morphology was examined through scanning electron microscopy (SEM) to detect the effect of the adsorbing films.

## 2. Experimental

### 2.1. Preparation of PILs inhibitors

**2.1.1. Materials.** Chitosan (CS), dicyclohexyl carbodiimide (DCC), stearic acid, palmitic acid, myristic acid, lauric acid, and *p*-toluene sulfonic acid (PTSA) were acquired from Aldrich Chemical Co. Hydrogen chloride was attained from Merck Company.

Carbon steel (X-65 type) was the working electrode used with a composition: C 0.10, Si 0.21, Mn 1.53, P 0.02, S 0.06, Ni 0.03, Cr 0.05, Mo 0.003, V 0.004, Cu 0.05, Al 0.01, and the rest was Fe. These samples were obtained from an unused petroleum pipeline.

#### 2.1.2. Procedure

(a) *Preparation of chitosan-p-toluene sulfonate salt (CSPTA).* First, 4.8 g of PTSA was solubilized in 80 ml distilled water and then mixed with CS (3.8 g) in a flask connected to a reflux condenser with stirring under an inert atmosphere. Then, the temperature was raised to 70 °C and sustained for 24 h. Last, acetone was added to the reacted mixture, which was washed with ether and dried in a vacuum oven at 40 °C.

(b) *CSPTA amidation reaction with fatty acids.* First, 2 g of CSPTA was stirred with 100 ml of water until dissolved and 1.3 g of fatty acid was mixed with 80 ml methanol then added with stirring well. The fatty acid/CSPTA solution molar ratio was 0.34 : 1. DCC (15 ml) dissolved in methanol (0.08 g l<sup>-1</sup>) was further added at room temperature. The reaction was finished under shaking in a nitrogen atmosphere for 20 h. Finally, precipitation was performed using 250 ml acetone, followed by filtration, washing with diethyl ether, absolute methanol, and finally vacuum drying at 20 °C.

**2.1.3. Spectroscopic characterization.** Fourier transform infrared (FTIR) spectrophotometry (Bruker) was used to complete the infrared (IR) analyses and <sup>1</sup>H-NMR spectra were recognized on a spectrometer 400 MHz Avance Bruker DRX-400 using DMSO-d<sub>6</sub> solvent.

### 2.2. Hydrogen evolution measurements

A water replacement method was conducted to evaluate the hydrogen evolution rate similar to that formerly described.<sup>43,44</sup>

Initially, 100 ml of 1 M HCl (the corrosive medium) was placed into a container and a carbon steel coupon with dimensions of 3.5 cm × 2.5 cm was dipped in to the test solution. Then, the container was quickly closed to stop any loss of hydrogen gas. Finally, at fixed time periods, the hydrogen gas volume generated throughout the corrosion reaction was practically recorded from the volume of gas replacing the water level in the burette (in cm<sup>3</sup>).

### 2.3. Electrochemical measurements

A Voltalab 80 Potentiostat PGZ 402 Tacussel Radiometer was used to quantify the electrochemical experiments. The technique was controlled by Voltmaster-4 software. Tests were performed *via* an electrochemical cell of 100 ml glass with spaces for 3 electrodes filled with 100 ml of corrosive medium (1 M HCl). The three electrodes comprised a saturated calomel electrode (SCE) as the reference electrode, a platinum electrode as an auxiliary electrode, and carbon steel working electrode (WE).

Before all the experiments, the steel electrode surface was polished manually with different emery papers below 2500 grade. In order to gain the open-circuit potential, the electrode potential was steadied for 1 h in test solution previously to initiate the measurements. The electrode zone subjected to the destructive acidic media was 1 cm<sup>2</sup> and the total measurements were employed at room temperature. Besides, the potential was altered from -900 to -300 mV vs. SCE, with a scan rate of 2 mV s<sup>-1</sup> to acquire the potentiodynamic curves.

Nyquist and Bode plots were utilized assumed to define the EIS with a frequency range between 100 kHz and 50 mHz with an amplitude of 10 mV.

### 2.4. Gravimetric experiments

The effect of various temperatures on the corrosion rate (mg cm<sup>-2</sup> h<sup>-1</sup>) of the carbon steel was attained by performing the weight loss test for the carbon steel samples in 1 M HCl solution with and without inhibitors (250 ppm). The carbon steel sheets with the dimension of 2 cm × 2.5 cm × 0.3 cm were immersed in test solution for 8 h. After that, the carbon steel samples were washed by distilled water and ethanol, dried in cold air, and then weighed using an analytical balance.

### 2.5. Surface morphology studies

A scanning electron microscopy (SEM) instrument Quanta Model 250 FEG joined with EDX Part (energy dispersive X-ray) analyses was utilized for this examination with an accelerating voltage of 30 kV and a magnification force of *X* = 2000. Surface morphology was tested after dipping in the uninhibited solution (blank) and in an inhibited solution of 100 ppm of the CSPTA-lauric acid inhibitor.

### 2.6. Chemical quantum calculations

PILs molecular structures were geometrically modernized by density functional theory (DFT) method by means of the 3-21 G\*\* premise set with Hyperchem 7.5. Chemical quantum

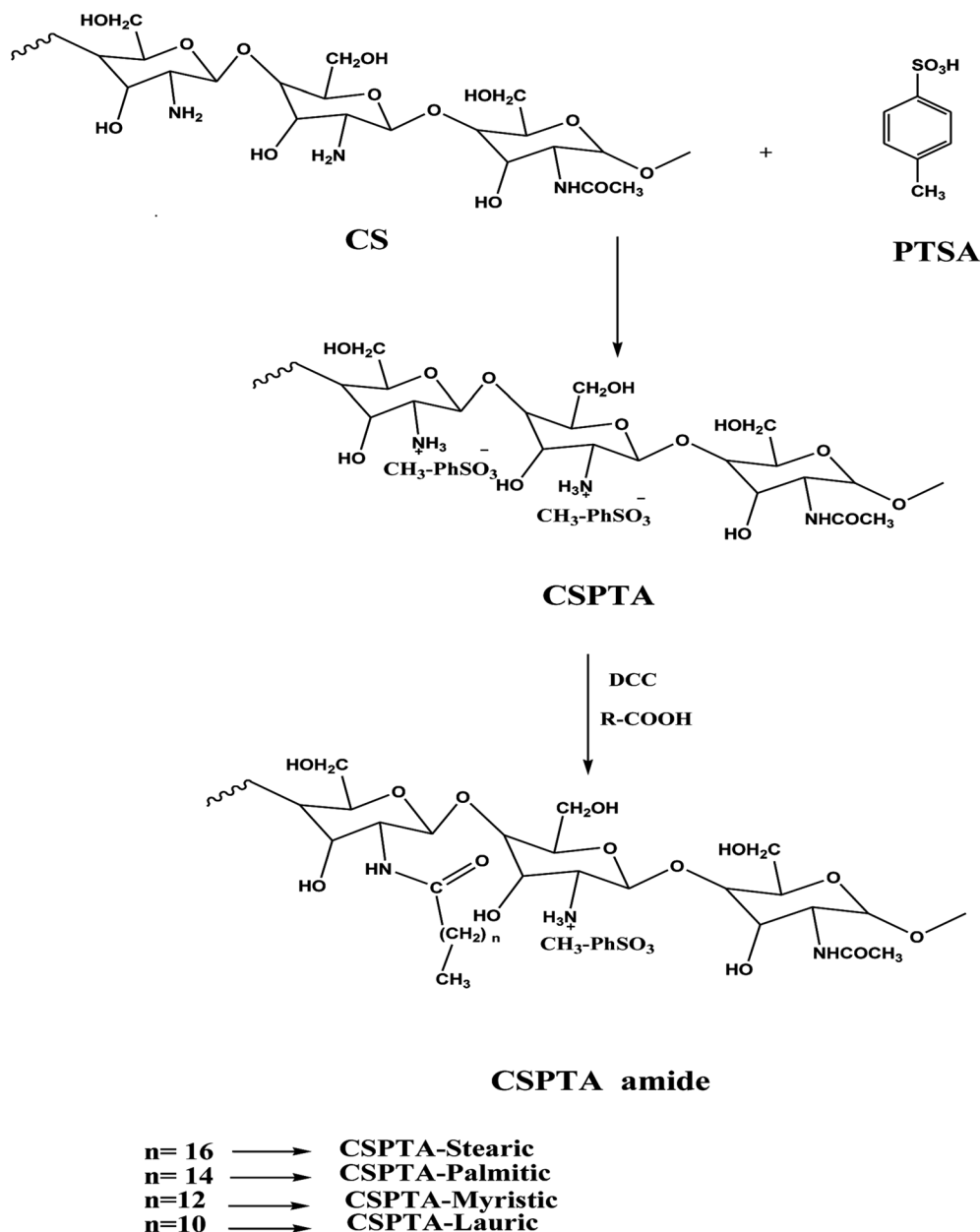
calculation parameters (*e.g.*,  $E_{\text{HOMO}}$  (the highest occupied molecular orbital energy),  $E_{\text{LUMO}}$  (the lowest unoccupied molecular orbital energy), energy gap (energy difference between the LUMO and HOMO energy levels),  $X$  (electronegativity),  $\eta$  (hardness), softness ( $\sigma = 1/\eta$ ), chemical potential ( $\pi = -X$ ), and number of electron transferred ( $\Delta N$ )) were assessed for the prepared PILs molecules.

