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SOME WATER-SOLUBLE POLYMERS AS CORROSION INHIBITORS FOR CARBON STEEL IN ACIDIC MEDIUM

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ABSTRACT

Encouraged by the excellent performance of organic polymers as scale inhibitors and anticorrosive coatings, we reasoned that polymers with properly designed structures might also be used as efficient water-soluble corrosion inhibitors. New kind of water soluble hyperbranched polyamide ester (HPAE) synthesized using trimethylol propane as a precursor via Michael addition reaction with diethanolamine. The chemical structure of the prepared polymers was determined by FT-IR. The inhibitive effect of HPAE on the corrosion of steel in 1 M HCl solution has been investigated using potentiodynamic polarization and electrochemical impedance spectroscopy (EIS) techniques. The results indicated that HPAE inhibited the corrosion of steel and the extent of inhibition increased with HPAE concentrations. Potentiodynamic polarization data suggests mixed-mode of corrosion inhibition.

Keywords : Polarization, EIS, Electrochemical, Steel, Corrosion, Acid inhibition

1. INTRODUCTION

Corrosion is gradual destruction of a material because of its reaction with environment. It is a major industrial problem that has attracted a lot of investigators in recent years [1,2]. In the fields of petroleum and oil industry, corrosion represents a huge percentage of the annual economic losses. The most significant corrosion types in petroleum industry are local and general corrosion. Another problem in the operation of pipelines is internal corrosion. The corrosion problems are always associated with the equipment maintenance and operation. This lead to stopping the process partially or even completely, resulting in economic losses ranging between 10% and 30% of the whole maintenance budget. Indeed, corrosion control is an essential issue from application point of view and it has been reported that inhibitors are needed to be used which act as a barrier to reduce the aggressiveness of the environments against the corrosion attack [3-5]. Most common type of corrosion inhibitors are organic compounds. Their inhibition property is

dependent upon their functional group, which adsorb on the metal surface. Most of the efficient organic compounds acts as inhibitors have oxygen, sulphur, nitrogen atoms and multiple bonds through which they adsorb on metal surface [6-8]. The inhibition efficiency of inhibitors increases in the order of: O < N < S < P [9-11].

The use of polymers as corrosion inhibitors has attracted considerable attention recently [12]. Polymers are used as corrosion inhibitors because, through their functional groups they form complexes with metal ions and on the metal surface these complexes occupy a large surface area [13,14], thereby blanketing the surface and protecting the metal from corrosive agents present in the solution [15-18]. The inhibitive power of these polymers is related structurally to the cyclic rings, heteroatom (oxygen and nitrogen) that are the major active centers of adsorption [19,20]. Because of the unique characteristics of hyperbranched polymers which have become widely used, they have attracted more attention especially in scientific research and industrial fields such as

oil industry, corrosion inhibitor, and coating of pipelines [21-23]. The hyperbranched polymers have many advantages, the most significant of which is that it easy to prepare, high solubility in addition low viscosity [24].

In this paper, we have to synthesize a hyperbranched poly amide-ester via Michael addition reaction. These polymers were then investigated for their ability to prevent corrosion of the steel in aggressive aqueous acidic medium.

2. EXPERIMENTAL

2.1. Materials

2.1.1. Chemicals

Diethanolamine (DEA), trimethylol propane (TMP), maleic anhydride(MA), methanol, p-toluene sulfonic acid (p-TSA), N,N-Dimethylacetamide (DMAC), ethyl ether, acetone.

2.1.2 Solutions

a-Hydrochloric acid solution

1 M HCl was used as a corrosive solution, and was prepared by diluting concentrated HCl (37%) to a required concentration using bidistilled water.

b- Inhibitor solutions

100 ml stock solutions of hyperbranched polymers were prepared by dissolving the desired weight of each compound in 100ml of absolute ethanol, and then the six different concentrations 50, 100, 200, 400, 500 and 1000ppm were prepared by dilution with known concentration of hydrochloric acid.

2.1.3. Carbon steel electrode

Corrosion tests were performed on carbon steel electrode having the following chemical composition (wt.%): 0.09% C, 0.09% Si, 0.46% Mn, 0.03% P and the remainder Fe.

2.2. Synthesis techniques

The synthetic procedure was described in the literature reported by CHEN Q. H. et al [25]

2. 2.1. Synthesis of AB₂ monomer

AB₂ monomer was prepared by amidation reaction of DEA and maleic anhydride at a molar ratio of 1/1. 0.1 mol (9.8060 g) maleic anhydride was dissolved with 20.0 g DMAC in a three-necked round-bottomed flask at 25°C constant temperature. 0.1 mol (10.5140 g) crystal DEA was heated to be liquid state and poured into the maleic anhydride solution rapidly, and then 10.0 g more DMAC was added to assure a proper viscosity of mixture for reaction. The reaction mixture was stirred with a magnetic stirrer at 25 °C for 2 hrs to obtain the DMAC solution of AB₂ monomer.

