



KINETICS AND THERMODYNAMICS OF THE CORROSION OF MILD STEEL IN THE PRESENCE OF *EUPHOBIA TIRUCALLI* GUMS

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Keywords: *Euphobia tirucalli*, adsorption isotherm, activation energy, corrosion inhibition

Understanding the mechanism of corrosion of steel in acidic environments along with the accompanying quest for corrosion inhibitors that are nontoxic to the environment is of significant industrial concern. This work sets to investigate the corrosion of mild steel in 1.0 M solution of HCl in the presence of *Euphobia tirucalli* gums (ET) using weight loss and linear polarization measurements. The study was carried out within the inhibitor concentration range of 0.2-1.0 g L⁻¹ and at the temperature range of 303 - 333 K. From the results obtained, it was observed that ET gums inhibited the dissolution of mild steel in the acid medium. Thermodynamic parameters obtained from the studies revealed the corrosion inhibition process to be spontaneous, exothermic and physisorptive. The adsorption of ET gums on the steel surface aligned well with the Langmuir, Freundlich and Temkin isotherm models. Linear polarization results revealed that ET functioned as a mixed-type inhibitor by inhibiting both the cathodic and anodic reactions on the mild steel surface.

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Introduction

Corrosion is the process through which metals/alloy deteriorates. The presence of moisture, acids, bases, salts, aggressive metal polishes and electrolytes generally would lead to the deterioration of most metals/alloys. According to Ahmad and Roberge,^{1,2} the tendency of a metal/alloy to corrode to a large extent is a function of the type of metal/alloy and the aqueous medium.

Mild steel is known for its widespread industrial applications due to its structural and mechanical properties. It is, however, susceptible to corrosion when it gets in contact with electrolytes resulting in a relatively short life span.^{1,2} Corrosion of steel structures and pipes result in significant losses of products in industries, environmental pollution and ecological disasters which have attracted much attention and investigations.^{3,4}

A substance is said to be a corrosion inhibitor when on its addition to a corrosive environment, it either slows down or prevents corrosion. These substances can function by adsorbing both physically and chemically at the metal/solution interface thereby serving as a barrier between the metal and the corrosive species in the environment.⁵⁻⁷ Peter et al. and Ramyan *et al.*,^{7,8} described corrosion inhibitors as usually possess heteroatoms such as oxygen, nitrogen and sulfur that have hydrocarbon parts that are typically attached to or part of a polar group, an aromatic ring or a conjugated system.

Many inorganic and organic compounds have been used and are still being employed in mitigating corrosion of metals and alloys. However, most of these compounds are expensive coupled with the fact that they pose health and

environmental challenges hence the quest for cheap and environmentally friendly inhibitors as substitutes.^{5,7,8}

Plant gums have been reported as good corrosion inhibitors against the dissolution of steel in acidic environments.⁹⁻¹⁴ Umoren has investigated the inhibitive effects of plant gum exudates from *Raphiahookeri* (RH) against the corrosion of mild steel in H₂SO₄ and Ebenso.⁹ The results obtained revealed that RH acted as a suitable corrosion inhibitor for the corrosion of mild steel in sulphuric acid medium. Abdallah¹⁰ investigated the potential of *Guar gum* (GG) as a corrosion inhibitor for carbon steel in 1.0 MH₂SO₄ solution using weight loss and Tafel polarization techniques. The results showed that GG inhibited the corrosion of carbon steel in the acid medium and acted as a mixed type inhibitor. In other studies, other gums such as *Gum arabic* (GA), *Albizia zygia* (AZ), *Anogessus leiocarpus* (Al) and *Ficus platphylla* (FP) have also been reported to be good corrosion inhibitors.¹¹⁻¹⁴

The results of phytochemical and pharmacological studies carried out by Prabha *et al.*, and Prashant and Shital on *Euphobia tirucalli* (ET) gum exudates showed that the gum exudates contain triterpenes, diterpene esters, steroids, alkaloids, flavonoids, tannins, phenols, anthraquinones and cardiac glycosides^{15,16} which are sources of heteroatoms.

This study sets to investigate the kinetics and thermodynamics of the dissolution of mild steel in 1.0 M solution of HCl in the absence and presence of *Euphobiatirucalli* gums using weight loss and linear polarization measurements.