### 3. Results and discussion

Chitosan is sparingly soluble in water but its solubility can occur in basic aqueous solution through deprotonation of the CS amino group or it is soluble in slightly acidic medium.<sup>45,46</sup> The present work aimed to make CS soluble in water by reacting

the amine group with PTSA. Then, the ionic liquid product was amidated through stearic, palmitic, myristic, and lauric acid with the addition of DCC as a catalyst to yield the CS polymeric ionic liquid, as represented in Scheme 1.

FTIR and <sup>1</sup>H-NMR analyses were used to elucidate the chemical structures of the formed CSPTA and CSPTA-lauric. The synthesis of CSPTA was elucidated by FTIR spectroscopy (S1a†), which displayed the distinguish broad -OH band at 3444 cm<sup>-1</sup>, bands at 3100, 1637, and 1010 cm<sup>-1</sup> for C-H stretching, C=C bonds, and bending of phenyl group of PTSA, respectively. Also, it showed a band at 2923 cm<sup>-1</sup> assigned to the CH aliphatic of CS and a band at 1210 cm<sup>-1</sup> for stretching C-N. The appearance of a band at 1070 and 1345 cm<sup>-1</sup> could be recognized as due to S=O bond in PTSA, which points to CSPTA formation.



Scheme 1 Quaternization of CS amine group by PTSA, followed by amidation with different saturated fatty acids.

Fig. 1(a) shows the FTIR spectrum of CSPTA-lauric, which displays a strong absorption band at  $1693\text{ cm}^{-1}$ , indicating C=O amides. The strong band at  $1693\text{ cm}^{-1}$  proves the amide linkage formation between the CS amino group and lauric acid carboxyl group. A band appearing at  $3322\text{ cm}^{-1}$  was assigned to stretching N-H. Peaks at  $1625$  and  $937\text{ cm}^{-1}$  were recognized due to C=C bonds and bending of CH for the aromatic ring in PTSA. Also, peaks appearing at  $1217$  and  $1301\text{ cm}^{-1}$  are for C-N and S=O stretching, respectively.

In addition, the  $^1\text{H-NMR}$  spectra for CSPTA and CSPTA-lauric are given in S1b† and Fig. 1(b), respectively. These display signals at shifts of  $7.1\text{ ppm}$  (dd,  $2\text{H}$ ,  $J = 6.03\text{ Hz}$ ), and  $7.5\text{ ppm}$  (dd,  $2\text{H}$ ,  $J = 6.25\text{ Hz}$ ) and peaks at  $8.2\text{ ppm}$  signifying  $^+\text{NH}_3$ - and proving the reaction of amino groups of CS with PSTA in S1b.† This peak at  $8.2\text{ ppm}$  was slight in Fig. 1(b), indicating that the amidation occurs but not with all  $-\text{NH}_2$  groups. In Fig. 1(b), the appearance of new peaks at  $1.9$  (t,  $3\text{H}$ ,  $\text{CH}_3$ - $(\text{CH}_2)_{10}$ ) and  $3.3$  (m,  $\text{CH}_3$ - $(\text{CH}_2)_{10}$ ) confirmed the amidation. The disappearance of the signal at  $2.8\text{ ppm}$  (s,  $2\text{H}$ , C- $\text{NH}_2$ ) pointed to that there were

no available amine groups after reacting the CS with PTSA and lauric acid. In S1b,† the signals at  $\delta$   $3.5$ – $3.9\text{ ppm}$  confirmed the methylene groups of CS, with a peak at  $1.84\text{ ppm}$  (s,  $3\text{H}$ ,  $-\text{NHCO}-\text{CH}_3$ ) in CSPTA and signal at  $2.5\text{ ppm}$  (s,  $3\text{H}$ ,  $\text{CH}_3$ -Ph) in PTSA, while the signals at  $2.8$  and  $4.7\text{ ppm}$  could be attributed to  $-\text{NH}$  and  $-\text{OH}$ , respectively.

### 3.1. Hydrogen evolution measurements

Fig. 2 and S2† show the volume of hydrogen progressed from carbon steel corrosion in solution of  $1\text{ M HCl}$  with time in the absence and presence of inhibitors. There was a clear and remarkable increase for hydrogen production with the increasing steel immersion time. From the volume produced, the hydrogen generation rate ( $H_r$ ) was evaluated *via* the next expression:<sup>47</sup>

$$H_r = (V_2 - V_1)/(t_2 - t_1) \quad (1)$$

where  $V_2$  and  $V_1$  are the hydrogen-evolved volumes at  $t_2$  and  $t_1$  time, respectively. Adding many concentrations of CSPTA-

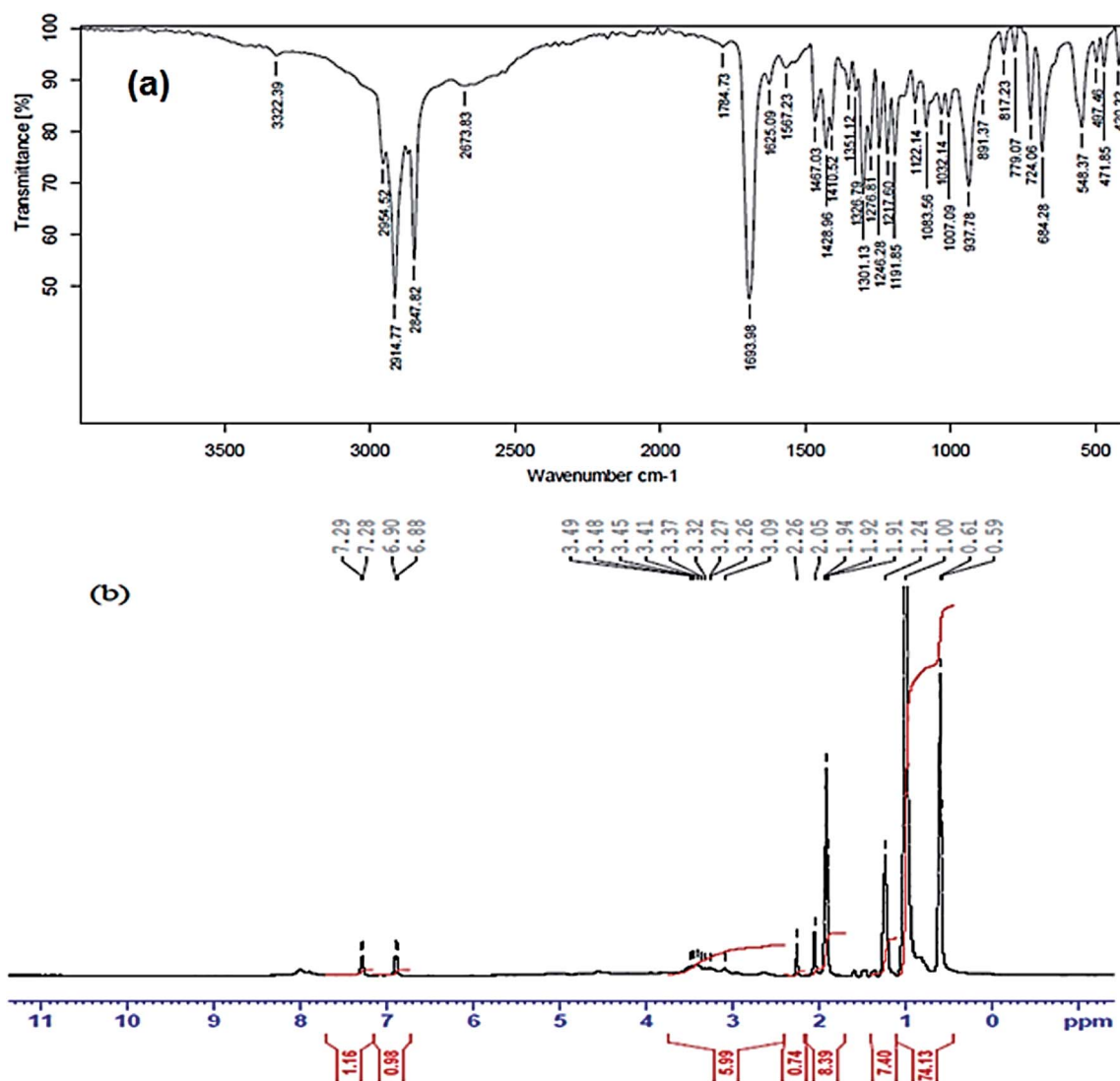


Fig. 1 Chemical structure characterization of CSPTA-lauric inhibitor: (a) FTIR, (b)  $^1\text{H-NMR}$ .