2. 2.2. Synthesis of hydroxyl terminated fourth generation hyperbranched poly amide-ester [4G-(OH)]

Hyperbranched poly amide-esters were synthesized through “one-step process” or “pseudo one-step process”. TMP was added as a center core. In 100 mL three-necked round-bottomed flasks, calculated amounts of P- TSA and the obtained AB₂ monomer solution were introduced according to various formulae, respectively. A neck mouth of this flask was ventilated a vacuum pump and installed a reflux distillation equipment with water condenser. The flask was put into an oil bath with constant temperature of 120°C. The mixture was stirred with a magnetic stirrer for 6 hrs under vacuum condition. The crude products were dissolved in 30.0g DMAC, and then precipitated and washed by the mixture solvents of ethyl ether- acetone (vol. ratio is 1/1). The process was carried out for several times before the precipitate was dried in vacuo. The synthesized inhibitors were illustrated in Scheme 1, and the chemical structure was confirmed by FT-IR spectroscopic analysis using Nicolet iS10 FT-IRspectrometer, Thermo Fisher Scientific (USA).

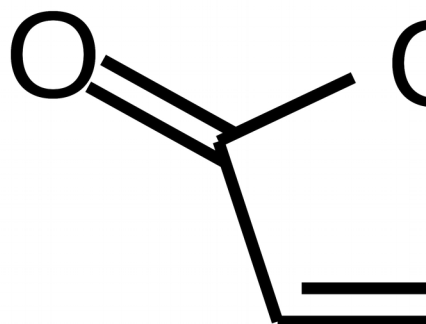
2.3- Method used for Corrosion Measurements

Electrochemical techniques

Electrochemical measurements were performed with a traditional three-electrodes cell using Volta lab 40 (Tacussel-RadiometerPGZ301) potentiostat and controlled by Tacussel corrosion analysis software model (Voltmaster 4) at under static condition. All experiments were conducted in a

electrode, saturated calomel electrode (SCE) as reference electrode and carbon steel as working electrode.

A- Potentiodynamic polarization curves were obtained by varying the potential automatically from -800 to -300 mV in relation to a steady-state open circuit potential (EOCP) with the scan rate 2 mV s^{-1} . The polarization curves were obtained after 1h in the open-circuit potential.



maleic

Scheme 1 Synthesis of hyperbranched poly amide-ester

conventional three-electrodes glass cell assembly with a platinum wire as auxiliary

B- Electrochemical impedance spectroscopy (EIS) measurements were carried out using a

potentiostat (VoltaLab PGZ-301) attached to the Zsimpwin software program. The measurements were carried out using AC signal (10 mV) peak to peak at OCP in the frequency range of 100 kHz–50 mHz. In all experiments, the carbon steel electrode was also allowed to reach its stable open-circuit potential (OCP), which occurred after 1h. EIS diagrams are given in the Nyquist representation.

2.4. Scanning Electron Microscope (SEM) Analysis

The surface films were formed on the carbon steel specimens by immersing them in 1M HCl solution with and without different concentrations of inhibitors for a period of time of 12hrs. After the immersion period, the specimens were taken out, dried and the nature of the film formed on the surface of the metal specimen was analyzed by SEM techniques to examine the surface morphology. JEOL JSM-5500 scanning electron microscope was used for this investigation.

3. RESULTS AND DISCUSSION

3.1. Chemical structure confirmation of the synthesized Inhibitors

The structural characteristics of the synthesized hyperbranched poly (amide-ester)

by FT-IR spectroscopy in the range 4000–500 cm^{-1} .

AB₂ monomer

Fig. 1 illustrates the FTIR spectra of AB₂ monomer. It shows that in the AB₂ monomer spectrum, strong absorption of tertiary amine and hydroxyl group stretching vibration occur at about 1614 and 3387 cm^{-1} , respectively. Band at 1714 cm^{-1} is ascribable to isolated aromatic carboxylic acid carbonyl group [26]. The results show that the reaction between diethanolamine and maleic anhydride is amidation but not esterification under experimental conditions and the prepared AB₂ monomer is an amine compound containing hydroxyl groups. The structure of AB₂ monomer is comparable with the theoretic one. And so, the synthesis process is sound.