Experimentals

Mild steel coupons of dimension 3x2 cm² and percentage composition: Si-0.03 %, P-0.04 %, S-0.04 %, Mn- 0.60 % and the rest Fe, were used for this study. Analar grade reagents were used for the study.

Purification of *Euphobia tirucalli* gum.

The procedure adopted for the purification of gum has been reported elsewhere by Femi et. al., and other researchers.^{17,18} The gums were oven dried at 40 °C for two hours after which they were pulverized with a blender and hydrated in chloroform water of double strength for five days with intermittent stirring such that all the gums particles dissolve completely. A 75 μ m sieve was used to strain the hydrated gums to obtain a pure slurry which was allowed to sediment. Absolute ethanol was used to precipitate the gum sediment from the slurry which was later filtered and defatted using diethyl ether. The precipitate was dried again by placing it in an oven at the temperature of 40 °C for 48 hours, after which the dried flakes were pulverized using a blender. The pulverized gums were kept in a dry and air tight container. Inhibitor test solutions (0.2, 0.4, 0.6, 0.8 and 1.0 g L⁻¹ concentrations)¹⁹ were prepared by dissolving appropriate quantities (0.05 g - 0.25 g) of the gum exudates in 250 mL of 1.0 M HCl solution.

Weight loss measurement

Mild steel coupons were cleaned, weighed and immersed in 250 mL test solutions in the absence and presence of ET gum exudates (0.2, 0.4, 0.6, 0.8 and 1.0 g L⁻¹) as reported in previous studies.^{19,20} They were removed at specified time intervals, washed in water, cleaned with ethanol, dried with acetone and reweighed. The average loss in weight was taken as the difference between the initial and final weights of the mild steel (in duplicates).

The corrosion rate of mild steel as a function of time was investigated by carrying out the experiment for 24 – 168 hours. This was done in the absence and presence of ET exudates. Temperature studies were done in a similar manner for three hours with the water bath maintained at 303- 333 K.

By carrying out the necessary substitutions in equations 1 to 3, weight loss data were used to evaluate the weight loss (g h⁻¹), corrosion rate (g h⁻¹ cm⁻²), inhibition efficiency (% *IE*) and the degree of surface coverage (θ).

$$CR(\text{g h}^{-1} \text{cm}^{-2}) = \frac{\Delta W}{At} \quad (1)$$

$$IE(\%) = 100 \left(1 - \frac{W_1}{W_2} \right) \quad (2)$$

$$\theta = 1 - \frac{W_1}{W_2} \quad (3)$$

where $\Delta W = W_2 - W_1$ is the weight loss of mild steel after time, t , W_1 and W_2 are the weight loss (g) of mild steel with and without the inhibitor, respectively, θ is the extent of surface coverage of the inhibitor, A is the area of the metal coupon (in cm²) and t is the immersion period (in hours).⁶

Linear polarization measurements

The coupons of 3 x 2 cm² dimension were sealed with epoxy resin in such a way that only 1 cm² was left exposed. A three-electrode system consisting of mild steel as working electrode, platinum (Pt) as an auxiliary electrode and saturated calomel electrode (SCE) as a reference electrode was used. Linear polarization studies were carried out using Metrohm Autolab AUT50280 in the potential range of -1500 to 1500 mV with a scan rate of 0.012 V s⁻¹ at room temperature.

The linear Tafel segments of the anodic and cathodic curves were extrapolated to corrosion potential to obtain the corrosion current densities (I_{corr}). The % *IE* was obtained from the equation below:

$$\%IE = 100 \frac{I_{\text{corr}(\text{blank})} - I_{\text{corr}(\text{inh})}}{I_{\text{corr}(\text{blank})}} \quad (4)$$

where $I_{\text{corr}(\text{blank})}$ is the corrosion current density of MS in the free acid solution and $I_{\text{corr}(\text{inh})}$ is the corrosion current density of MS in the inhibited acid solution.⁷

Optical microscopy

The morphologies of the surface of the polished mild steel coupons before immersion in the test solutions and after immersion in the test solutions were investigated using a metallurgical microscope (Tsview digital metallurgical microscope, model: Tucsen 0923502).