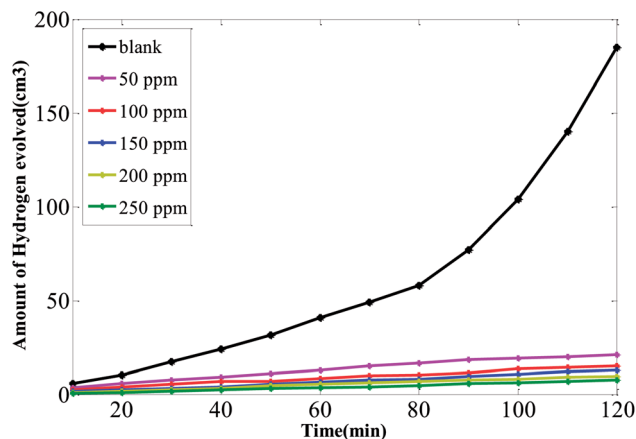


Fig. 2 Volume of hydrogen evolved with time for carbon steel in 1 M HCl at various concentrations of CSPTA-lauric.

stearic, CSPTA-palmitic, CSPTA-myristic, and CSPTA-lauric affect the hydrogen generation rates ( $H_T$ ), as revealed in Fig. 3. The data show that  $H_T$  is reduced with the rise in the prepared PILs concentrations. Moreover, the efficiency of PILs ( $I_H\%$ ) for governing the hydrogen evolution was considered using the next relationship:<sup>44</sup>

$$I_H\% = [1 - (H_T/H_{T_0})] \times 100 \quad (2)$$

where  $H_{T_0}$  and  $H_T$  are the hydrogen generation rates in the absence and presence of the PIL inhibitors, respectively. Fig. 4 illustrates the plot of inhibitors efficiency ( $I_H\%$ ) versus logarithmic concentrations of the PILs inhibitors. The plot demonstrates that the inhibition effect of the PILs rises as their concentrations increase. Besides, the order of effectiveness of the inhibitors was CSPTA-lauric > CSPTA-myristic > CSPTA-palmitic > CSPTA-stearic. This indicates that CSPTA-lauric is the most efficient inhibitor, which has the shortest chain among the prepared PILs. The difference between the inhibitors' efficiency may be attributable to the spacer length, whereby an inhibitor with a longer spacer has low performance due to the longer distance between the two head hydrophobic groups

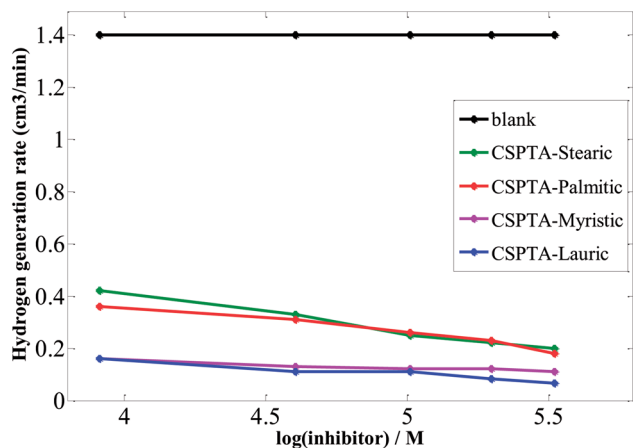


Fig. 3 Relation between hydrogen generation rates vs. logarithmic inhibitor concentrations for carbon steel in 1 M HCl.

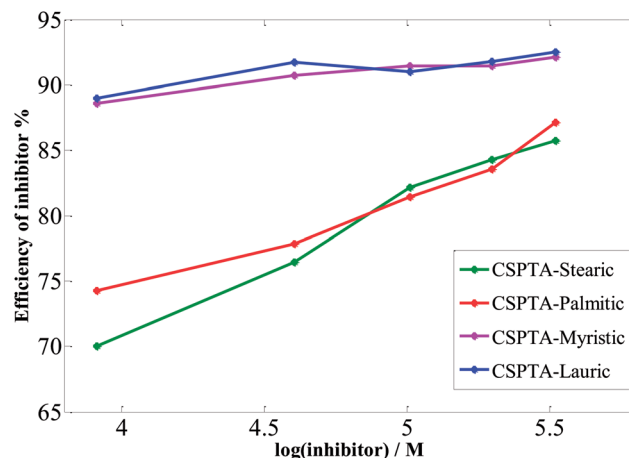


Fig. 4 Variation of the efficiency of inhibitors with logarithmic inhibitor concentrations in 1 M HCl.

in the structure, which decreases the charge density of the head groups and lowers the adsorption of the inhibitor.<sup>48–52</sup>

These inhibitors inhibit the dissociation of carbon steel in HCl and accordingly, hinder the cathodic hydrogen evolving reaction *via* adsorption at the metal/acid solution interface.<sup>53,54</sup> The ability of these PILs inhibitors as hydrogen evolving inhibitors is influenced by their structures. Inhibitors are able to arrange a thin film on the metal surface. Additionally, the chemisorption on the surface was increased by the PILs' chemical structures owing to coordination bonds among the metal surface and the electron pairs on N, O, and S atoms in the PILs' structures, the CO group *p*-electron, and as protonated moieties.<sup>55</sup>

Table 1 Values of the mean and standard deviation (SD) for the amount of hydrogen evolved for carbon steel electrode at different concentrations of PILs inhibitors in 1 M HCl solution

Inhibitor	Concentration (ppm)	Mean value	SD value
Blank	000	61.95	±55.64
CSPTA-stearic	50	27.57	±15.03
	100	21.23	±12.08
	150	16.69	±8.864
	200	15.79	±8.12
	250	12.63	±7.35
CSPTA-palmitic	50	22.45	±13.01
	100	19.87	±11.24
	150	16.15	±9.284
	200	13.48	±8.327
CSPTA-myristic	250	10.25	±6.4
	50	11.28	±5.598
	100	8.642	±4.408
	150	8.442	±4.539
CSPTA-lauric	200	7.825	±4.236
	250	7.283	±3.932
	50	13.29	±5.94
	100	9.067	±4.126
	150	7	±3.85
	200	5.5	±2.922
	250	3.917	±2.318

Values of the mean and standard deviation (SD) for the amount of hydrogen evolved at different concentrations of PILs inhibitors in 1 M HCl solution are recorded in Table 1. In addition, values of the mean and standard deviation (SD) for the hydrogen evolution rate for the carbon steel electrode in 1 M HCl solution are recorded in Table 2.

### 3.2. Measurements of potentiodynamic polarization (PDP)

The curves of steel electrochemical polarization in 1 M HCl solution in the presence and absence of the produced PILs with the varied inhibitor concentrations are revealed in Fig. 5 and S3.† A drop in anodic and cathodic currents was presented in the presence of the inhibitor. The decrease was more obvious at higher inhibitor concentrations. This may be recognized as due to a defensive film protecting the steel surface from the corrosive medium. Adsorbing of the inhibitor on the steel surface reduces the hydrogen evolution (cathodic reaction) and reduces the iron metal dissolution (anodic reaction).

The polarization parameters, *i.e.*, corrosion potential ( $E_{\text{corr}}$ ), current density of corrosion ( $I_{\text{corr}}$ ), slopes of cathodic Tafel ( $\beta_c$ ), slopes of anodic Tafel ( $\beta_a$ ), were obtained from the polarization curves and are recorded in Table 3. The surface coverage degree ( $\theta$ ) and the inhibition efficiency (IE %) were considered from the  $I_{\text{corr}}$  values *via* the next relation:<sup>56,57</sup>

$$\theta = [1 - (I_{\text{corr}(2)}/I_{\text{corr}(1)})] \quad (3)$$

$$\text{IE \%} = [1 - (I_{\text{corr}(2)}/I_{\text{corr}(1)})] \times 100 \quad (4)$$

where  $I_{\text{corr}(1)}$  and  $I_{\text{corr}(2)}$  are the current densities for corrosion in the absence and presence of the inhibitor, respectively. The IE% values with changed inhibitor concentrations are recorded in Table 3. It was found that the IE% values increased as the inhibitor concentrations increased because of the increase in the amount of inhibitor arranged and adsorbed on the steel surface. Accordingly, this process resulted in high corrosion inhibition. The maximum effective inhibition was about 96.2% at a concentration of 100 ppm for CSPTA-lauric.

The values of IE% tracked the matching trend with those attained from the hydrogen evolution measurements but with dissimilar values related with the various techniques. Furthermore, the type of inhibitor can be regarded as an anodic or cathodic if the  $E_{\text{corr}}$  value exceeded a value of 85 mV.<sup>58-60</sup> From Table 3, it can be seen that the  $E_{\text{corr}}$  value varies with a maximum shift in  $E_{\text{corr}}$  less than 85 mV, demonstrating a mixed mode for corrosion (disturbs the anodic and cathodic reaction together). Nevertheless, the cathodic direction was more noticeable as a minor shift of  $E_{\text{corr}}$  in the cathodic trend. The potentiodynamic results reveal that the prepared PILs efficiently reduced the corrosion of steel, even used in lesser concentrations in 1 M HCl.