Structural description for structural description for hyperbranched poly amide-ester generation (G4)

Fig. 2 reveals the FTIR spectra of hyperbranched poly amide-ester generation 4 (G4), the spectrum of (G4) possesses strong absorption bands at 1726, 1657, 1176 and 1054 cm^{-1} for tertiary amide-ester group, indicating that the polymerization reaction has been

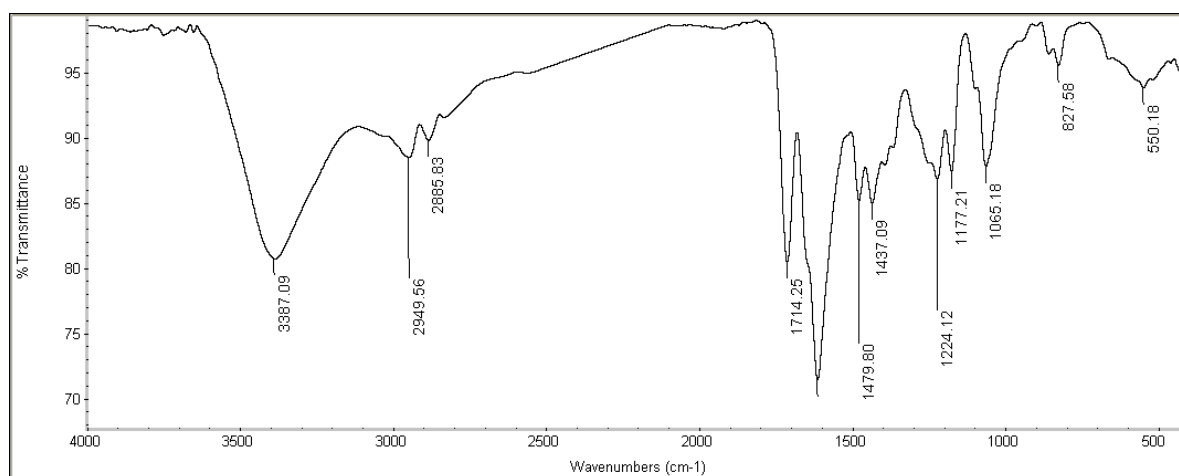


Fig. 1 FT-IR spectra of AB₂ monomer

generation 3 and 4 G3 & G4. were confirmed

carried out among the prepared AB₂ monomers

under the proposed condensation condition, and the (G4) product is confirmed to be an amide-ester compound.

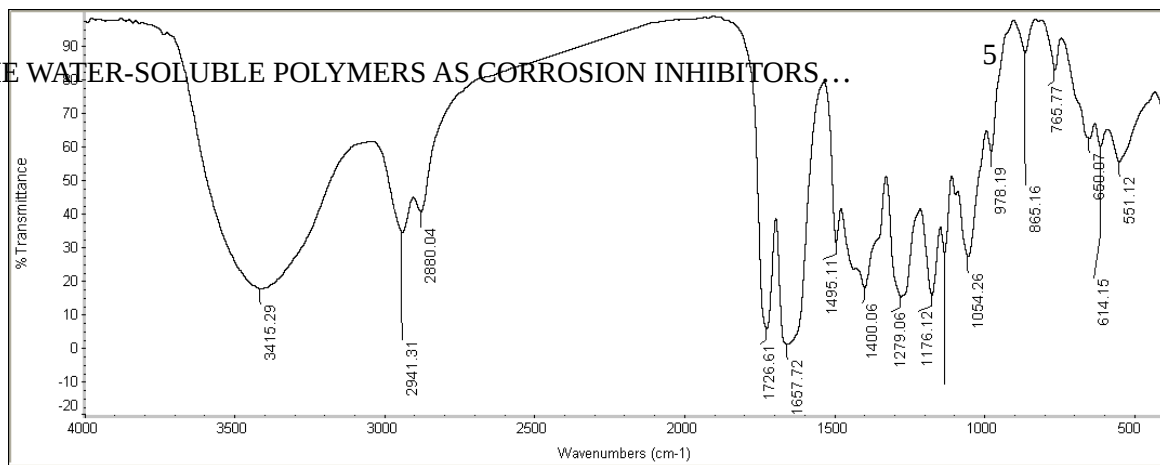


Fig. 2 FT-IR spectra of hyperbranched poly amide-ester generation 4 (G4)

The absorption band at 3415 cm^{-1} range is ascribable to the stretching vibration of hydroxyl group, showing that the hydroxyl groups of AB_2 monomers have not completely reacted with the carboxyl groups on them or their oligomers. The results indicate that AB_2 monomer can polycondense by self-catalysis [27], and trimethylol propane serving as a “center core” leads to few changes in the structures of hyperbranched macromolecules

3.2. Study the protection efficiency of C-Steel electrode by the electrochemical technique

3.2.1. Potentiodynamic polarization

The electrochemical polarization measurements of carbon steel in 1 M HCl without and with various concentrations (50,100,200,400,500 and 1000ppm) of HPAE generation 3 and 4 (G3 & G4) are shown in Fig. 3. The values of electrochemical parameters such as corrosion potential (E_{corr}), cathodic and anodic Tafel slopes (β_a and β_c) and corrosion current density (I_{corr}) were extracted by Tafel extrapolating the anodic and cathodic lines and are listed in Table 1. The percentage of inhibition efficiency (IE %) was calculated using the following equation [28]:

$$\text{IE}\% = \frac{I_0 - I}{I_0} \times 100 \quad (1)$$

where I_0 and I are the corrosion current densities in the absence and presence of inhibitors respectively.