Results and discussion

The corrosion rate of mild steel and inhibition efficiency of ET gums

Figure 1 showed the rate of corrosion of mild steel in the free acid and inhibited 1.0 M HCl solution as a function of immersion time. It can be observed from the plot in Figure 1 that the rate of corrosion of the steel decreased with immersion time and as the concentrations of ET gums increased.

Figure 2 illustrates the trend of inhibition efficiency (% *IE*) of various concentrations of ET gums with immersion time. Inhibition efficiency of the gums increased as the concentrations of ET gums increased with % *IE* > 98 up to the immersion time of 96 h before decreasing.

This trend may be due to the desorption of the constituents of ET gums from the surface of the steel over a prolonged exposure resulting in the inhibition efficiency been lowered as observed in the plots.

Olasehinde *et al.*, and Momoh-Yahaya *et al.*^{20,21} in previous studies reported the decrease in inhibition efficiency of *Nicotiana tabacum* extracts and adenine

molecules in acidic environments at more extended immersion periods.

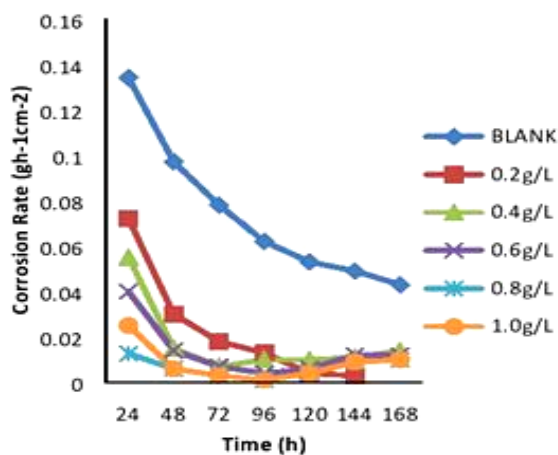


Figure 1. The corrosion rate of mild steel as a function of time in the absence and presence of ET gums

In both studies, the desorption of the constituents of *Nicotiana tabacum* extracts and adenine molecules from the surface of the mild steel was attributable to the decrease in inhibition efficiency observed.

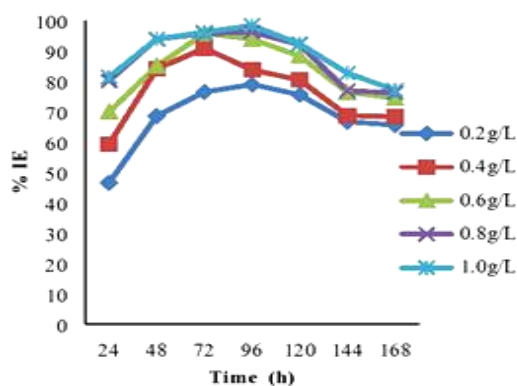


Figure 2. Inhibition efficiency of ET gums as a function of time (h).

Temperature considerations

Figure 3 showed the trend in the rate of corrosion of mild steel in uninhibited and inhibited 1.0 M solution of HCl at 303 - 333 K. The plots reveal that the rates of corrosion of mild steel increased generally at higher temperature values both in the free acid solution and in the presence of ET gums. Thermodynamically, the average kinetic energy of reacting species in a system is a function of the temperature. Reacting species become more energetic with a rise in temperature, this explains the increased rates of dissolution and corrosion rates of the mild steel in both the uninhibited and inhibited test solutions at higher temperatures. Other researchers corroborate this observation.²⁰⁻²³ However, the rate of corrosion of mild steel was relatively lower in the inhibited test solutions compared with the uninhibited acid solution. This signifies the inhibitive effect of ET gums on the acid corrosion of mild steel.