Values of the mean and standard deviation (SD) corrosion current density for the carbon steel electrode at different concentrations of PILs inhibitors in 1 M HCl solution are indicated in S4.†

Table 2 Values of the mean and standard deviation (SD) for the hydrogen evolution rate for carbon steel electrode in 1 M HCl solution

Inhibitor	Mean value	SD value
Blank	1.4	0
CSPTA-stearic	0.284	±0.09072
CSPTA-palmitic	0.268	±0.06979
CSPTA-myristic	0.128	±0.01924
CSPTA-lauric	0.105	±0.03651

### 3.3. Electrochemical impedance spectroscopy (EIS) measurements

The inhibition behavior in 1 M HCl for carbon steel in the absence and presence of several concentrations of CSPTA-stearic, CSPTA-palmitic, CSPTA-myristic, and CSPTA-lauric

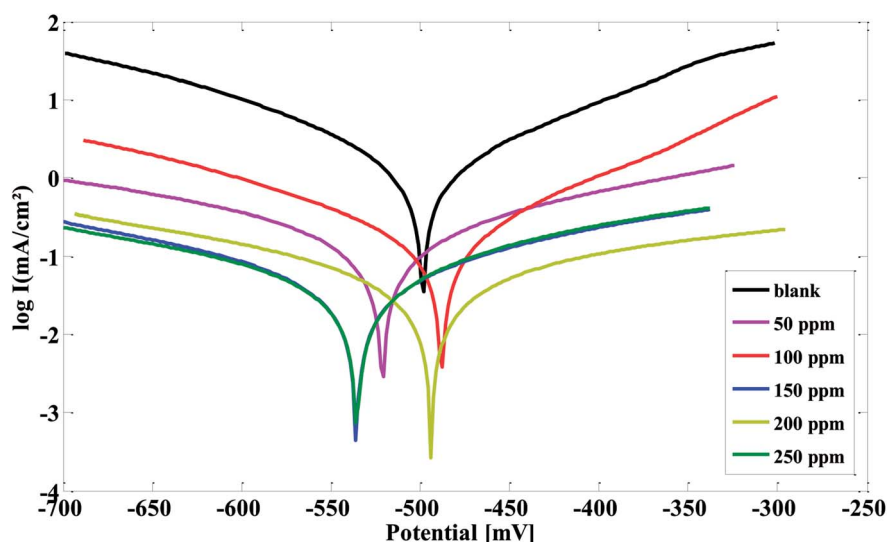


Fig. 5 Polarization plots of steel electrode obtained in 1 M HCl solution and containing various concentrations of CSPTA-lauric.

**Table 3** Corrosion parameters for carbon steel electrode in the absence and presence of different concentrations of PILs inhibitors in 1 M HCl solution

Inhibitor	Concentration (ppm)	$\beta_a$ (mV)	$\beta_c$ (mV)	$E_{\text{corr}}$ (mV)	$I_{\text{corr}}$ (mA cm <sup>-2</sup> )	$\theta$	IE%
Blank	000	162.1	-188.5	-493	3.47	—	—
CSPTA-Stearic	50	94.5	-162.6	-536	0.9	0.7406	74
	100	173.8	-155.4	-628	0.8	0.7694	76.9
	150	193.2	-143.3	-550	0.7	0.7982	79.8
	200	196.3	-148.5	-551	0.6	0.8270	82.7
	250	149.9	-149.9	-553	0.4	0.8847	88.4
CSPTA-palmitic	50	172.3	-169.4	-551	0.7	0.7982	79.8
	100	179.5	-214.6	-546	0.59	0.8299	82.9
	150	119.4	-159.3	-543	0.34	0.9020	90.2
	200	131.6	-212	-548	0.33	0.9048	90.4
	250	126.2	-173.4	-543	0.24	0.9308	93
CSPTA-myristic	50	333.8	-143.7	-518	0.4	0.8847	88.4
	100	167.8	-145.6	-504	0.3	0.9135	91.3
	150	224.5	-264.9	-519	0.19	0.9452	94.5
	200	219.9	-250.5	-521	0.18	0.9481	94.8
	250	103.4	-132.5	-488	0.14	0.9596	95.6
CSPTA-lauric	50	227.3	-260.8	-513	0.17	0.9510	95.1
	100	105	-130.5	-490	0.13	0.9625	96.3
	150	290.3	-326.2	-537	0.089	0.9743	97.4
	200	300.7	-230.2	-497	0.055	0.9841	98.4
	250	189.8	-226.5	-536.5	0.046	0.9867	98.6

(the prepared PILs) were also explored using EIS. Fig. 6 and S5† show the EIS Nyquist plots in various concentrations of the PILs inhibitors.

In Fig. 6, the diameter of capacitive loop in the Nyquist plot is enlarged as the inhibitor concentration is increased. This designates that the charge-transfer reaction is mainly controlling the steel corrosion.<sup>61</sup>

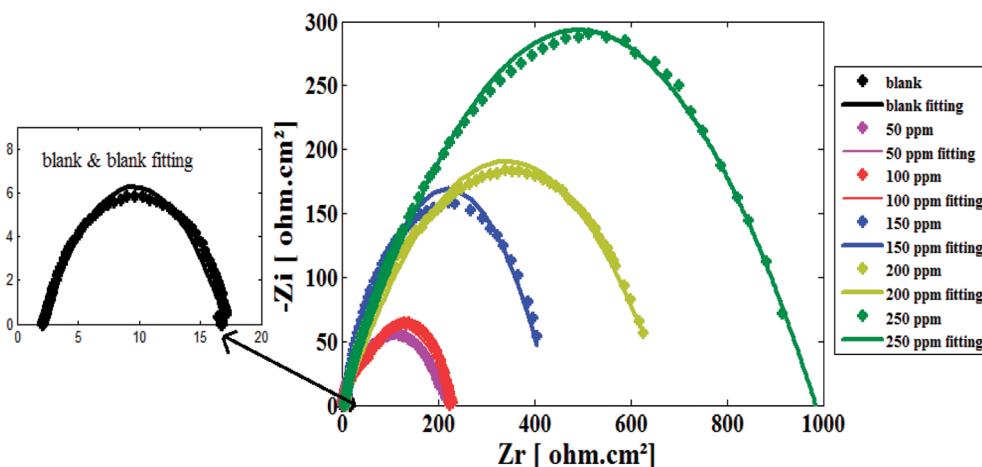
Values of  $R_c$  (charge-transfer resistance) and  $C_{dl}$  (double-layer capacitance) resulting from Nyquist plots in the absence and presence of the varied concentrations of PILs inhibitors are listed in Table 4. Values of  $C_{dl}$  are reduced, but the values of  $R_c$  are enlarged in the presence of PILs inhibitors, confirming the protecting efficiency of PILs, which is greatly reliant on their concentration. Additionally, the decrease in  $C_{dl}$  values can occur as a result of the electrical double-layer thickness, which exchanges the PILs moieties (that have a lower dielectric

constant) with the water molecules (that have a higher dielectric constant). The corrosion occurrence is typically associated with the double-layer performance.<sup>62</sup> Besides, working as inhibitors occurs by arranging and adsorption *via* substituting the water molecules existing at the steel/corrosive medium interface.<sup>63</sup>

Both the  $R_c$  and IE% values improved as the inhibitor concentration is increased, as shown in Fig. 6 and pointed out in Table 4. The inhibition effectiveness (IE%) can be estimated from the results gained by using the next relation:<sup>64,65</sup>

$$\text{IE}\% = 1 - (R_c^1/R_c^2) \times 100 \quad (5)$$

where  $R_c^1$  and  $R_c^2$  are charge-transfer resistances in the absence and presence of the inhibitor, respectively. Values of  $R_c$  were evaluated in impedances *via* variance at lower and greater frequencies. Similarly, values of IE% tracked the matching

**Fig. 6** Nyquist plots for carbon steel electrode in 1 M HCl solution with and without various concentrations of CSPTA-lauric.

**Table 4** Impedance parameters for carbon steel electrode in the absence and presence of different concentrations of inhibitors in 1 M HCl solution

Inhibitor	Concentration (ppm)	$R_c$ (Ohm)	$C_{dl}$ ( $\mu\text{F cm}^{-2}$ )	$\theta$	IE%
Blank	000	16.97	680.2	—	—
CSPTA-stearic	50	35	626.1	0.5151	51.5
	100	39	466.3	0.5648	56.4
	150	62	120.6	0.7262	72.6
	200	84	94.53	0.7979	79.7
	250	86	88.34	0.8026	80.2
CSPTA-palmitic	50	40	190.6	0.5757	57.5
	100	62	120.6	0.7262	72.6
	150	65	118.1	0.7389	73.8
	200	107	62.34	0.8414	84.1
	250	122	62.99	0.8609	86
CSPTA-myristic	50	83	88.06	0.7955	79.5
	100	160	72.09	0.8939	89.3
	150	220	45.04	0.9228	92.2
	200	248	33.01	0.9315	93.1
	250	320	40.53	0.9469	94.6
CSPTA-lauric	50	218	45	0.9321	93
	100	235	33.01	0.9352	93.5
	150	425	47.4	0.9600	96
	200	668	41.6	0.9745	97.4
	250	991	36.9	0.9828	98.2

trend stated for the polarization measurements and were in reasonable agreement with those achieved by the hydrogen evolution measurements, but with the dissimilar values related to the various techniques. The extreme inhibition efficacy (93%) was reached at a concentration of 100 ppm of CSPTA-lauric. EIS spectra were analyzed using the equivalent circuit, as illustrated in Fig. 7. This figure reveals a single charge-transfer reaction.