It was observed that the polarization resistance (R_p) and inhibition efficiency (IE%) increase as the concentration of HPAE increase. The highest inhibition efficiency was obtained at 1000 ppm concentration. This also indicates that more inhibitor particles were adsorbed on the surface of the metal thus providing a wider surface coverage and the extract acts as an adsorption inhibitor [29]. It can be observed from Table 1 that the corrosion current density (I_{corr}) decreases with increases in the concentration of HPAE which is due to the increase in the blocked fraction of the metal surface by adsorption [30]. The shift in the Tafel slopes of both the cathodic reaction (β_c) and anodic reaction (β_a) as shown in Fig. 3 in the presence and absence of HPAE suggest that the inhibitor affects both the cathodic and anodic reactions. This also suggests that HPAE is a mixed type of inhibitor. It can also be observed from Fig. 4 that as the concentration of the HPAE increases, there appears to be a shift towards more negative potential indicating that the HPAE acts as a good corrosion inhibitor. It can be observed that as the corrosion current density (I_{corr}) decreases, the inhibition efficiency (IE)% increases as the concentration of the HPAE is increased. This suggests that the HPAE retards the reactions at both the cathodic and the anodic sites as a result of coverage of these sites by the molecules of the HPAE [31]. The shift in β_c and β_a values as shown in Table 1 indicates that

adsorption of HPAE modifies the mechanism of anodic dissolution and the cathodic hydrogen evolution, this reveals that the inhibition mechanism occurred by simply blocking the available cathodic and anodic sites of the metal surface [32]. The inhibition efficiency obtained from polarization measurements is found to follow the order: G4 > G3, indicating the more beneficial effect of compound G4 on corrosion inhibition of carbon steel.

3.2.2. Electrochemical impedance spectroscopic measurements (EIS)

EIS measurements provide a better understanding of the corrosion mechanism taking place at the electrode surface including the kinetics of the electrode processes and simultaneously about the surface properties of the investigated systems. The method is widely used for investigation of the corrosion inhibition processes [33]. The Nyquist plots of carbon steel in 1 M HCl solution in the absence

Table 1. Electrochemical polarization parameters for the corrosion of carbon steel in 1 M HCl containing various concentrations of G3 and G4.

Inhibitor	Conc. (ppm)	R_p (Ωcm^2)	$E_{\text{corr.}}$ (mv)	I_{corr} (mAcm^{-2})	β_a (mv dec^{-1})	β_c (mv dec^{-1})	IE%	θ
G3	Blank	69.09	-522.6	1.0405	199.9	-188.4	-	-
	50	153.6	-510.4	0.2394	144.9	-127.9	76.99	0.769
	100	166.62	-499.6	0.1904	88.6	-103.1	81.70	0.817
	200	171.06	-580.6	0.1772	118.4	-110.0	82.96	0.829
	400	346.91	-523.1	0.1477	154.3	-137.7	85.80	0.858
	500	478.37	-525.1	0.1203	151.0	-144.6	88.43	0.884
	1000	487.13	-523.4	0.0998	129.2	-137.5	90.40	0.904
G4	50	240.50	-525.1	0.1997	141.5	-144.4	80.80	0.808
	100	268.18	-524.0	0.1756	126.6	-140.6	83.12	0.831
	200	344.83	-526.7	0.1455	143.0	-141.8	86.01	0.860
	400	441.34	-529.1	0.1257	144.5	-141.5	87.91	0.879
	500	398.39	-536.8	0.1068	117.4	-134.6	89.73	0.897
	1000	611.03	-540.7	0.0714	122.2	-132.7	93.13	0.931