Values of the mean and standard deviation (SD) for  $Z_i$  [ $\text{ohm cm}^2$ ] for the carbon steel electrode at different concentrations of PILs inhibitors in 1 M HCl solution are indicated in S6.†

Fig. 8 and S7† display the Bode and phase angle plots for the PILs inhibitors for carbon steel in 1 M HCl. In order to define the improved phenomena happening at the interfaces, a frequency range for the Bode-phase curve was applied. At high frequencies, the phase angle was utilized to present an overall indication of the inhibition performance. It is well known that the phase angle value of  $-90^\circ$  represents the perfect capacitive action.<sup>66,67</sup>

Fig. 8 illustrates a growth of the phase angle variation as the concentration of the inhibitor is increased and consequently, the phase angle shifts regularly near to the effective capacitive action. In the Bode plot, at lower frequencies, the absolute impedance improved. This confirmed that the developed

surface was protected as the concentration of inhibitor increased, which is correlated to the adsorption effect of the inhibitor on the carbon steel surface.<sup>68</sup>

### 3.4. Gravimetric experiments

The effect of various temperatures on the corrosion rate ( $\text{mg cm}^{-2} \text{h}^{-1}$ ) of the carbon steel with and without the PILs inhibitors at a concentration of 250 ppm is indicated in Table 5 and Fig. 9. The inhibitory efficiency of an IL is calculated as follows:

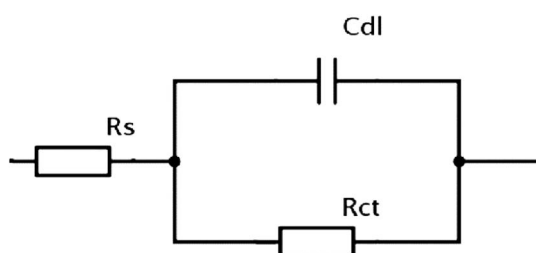
$$\text{CR} = \Delta W/At \quad (6)$$

$$\text{IE \%} = [1 - (\text{CR}/\text{CR}_0)] \times 100 \quad (7)$$

where  $\Delta W$  is the weight loss of carbon steel specimens,  $A$  is the total superficial area,  $t$  is the immersion time, and  $\text{CR}_0$  and  $\text{CR}$  are the corrosion rates of metal samples in the absence and presence of 250 ppm of inhibitors, respectively. The results showed that the percent inhibition was greatly affected by the increase in temperature (318 to 348 K). As can be seen from Table 5, the corrosion rate of carbon steel increased as the temperature increased and the performance of inhibitors became poor gradually when the temperature began to rise. It is clear that CSPTA-lauric had the better inhibitory effect for carbon steel in 1 M HCl solution at elevated temperatures. In addition, the difference in inhibiting ability is strongly affected by the temperature.

### 3.5. Surface morphology investigations

Scanning electron microscopy (SEM) and energy dispersive X-ray (EDX) investigations were done to examine the surface morphology and to detect the elements composition formed on the metal surface before and after the involvement of the inhibitor in the corrosive medium.



**Fig. 7** Equivalent circuit used to fit the impedance data for carbon steel in 0.1 M HCl solution.



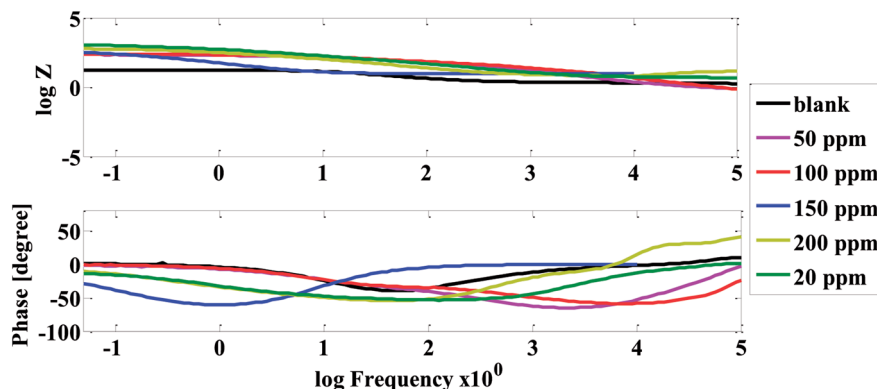


Fig. 8 Bode plots for carbon steel electrode in 1 M HCl solution with and without various concentrations of CSPTA-lauric.

Fig. 10(a) exhibits the EDX bands for elements adsorbed on the carbon steel surface in the absence of inhibitors (blank). Signals of Fe and O ascertain that iron oxide is present, due to metal dissolution (anodic reaction). Also, Fig. 10(a) displays the SEM picture after dropping in 1 M HCl without inhibitor. A coarse severely corroded surface with annihilation was noted on the steel surface.

Upon the addition of 100 ppm of CSPTA-lauric inhibitor, the EDX spectrum in Fig. 10(b) displayed additional signals, confirming the existence of C and N atoms found in the CSPTA-lauric inhibitor. Additionally, the O signal is developed owing to the oxygen atoms present in the inhibitor. Also, the Fe signals are significantly inhibited regarding the samples in Fig. 10(a) due to the creation of a protective inhibitor film. The SEM photo

Table 5 Effect of various temperatures on the corrosion rate ( $\text{mg cm}^{-2} \text{h}^{-1}$ ) of carbon steel with and without PILs inhibitors at a concentration of 250 ppm

Inhibitor	Temperature (T)/K							
	318		328		338		348	
	Corr. rate (CR) ( $\text{mg cm}^{-2} \text{h}^{-1}$ )	IE %	CR	IE %	CR	IE %	CR	IE %
Blank	7.25	—	10.23	—	13.51	—	19.21	—
CSPTA-stearic	1.672	76.9	3.521	65.58	5.75	57.43	9.51	50.49
CSPTA-palmitic	0.723	90.02	1.082	89.42	2.51	81.42	6.130	68.08
CSPTA-myristic	0.451	93.77	0.872	91.47	2.053	84.80	5.752	70.05
CSPTA-lauric	0.253	96.51	0.502	95.09	1.751	87.03	3.271	82.97

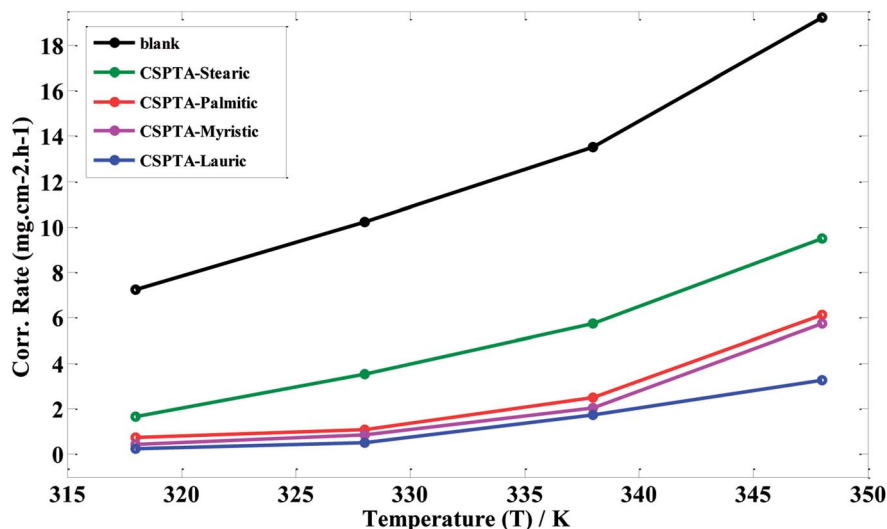


Fig. 9 Effect of various temperatures on the corrosion rate ( $\text{mg cm}^{-2} \text{h}^{-1}$ ) of carbon steel with and without PILs inhibitors at a concentration of 250 ppm.

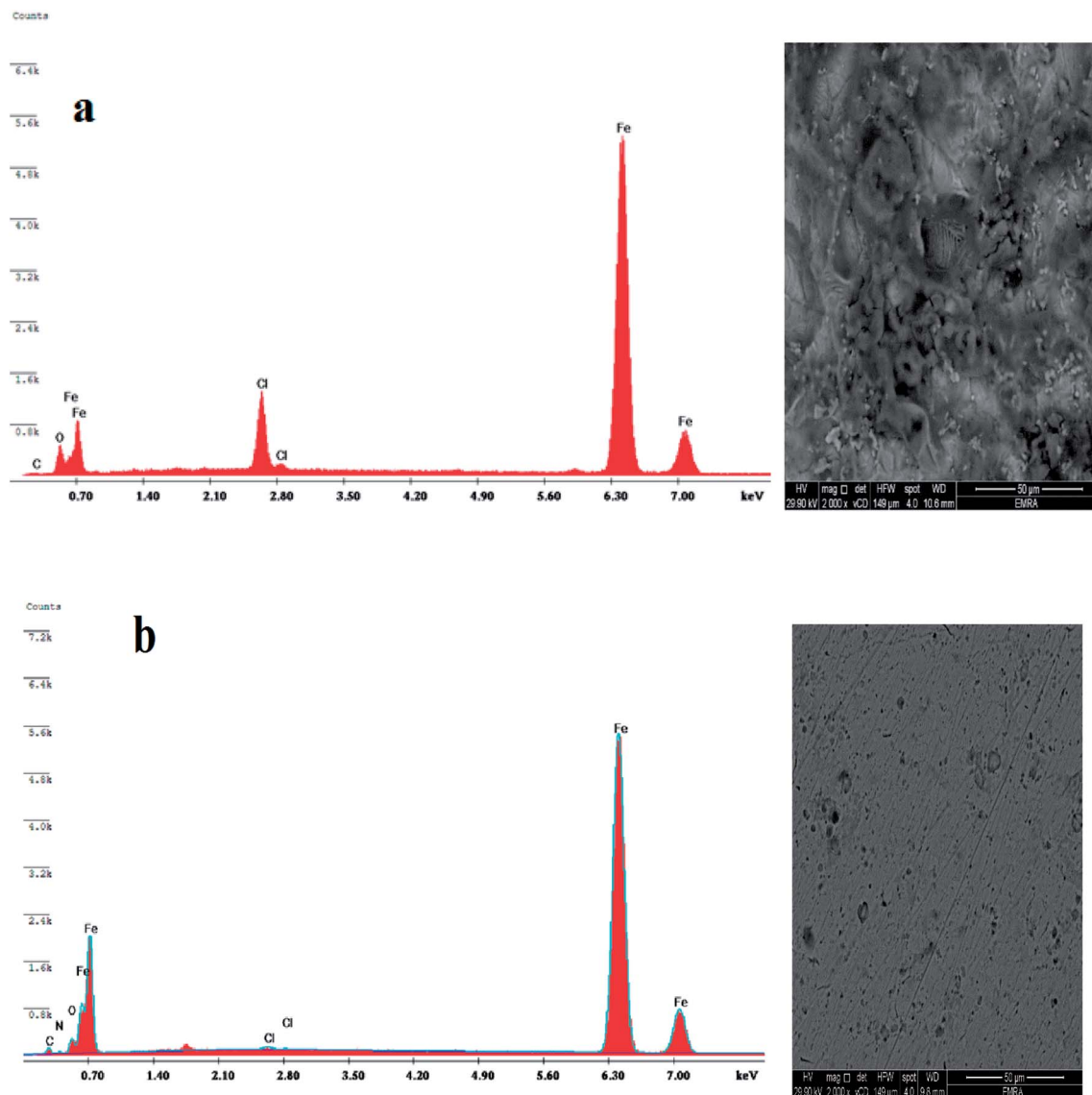


Fig. 10 EDX & SEM for (a) sample after immersion in 1 M HCl without inhibitor (blank) and (b) sample after immersion in 1 M HCl solution containing 100 ppm of CSPTA-lauric inhibitor.

in Fig. 10(b) shows a clear diminishing in the corroded zones caused by adsorbing the molecules inhibiting the carbon steel surface. As a result, a defensive film was produced on the steel surface compared to the sample immersed in acidic medium without an inhibitor.

The SEM and EDX investigations proved the development of a corrosion-inhibitive film on the carbon steel surface, which therefore inhibited the Fe dissolution and hindered the hydrogen evolution resulting from corrosion.

### 3.6. Quantum chemical estimations

The chemical quantum approaches can be used to study the correlation relating to the organic molecular construction and the inhibition effect.<sup>69,70</sup> Tables 6 and 7 show the most essential quantum variables. The results for the molecular frontier orbitals density motions for the PILs molecules are exposed in Fig. 11 and 12. The principle variables are the highest occupied

molecular orbital energy ( $E_{\text{HOMO}}$ ) and the lowest unoccupied molecular orbital energy ( $E_{\text{LUMO}}$ ), which play remarkable roles in detecting the molecules adsorption centers, *i.e.*, the higher the  $E_{\text{HOMO}}$  values are, the easier it is to donate electrons from the inhibitor to the vacant metal d-orbitals. In addition, the capability to gain electrons is governed by the  $E_{\text{LUMO}}$  values, *i.e.*, the lower the  $E_{\text{LUMO}}$  values are, the easier it is to add more negative charge *via* the molecules.

The energy difference ( $\Delta E_{\text{L-H}}$ ) is recognized as the stability key of any inhibitor. So, the higher the values of the inhibitor  $\Delta E$ , the lower its efficiency as the ionization potential will increase and the energy needed for eliminating an electron from the filled outer orbital will increase.<sup>71</sup> Values of low ionization potential represent that the energy needed to eliminate an electron from the occupied exterior orbital is low unlike the values of high dipole moment, which would improve the corrosion inhibition capability.<sup>72</sup> This pointed out that the

**Table 6** Quantum chemical parameters of the investigated compounds

Compound	$E_{\text{HOMO}}$ (eV)	$E_{\text{LUMO}}$ (eV)	$\Delta E$ (eV)	$\mu$ (debye)	$\log P$	$\Delta N$
I	-5.941	0.228	6.169	7.37	7.28	0.671
II	-5.954	0.2146	6.1686	5.91	4.90	0.669
III	-5.945	0.2197	6.1647	5.43	3.04	0.671
IV	-5.985	0.178	6.163	5.21	1.73	0.664

inhibition effectiveness grew in the order: CSPTA-lauric > CSPTA-myristic > CSPTA-palmitic > CSPTA-stearic, which is in complete agreement with the experimental work.

Both the absolute electronegativity ( $X$ ) and global hardness ( $\eta$ ) of the inhibitor molecule are governed by electron affinity ( $A$ ) value and ionization potential ( $I$ ) value in this manner:

$$X = (I + A)/2 \quad (8)$$

$$\eta = (I - A)/2 \quad (9)$$

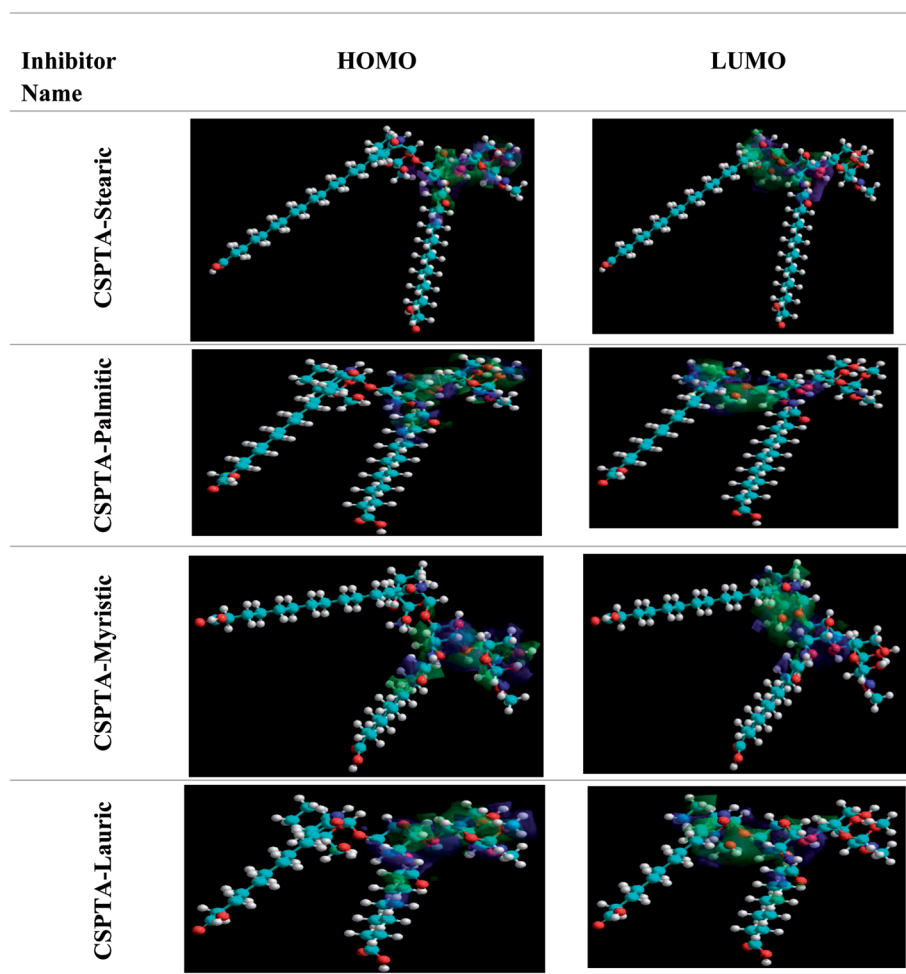
Then  $I$  and  $A$  were evaluated from  $E_{\text{HOMO}}$  to  $E_{\text{LUMO}}$  in this manner:

$$I = -E_{\text{HOMO}} \quad (10)$$

$$A = -E_{\text{LUMO}} \quad (11)$$

**Table 7** Other calculated quantum chemical parameters of the investigated inhibitors

Inhibitor	Ionization potential, $I$ (eV)	Electron affinity, $A$ (eV)	Electronegativity ( $\text{eV mol}^{-1}$ )	Global hardness, ( $\text{eV mol}^{-1}$ )	Softness, $\sigma = 1/\eta_{\text{inh}}$ ( $\text{eV}^{-1}$ )
I	5.941	-0.228	2.8565	3.0845	0.324202
II	5.954	-0.2146	2.8697	3.0843	0.324223
III	5.945	-0.2197	2.86265	3.08235	0.324428
IV	5.985	-0.178	2.9035	3.0815	0.324517

**Fig. 11** Frontier molecule orbital density distributions of the prepared inhibitors.

The number of transferred electrons ( $\Delta N$ ) from the inhibitor molecules to the carbon steel was estimated using the subsequent equation:<sup>73,74</sup>

$$\Delta N = (x_{\text{Fe}} - x_{\text{inh}})/2(\eta_{\text{Fe}} + \eta_{\text{inh}}) \quad (12)$$

where  $x_{\text{Fe}}$  and  $x_{\text{inh}}$  denote the absolute electronegativity of the iron and inhibitor molecule, respectively, and  $\eta_{\text{Fe}}$  and  $\eta_{\text{inh}}$  denote the absolute hardness of the iron and the inhibitor molecule, respectively. Note that the theoretic values of absolute electronegativity of iron ( $\chi_{\text{Fe}}$ ), iron absolute hardness ( $\eta_{\text{Fe}}$ ) and the chemical electronic potential of iron ( $\mu_{\text{Fe}}$ ) are 7, 0, and  $-7$  eV mol<sup>-1</sup>, respectively.