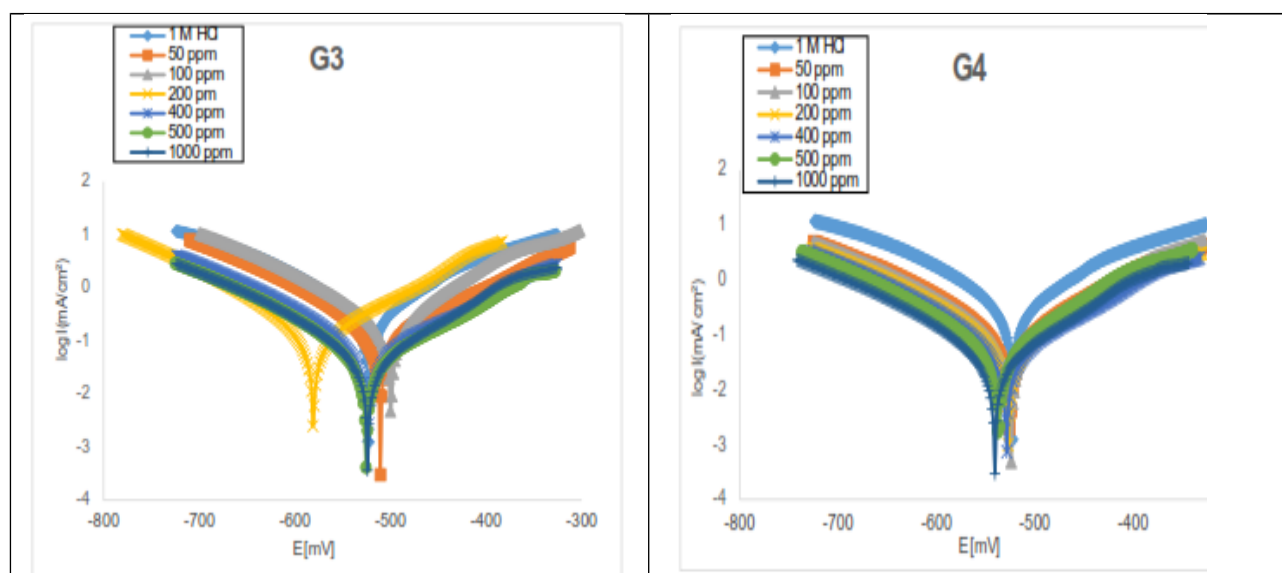


Fig. 3. Potentiodynamic polarization curves for carbon steel in 1 M HCl without and with various concentrations of G3 and G4.

and presence of different concentrations of the synthesized inhibitors given in Fig. 4. It is clear from the figure that all impedance spectra have a depressed semicircular shape in the complex impedance plane, with the center under the real axis. Deviations from a perfectly circular shape indicate the frequency dispersion of the interfacial impedance [34].

The values of percentage inhibition efficiency were calculated from the values of charge transfer resistance (R_{ct}) according to the following equation [35]:

$$IE\% = \frac{R_{ct} - R_{ct}^0}{R_{ct}} \times 100 \quad (2)$$

where R_{ct} and R_{ct}^0 are the charge transfer resistances with and without the inhibitors, respectively. Inhibition efficiencies and other calculated impedance parameters obtained from fitted spectra are given in Table 2. As seen from Table 2, the charge transfer resistance (R_{ct}) values of inhibited substrates increase with the concentration of inhibitors. The most pronounced effect and the highest R_{ct} are obtained by inhibitor (G4). A large R_{ct} values are generally associated with a slower corroding system as the R_{ct} value is a measure

inversely proportional to the corrosion rate [36]. On the other hand, the values of C_{dl} decrease with an increase in inhibitor concentration. This situation was the result of an increase in the surface coverage by these inhibitors which led to an increase in the inhibition efficiency. Also, this effect leads to a protective film, inhibiting species or both on carbon steel surface.

This decrease in the C_{dl} , which can result from a decrease in local dielectric constant and/or increase in thickness of the electrical double layer, suggested that the compounds act via adsorption at the metal/solution interface [37]. It could be assumed that the decrease of C_{dl} values is caused by the gradual replacement of water molecules by adsorption of organic molecules on the electrode surface, which decreases the extent of the metal dissolution [38]. The adsorption can occur either directly on the basis of donor-acceptor interaction between the unshared electron pairs and/or p-electrons of inhibitor molecule and the vacant d-orbitals of the metal surface or by interaction of the inhibitors with already adsorbed chloride ions [39]. These results confirm that all of the synthesized inhibitors exhibit good inhibiting

Table 2. Electrochemical impedance parameters and inhibition efficiency for the corrosion of carbon steel in 1 M HCl containing various concentrations of G3 and G4.

Inhibitor	Conc.(ppm)	R_s ohm.cm ²	R_{ct} ohm.cm ²	C_{dl} μF/cm	(%)θ
G ₃	Blank	3.299	60.92	653.0	-
	50	3.732	247.5	179.9	75.38
	100	4.009	287.9	87.34	78.83
	200	5.704	333.6	84.90	81.73
	400	4.227	390.7	64.34	84.40
	500	4.443	467.3	51.02	86.96
G ₄	1000	9.554	555.1	38.14	89.02
	50	6.011	311.2	64.68	80.42
	100	6.613	352.1	51.48	82.69
	200	7.473	389.4	51.13	84.35
	400	6.143	476.6	46.74	87.21
	500	10.21	558.6	45.19	89.09
	1000	14.04	688.9	31.90	91.15

of electron transfer across the surface and is

performance of carbon steel in HCl solution,

and the more efficient to inhibit the corrosion of carbon steel is compound (G4). It is worth noting that the inhibition efficiencies calculated from EIS measurements are in good agreement with that obtained from potentiodynamic polarization measurements.