A positive number of electrons transferred ( $\Delta N$ ) as shown in Table 6 proves that the molecules work as electron donors, and the developed  $\Delta N$  indicates an extensive readiness to react with atoms of the metal surface. As stated by Lukovits' study, if the  $\Delta N$  value is less than 3.6, this indicates that the inhibition ability improves with the growing donation capability of electron of the inhibitor at the steel surface.<sup>70</sup>

From these calculations, it is predicted that the inhibitors act as the electron donor, and the iron surface behaves as the electron acceptor. Inhibitors worked at the metal surface steel by adsorbing and consequently developing an inhibitive adsorbed layer that reduces the corrosion. Besides, the

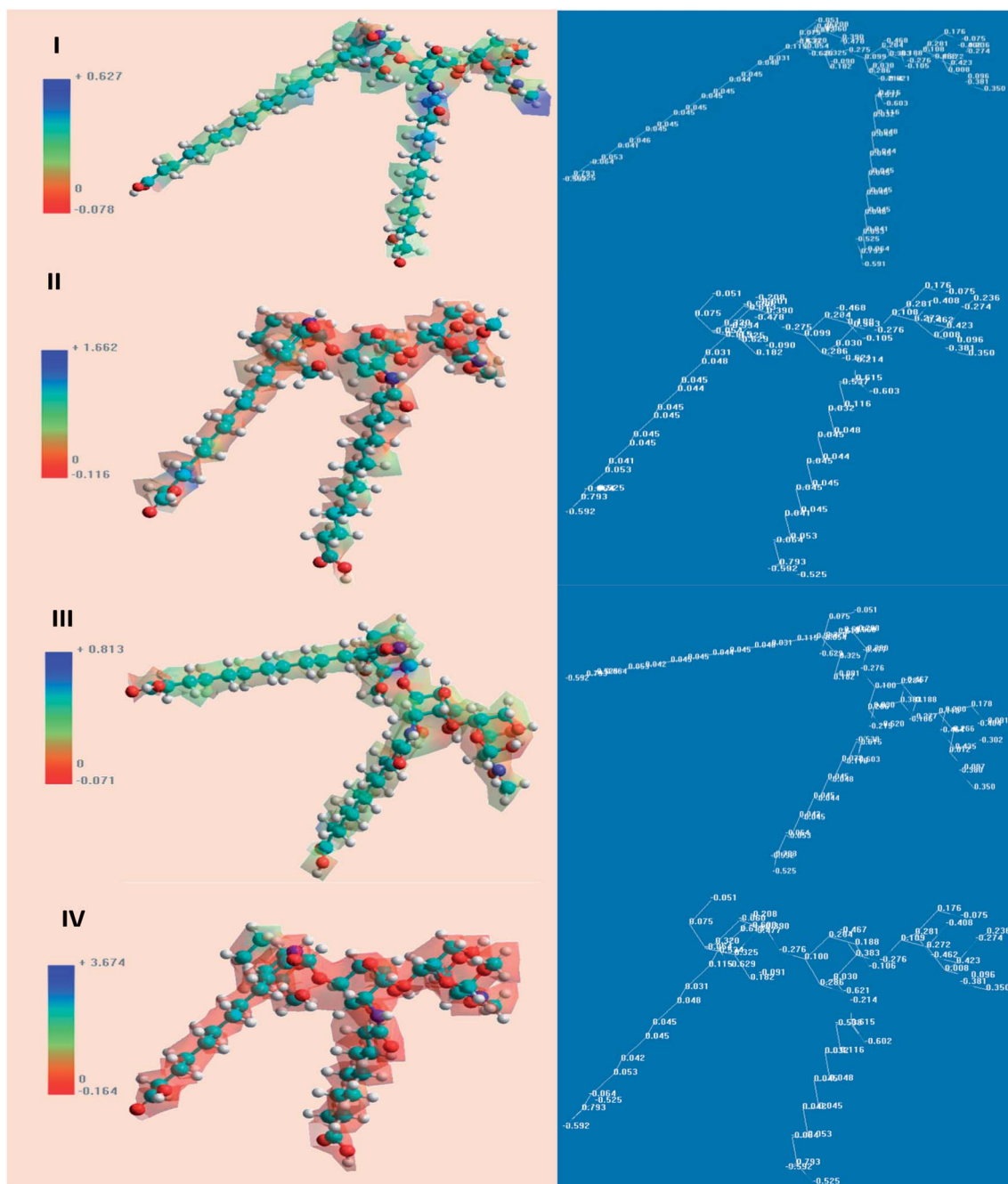


Fig. 12 Molecular electrostatic potential map of: (I) CSPTA-stearic, (II) CSPTA-palmitic, (III) CSPTA-myristic, and (IV) CSPTA-lauric.

produced inhibitors were adsorbed on the iron surface to form an anticorrosion adsorption inhibitive layer.

## 4. Conclusions

(1) Four PILs were prepared based on chitosan linked with four different fatty acids and examined as anticorrosion inhibitors through several techniques.

(2) Corrosion of carbon steel was considered in 1 M HCl, a source for producing hydrogen.

(3) The addition of the prepared PILs reduced the hydrogen generation rate; besides, the rate was further reduced as the concentration of inhibitor increased.

(4) Electrochemical measurements exhibited that the synthesized PILs had enhanced anticorrosion protection for carbon steel surface in acidic corrosive media and that CSPTA-lauric was the most effective.

(5) The percentage inhibition efficiency increased with developing the inhibitors concentration in 1 M HCl, representing a drop in the corrosion rate of carbon steel. On the other hand, the percentage inhibition decreased with the increase in temperature.

(6) The values attained from hydrogen evolution, polarization, impedance, and gravimetric approaches were in a good matching trend.

(7) SEM and EDX analyses demonstrated that the inhibition was due to forming a defensive film of PILs inhibitors on the surface.

(8) CSPTA-lauric inhibitor had the highest inhibitive effectiveness owing to having the lowest ( $\Delta E$ ) energy gap values, and hence, it was the most proficient one for the donation of electrons.

(9) ILs suggest a possible chance for innovatory applications for green chemistry. A great development in this area is expected thanks to the significantly enhanced ecofriendly benefits of these compounds in parallel with traditional inhibitors.

## Conflicts of interest

There are no conflicts to declare.

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## References

- 1 M. A. Deyab, *J. Power Sources*, 2015, **292**, 66–71.
- 2 A. Mohsenzadeh, Y. Al-Wahaibi, R. Al-Hajri and B. Jibril, *J. Pet. Sci. Eng.*, 2015, **133**, 114–122.
- 3 H. Mąka, T. Szychaj and J. Adamus, *RSC Adv.*, 2015, **5**, 82813–82821.
- 4 J. Saien and S. Hashemi, *J. Pet. Sci. Eng.*, 2018, **160**, 363–371.
- 5 B. Haddad, D. Mokhtar, M. Gousse, E. Belarbi, D. Villemin, S. Bresson, M. Rahmouni, N. R. Dhumal, H. J. Kim and J. Kiefer, *J. Mol. Struct.*, 2017, **1134**, 582–590.
- 6 A. S. Hanamertani, R. M. Pilus, A. K. Idris, S. Irawan and I. M. Tan, *J. Pet. Sci. Eng.*, 2018, **162**, 480–490.
- 7 T. Moumene, E. H. Belarbi, B. Haddad, D. Villemin, O. Abbas, B. Khelifa and S. Bresson, *J. Mol. Struct.*, 2015, **1083**, 179–186.
- 8 C. Wang, X. Hu, P. Guan, L. Qian, D. Wu and J. Li, *Int. J. Polym. Anal. Charact.*, 2014, **19**, 70–82.
- 9 S. L. Bacon, R. J. Ross, A. J. Daugulis and J. S. Parent, *Green Chem.*, 2017, **19**, 5203–5213.
- 10 A. M. El-Shamy, K. Zakaria, M. A. Abbas and S. Zein El Abedin, *J. Mol. Liq.*, 2015, **211**, 363–369.
- 11 M. S. Morad, *Corros. Sci.*, 2000, **42**, 1307–1326.
- 12 M. A. Deyab, M. T. Zaky and M. I. Nessim, *J. Mol. Liq.*, 2017, **229**, 396–404.
- 13 R. Bodirlau, C.-A. Teaca and I. Spiridon, *Int. J. Polym. Anal. Charact.*, 2010, **15**, 460–469.
- 14 M. Scendo and J. Uznanska, *Int. J. Corros.*, 2011, **2011**, 1–13.
- 15 D. Yang, M. Zhang, J. Zheng and H. Castaneda, *RSC Adv.*, 2015, **5**, 95160–95170.
- 16 M. Echeverría, C. M. Abreu, F. J. Deive, M. A. Sanromán and A. Rodríguez, *RSC Adv.*, 2014, **4**, 59587–59593.
- 17 Q. Zhang and Y. Hua, *Mater. Chem. Phys.*, 2010, **119**, 57–64.
- 18 K. Khaled, *Appl. Surf. Sci.*, 2004, **230**, 307–318.
- 19 P. Bonhote, A.-P. Dias, N. Papageorgiou, K. Kalyanasundaram and M. Grätzel, *Inorg. Chem.*, 1996, **35**, 1168–1178.
- 20 Q. B. Zhang and Y. X. Hua, *Electrochim. Acta*, 2009, **54**, 1881–1887.
- 21 X. Zhou, H. Yang and F. Wang, *Electrochim. Acta*, 2011, **56**, 4268–4275.
- 22 X. Li, S. Deng and H. Fu, *Corros. Sci.*, 2011, **53**, 1529–1536.
- 23 M. Taghavikish, S. Subianto, N. K. Dutta, L. de Campo, J. P. Mata, C. Rehm and N. R. Choudhury, *ACS Omega*, 2016, **1**, 29–40.
- 24 A. M. Atta, H. A. Al-Lohedan, M. M. S. Abdullah and S. M. ElSaeed, *J. Ind. Eng. Chem.*, 2016, **33**, 122–130.
- 25 E. S. H. El Tamany, S. M. ElSaeed, H. Ashour, E. G. Zaki and H. A. El Nagy, *J. Mol. Struct.*, 2018, **1168**, 106–114.
- 26 A. M. Fekry and R. R. Mohamed, *Electrochim. Acta*, 2010, **55**, 1933–1939.
- 27 G. A. El-Mahdy, A. M. Atta, H. A. Al-Lohedan and A. O. Ezzat, *Int. J. Electrochem. Sci.*, 2015, **10**, 5812–5826.
- 28 P. B. Raja, M. Fadaeinasab, A. K. Qureshi, A. A. Rahim, H. Osman, M. Litaudon and K. Awang, *Ind. Eng. Chem. Res.*, 2013, **52**, 10582–10593.
- 29 M. Sadeghi-Kiakhani and S. Safapour, *J. Ind. Eng. Chem.*, 2016, **33**, 170–177.
- 30 C. Choi, J.-P. Nam and J.-W. Nah, *J. Ind. Eng. Chem.*, 2016, **33**, 1–10.
- 31 Y. An, G. Jiang, Y. Ren, L. Zhang, Y. Qi and Q. Ge, *J. Pet. Sci. Eng.*, 2015, **135**, 253–260.
- 32 M. A. Elgadir, M. S. Uddin, S. Ferdosh, A. Adam, A. J. K. Chowdhury and M. Z. I. Sarker, *J. Food Drug Anal.*, 2015, **23**, 619–629.
- 33 N. E. Suyatma, A. Copinet, E. Legin-Copinet, F. Fricoteaux and V. Coma, *J. Polym. Environ.*, 2011, **19**, 166–171.