3.3 Scanning Electron Microscopy (SEM) Analysis.

In order to verify if the HBPA in fact adsorbed on the steel surface, SEM analysis was carried out. The SEM photograph for the polished steel surface before immersion in 1M HCl solution is shown in Fig. 5. The photograph shows the surface was smooth, clear and without pits. A photograph of the steel surface after immersion in 1 M HCl solution is shown in Fig. 6. The photograph revealed that, the surface was strongly damaged in the

absence of HBPA. A photograph of the steel surface after immersion in 1 M HCl solution containing 1000 ppm of HBPA is shown in Fig. 7. It was found that the faceting observed in figures disappeared and the surface was free from pits and it was smooth. It can be concluded from Figures (5-7) that corrosion decreased largely in the presence of HBPA hence corrosion was inhibited strongly when HBPA was present in the medium. In the presence of 1000 ppm of HBPA, there is much less damage on the steel surface, which further confirm the inhibition action. Also, there is an film adsorbed on mild steel surface.

In accordance, it might be concluded that the adsorption film can efficiently inhibits the corrosion of steel.

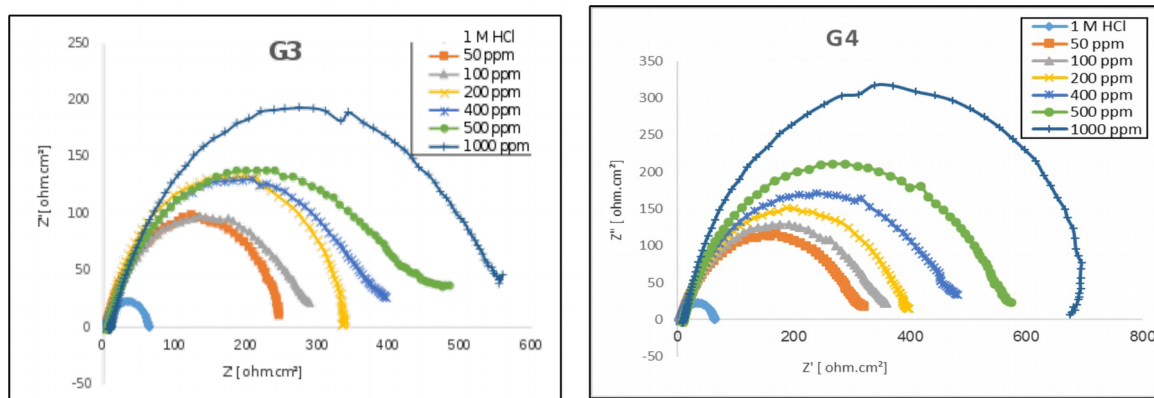


Fig. 4. Nyquist plots for carbon steel in 1 M HCl without and with various concentrations of G3 and G4.