- 34 S. Pokhrel, P. N. Yadav and R. Adhikari, *Nepal Journal of Science and Technology*, 2016, **16**, 99–104.
- 35 S. A. Umoren and U. M. Eduok, *Carbohydr. Polym.*, 2016, **140**, 314–341.
- 36 R. A. A. Muzzarelli, M. Guerrieri, G. Goteri, C. Muzzarelli, T. Armeni, R. Ghiselli and M. Cornelissen, *Biomaterials*, 2005, **26**, 5844–5854.
- 37 I. Aranaz, R. Harris and A. Heras, *Curr. Org. Chem.*, 2010, **14**, 308–330.
- 38 J. H. Hamman, *Mar. Drugs*, 2010, **8**, 1305–1322.
- 39 S. K. Shukla, A. K. Mishra, O. A. Arotiba and B. B. Mamba, *Int. J. Biol. Macromol.*, 2013, **59**, 46–58.
- 40 R. A. A. Muzzarelli and C. Muzzarelli, in *Polysaccharides I*, ed. T. Heinze, Springer-Verlag, Berlin/Heidelberg, 2005, vol. 186, pp. 151–209.
- 41 S. Mima, M. Miya, R. Iwamoto and S. Yoshikawa, *J. Appl. Polym. Sci.*, 1983, **28**, 1909–1917.
- 42 H. Zhang, S. Wu, Y. Tao, L. Zang and Z. Su, *J. Nanomater.*, 2010, **2010**, 1–5.
- 43 A. Chinnappan, A. H. Jadhav, J. M. C. Puguán, R. Appiah-Ntiamoah and H. Kim, *Energy*, 2015, **79**, 482–488.
- 44 I. B. Obot, S. A. Umoren and N. O. Obi-Egbedi, *J. Mater. Environ. Sci.*, 2011, **2**, 60–71.
- 45 D. de Britto, L. A. Forato and O. B. G. Assis, *Carbohydr. Polym.*, 2008, **74**, 86–91.
- 46 X. Fei Liu, Y. Lin Guan, D. Zhi Yang, Z. Li and K. De Yao, *J. Appl. Polym. Sci.*, 2001, **79**, 1324–1335.
- 47 Y.-I. Jeong, D.-G. Kim, M.-K. Jang and J.-W. Nah, *Carbohydr. Res.*, 2008, **343**, 282–289.
- 48 W. Huang and J. Zhao, *Colloids Surf., A*, 2006, **278**, 246–251.
- 49 C. Cao, *Corros. Sci.*, 1996, **38**, 2073–2082.
- 50 M. A. Migahed, M. M. EL-Rabiei, H. Nady, A. Elgendy, E. G. Zaki, M. I. Abdou and E. S. Noamy, *Journal of Bio-and Tribo-Corrosion*, 2017, **3**, 31.
- 51 M. A. Migahed, A. Elgendy, M. M. EL-Rabiei, H. Nady and E. G. Zaki, *J. Mol. Struct.*, 2018, **1159**, 10–22.
- 52 B. Lin, R. Hu, C. Ye, Y. Li and C. Lin, *Electrochim. Acta*, 2010, **55**, 6542–6545.
- 53 X.-H. Li, S.-D. Deng, H. Fu and G.-N. Mu, *J. Appl. Electrochem.*, 2009, **39**, 1125–1135.
- 54 M. Liang, H. Zhou, Q. Huang, S. Hu and W. Li, *J. Appl. Electrochem.*, 2011, **41**, 991–997.
- 55 X. Li, S. Deng, G. Mu, H. Fu and F. Yang, *Corros. Sci.*, 2008, **50**, 420–430.
- 56 V. M. Abbasov, H. M. Abd El-Lateef, L. I. Aliyeva, E. E. Qasimov, I. T. Ismayilov and M. M. Khalaf, *Egypt. J. Pet.*, 2013, **22**, 451–470.
- 57 Q. Qu, L. Li, W. Bai, S. Jiang and Z. Ding, *Corros. Sci.*, 2009, **51**, 2423–2428.
- 58 F. G. Liu, M. Du, J. Zhang and M. Qiu, *Corros. Sci.*, 2009, **51**, 102–109.
- 59 K. Ramya, K. K. Anupama, K. M. Shainy and A. Joseph, *Egypt. J. Pet.*, 2017, **26**, 421–437.
- 60 D. K. Yadav, B. Maiti and M. A. Quraishi, *Corros. Sci.*, 2010, **52**, 3586–3598.
- 61 M. Lebrini, F. Bentiss, N.-E. Chihib, C. Jama, J. P. Hornez and M. Lagrenée, *Corros. Sci.*, 2008, **50**, 2914–2918.
- 62 R. Solmaz, *Corros. Sci.*, 2014, **81**, 75–84.
- 63 S. Ramesh and S. Rajeswari, *Electrochim. Acta*, 2004, **49**, 811–820.
- 64 M. Bouklah, B. Hammouti, M. Benkaddour and T. Benhadda, *J. Appl. Electrochem.*, 2005, **35**, 1095–1101.
- 65 A. A. Abd-Elaal, N. M. Elbasiony, S. M. Shaban and E. G. Zaki, *J. Mol. Liq.*, 2018, **249**, 304–317.
- 66 M. A. Migahed, E. G. Zaki and M. M. Shaban, *RSC Adv.*, 2016, **6**, 71384–71396.
- 67 M. A. Hegazy, A. M. Hasan, M. M. Emara, M. F. Bakr and A. H. Youssef, *Corros. Sci.*, 2012, **65**, 67–76.
- 68 M. A. Migahed, M. M. EL-Rabiei, H. Nady, A. Elgendy, E. G. Zaki, M. I. Abdou and E. S. Noamy, *Journal of Bio-and Tribo-Corrosion*, 2017, **3**, 31.
- 69 D. F. V. Lewis, C. Ioannides and D. V. Parke, *Xenobiotica*, 1994, **24**, 401–408.
- 70 M. A. Migahed, A. M. Al-Sabagh, E. A. Khamis and E. G. Zaki, *J. Mol. Liq.*, 2015, **212**, 360–371.
- 71 L. Saqalli, M. Galai, F. Benhiba, N. Gharda, N. Habbadi, R. Ghailane and M. Ebn, *J. Mater. Environ. Sci.*, 2017, **8**(7), 2455–2467.
- 72 L. Adardour, H. Lgaz, R. Salghi, M. Larouj, S. Jodeh, M. Zougagh, O. Hamed and M. Taleb, *Pharm. Lett.*, 2016, **8**, 173–185.
- 73 I. Lukovits, E. Kálmán and F. Zucchi, *Corrosion*, 2001, **57**, 3–8.
- 74 E. S. H. El Ashry, A. El Nemr and S. Ragab, *J. Mol. Model.*, 2012, **18**, 1173–1188.