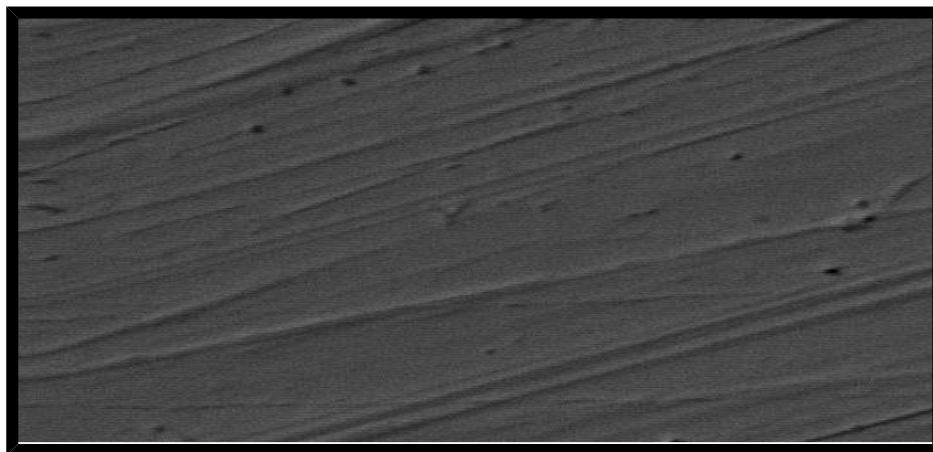


Fig. 5: SEM micrographs of steel surface before immersion in 1 M HCl

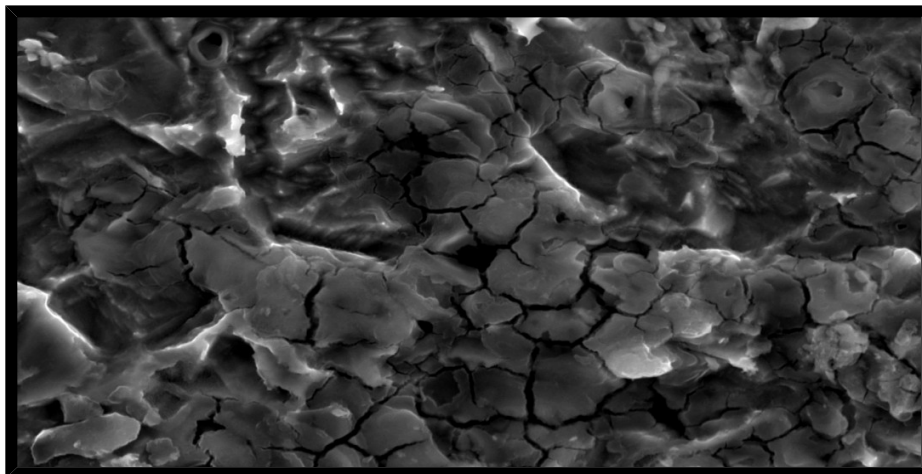


Fig. 6: SEM micrographs of steel surface after 12 hrs of immersion in 1 M HCl

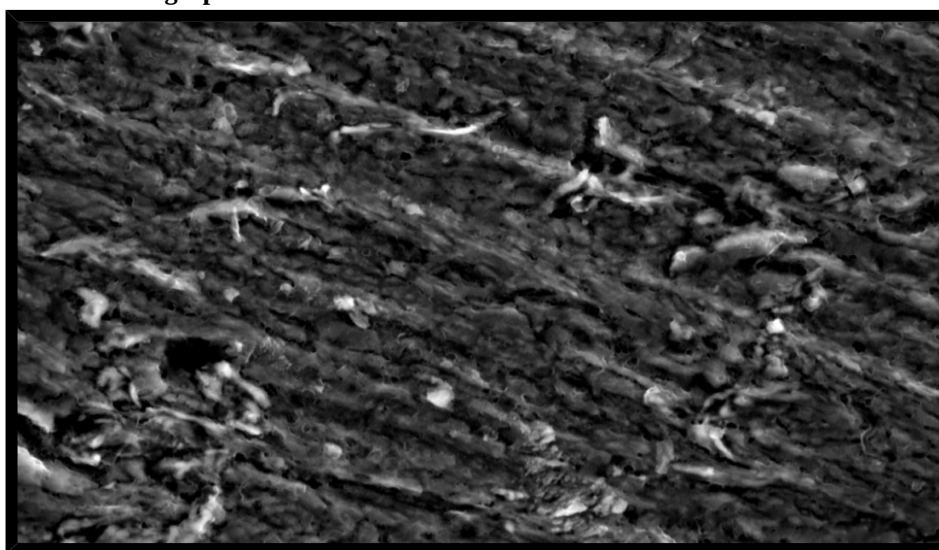


Fig. 7: SEM micrographs of steel surface after 12 hrs of immersion in 1 M HCl +1000 ppm of G4 HBPA

CONCLUSIONS

(1) The results showed that inhibitors HPAE generation 3 and 4 (G3 & G4) have excellent inhibition efficiency for the corrosion of carbon steel in 1 M HCl. The inhibition efficiency increases in the following order $G4 > G3$.

(2) The potentiodynamic polarization curves indicated HPAE generation 3 and 4 (G3 & G4) that act as a mixed-type of inhibitors by inhibiting both anodic metal dissolution and cathodic hydrogen evolution reactions.

(3) The impedance results indicate that the value of charge transfer resistance increased and double layer capacitance decreased. This result can be attributed to the increase of thickness of electrical double layer

(4) The results of different techniques (potentiodynamic polarization, (Tafel) and electrochemical impedance spectroscopy (EIS) showed a good agreement with each other. This agreement among the different independent techniques proves the validity of these results.

REFERENCES

1. Q.Liu, Z.Y. Zhu , W.Ke, C.I.Han and C.L. Zeng, *Corros. Nace.* 57(2001) 730.
2. D.Collins, R.E.Weyers and I.L. Al-Qadi, *Corros. Nace.* 49 (1993) 74.
3. F.Eghbali, M.H.Moayed, A.Davoodi and N.Ebrahimi, *Corros. Sci.* 53 (2011) 513.
4. A.S Fouda and A.S Ellithy., *Corros. Sci.* 51 (2009) 868.
5. S. Andreani, M. Znini, J. Paolini, L. Majidi, B. Hammouti, J. Costa¹ and A. Muselli¹, *J. Mater. Environ. Sci.* 1 (2016) 187.
6. M.A. Quraishi, I. Ahamad, A.K. Singh, S. K. Shukla, B. Lal and V. Singh, *Mater. Chem. Phys.* 112 (2008) 1035.
7. S. K. Shukla and M.A. Quraishi, *J. Appl. Electrochem.* 39 (2009) 1517.
8. S.K. Shukla ,M.A. Quraishi and R. Prakash, *Corros. Sci.* 50 (2008) 2867.
9. S.A. Ali, M.T. Saeed and S.U. Rahman, *Corros. Sci.* 45 (2003) 253.
10. M. Pardave, M. Romero, H. Hernandez, M. Quijano, N. Likhanova, J. Uruchurtu and J. Garcia, *Corros. Sci.* 54 (2012) 231.
11. A. Chetouani, B. Hammouti, T. Benhadda and M. Daoudi, *Appl. Surf. Sci.* 249 (2005) 375.
12. S.Rajendran, S.P. Sridevi, N. Anthony, A. John and M. Sundearavadivelu, *Anti-Corros. Methods Mater* 2 (2005) 52(2) 102.
13. G.A. Bereket, A. Yurt and H.Turk , *Anti-Corros. Methods Mater* 6 (2003) 422.
14. M.Daouadji and N.Chelali, *J Appl Polym Sci* 90 (2004) 1275.
15. S.Selvaraji, A.Kennedy, A. Amalraj, S .Rajendran and N.Palaniswamy, *Corros. Rev* 22 (2004) 219.
16. R. Manickavasagam, K. Jeya, M. Paramasivam and S. Venkatakrishnalyer , *AntiCorros. Methods Mater* 49 (2002)19.
17. M. Morooka, I. Sekine, T. Tanaki, N. Hirosett and M.Yuasa , *Corros. Eng* 50 (2001) 106.
18. J. Jianguo, W. Lin, V. Alego and D.Schweinsberg , *Corros. Sci* 37 (1995) 975.
19. A.Dubey and G.Singh , *Port Electrochim Acta* 25 (2007) 221.
20. S.Umoren, E.Ebenso, P.Okafor and O.Ogbobe , *Pigment Resin Technol* 35(2006) 346.
21. Y. Wu, *Journal of Applied Solution Chemistry and Modeling* 2(2013) 33-46.
22. A. Labena, K.I. Kabel, R.K. Farag, *Materials Science and Engineering* 58 (2016) 1150–1159.
23. H. Bergenudda, P. Erikssona, C. DeArmittb, B. Stenberga and E.M. Jonssona, *Polymer Degradation and Stability* 76 (2002) 503–509.
24. G. A. El-Mahdy, A.M. Atta, H. A. Al-lohedan and A. O. Ezzat, *Int. J. Electrochem. Sci.*, 9 (2014) 7925 – 7934.
25. Q.Chen, R.G.Chen, L.R.Xiao, Q.R.Qian and W. G. Zhang , 27 (2008) 877.
26. Y. Zheng, Y. G.Li, L.Pan and Y. S. Li, *Polymer* 48 (2007) 2496.
27. S. J. Wang, X. W. Ba, B. H. Zhao, S. W. Zhang and W. L.Hou, *Acta Polym. Sin.* (2004) 5, 634–639.
28. S. Issaadi, T. Douadi, A. Zouaoui, S. Chafaa, M.A. Khan and G. Bouet, *Corros. Sci.* 53 (2011).
29. O.Mokhtari, I.Hamdani, A.Chetouani, A.Lahrach, H.El Halouani, A.Aouniti and M.Berrabah, *Journal of Material Environmental Science*, 5 (2013)310.
30. A.S Fouda, Y.Aggour, G.Bekheit and M.A. Ismail, *International Journal of Advanced Research*, 2 (2014)1158.
31. A. El-Shafei, S.A. Abd El-Maksoud and A.S. Fouda, *Corrosion Science*, 46 (2004).579.
32. A. Abdel-Gaber, B.A. Abd-El-Nabey and M.Saadawy, *Corrosion Science*, 51 (2009)1038.
33. N.A. Negm, N.G. Kandile, E.A. Badr and M.A. Mohammed, *Corros. Sci.* 65 (2012) 94.
34. J.C. da Rocha, J.A.C.P. Gomes, E. D'Elia, *Corros. Sci.* 52 (2010) 2341.
35. M. Moradi, J. Duan and X. Du, *Corros. Sci.* 69 (2013) 338.
36. A. Istiaque, P. Rajendra and M.A. Quraisi, *Corros. Sci.* 52 (4) (2010) 1472.
37. B. Qian, J. Wang, M. Zheng and B. Hou, *Corros. Sci.* 75 (2013) 184.
38. N.A. Negm, E.A. Badr, I.A. Aiad, M.F. Zaki and M.M. Said, *Corros. Sci.* 65 (2012) 77.
39. M. Behpour, S.M. Ghoreishi, N. Soltani, M. Salavati-Niasari, M. Hamadani and A. Gandomi, *Corros. Sci.* 50 (2008) 2172..