The plot of the inhibition efficiency of ET gums as a function of temperature (303-333 K) is presented in Figure 4. The plot shows a decrease in inhibition efficiency of ET gums as the temperature of the system was increased. This could be attributed to the agitation of adsorbed ET gums and consequent desorption of ET gums from the mild steel surface as temperature increased.²⁰⁻²² The lowering of inhibition efficiency values with increasing temperature is associated with a physisorptive mechanism of adsorption as reported in other works.²⁰⁻²⁴

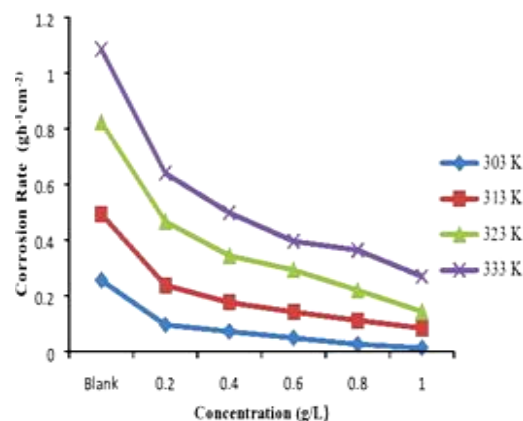


Figure 3. The corrosion rate of mild steel in 1.0 M HCl as a function of the concentration of ET gums at 303-333 K.

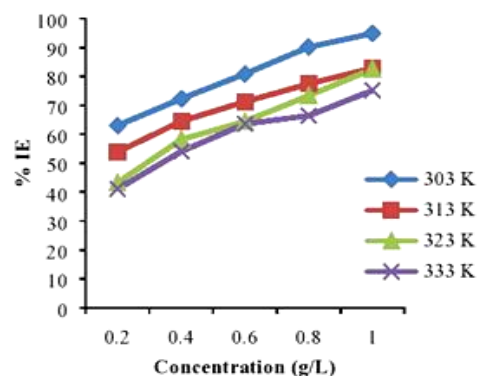


Figure 4. Inhibition efficiency of ET gums as a function of concentration at 303- 333 K.

The relationship between the corrosion rate of mild steel in the various test solutions and the temperature was evaluated using the Arrhenius equation (equation 5). Taking logarithm of both sides of equation 5, equation 6 is obtained.

$$CR = A \exp - E_a / RT \quad (5)$$

$$\log CR = \log A - \frac{E_a}{2.303RT} \quad (6)$$

where CR is the rate of corrosion of the metal, A is the Arrhenius or pre-exponential factor, E_a is the apparent effective activation energy, R is the universal gas constant and T is the absolute temperature of the system (K).

A plot of $\log CR$ versus $1/T$ represented on Figure 5 gave a straight line graph with a slope of $(-E_a/2.303RT)$ and an intercept of $[\log A]$ from which the values of the apparent activation energy (E_a) were evaluated.^{21,23,24} The values of E_a are shown in Table 1. It is evident from Table 1 that the value of E_a in the uninhibited system increased on the addition of ET gums as reflected in the values of E_a in the inhibited systems. This behavior is suggestive of a physisorptive kind of adsorption mechanism whereby an increase in temperature leads to the desorption of the adsorbed inhibitor from the steel surface thereby exposing the mild steel surface to the acid solution and leading to arise in the rate of corrosion of the steel as reported by Szauer and Brandt along with other researchers.^{25,23,24}

Enthalpy (ΔH^\ddagger) and entropy (ΔS^\ddagger) of activation of the corrosion inhibition were calculated using the Eyring equation;

$$\log \frac{CR}{T} = \left[\log \left(\frac{R}{nh} \right) + \left(\frac{\Delta S^\ddagger}{2.303R} \right) \right] - \frac{\Delta H^\ddagger}{2.303RT} \quad (7)$$

where CR is the corrosion rate at temperature T , R is the molar gas constant, n is Avogadro's constant 6.0225×10^{23} and h is the Planck's constant (6.6261×10^{-34} J s). A plot of $\log CR/T$ versus $1/T$ as shown in Figure 6 is a straight line graph with a slope of $(-\Delta H^\ddagger/2.303R)$ and an intercept of $[\log(R/nh) + \Delta S^\ddagger/2.303R]$ from which the values of ΔH^\ddagger and ΔS^\ddagger were calculated.^{21,23,24}

The results presented in Table 1 show that the enthalpies of activation are all negative which reveals the exothermic nature of the dissolution process of the mild steel. Also, the entropies of activation were all negative for the gums.⁷ Negative values of ΔS^\ddagger have been reported by Refat and Ishaq²⁵ to represent the association mechanism of the corrosion process whereby a decrease in disorder takes place ongoing from the reactants to the activated complex. Shukla and Ebenso corroborate this explanation in their findings on the adsorptive behavior and thermodynamic properties of streptomycin in the corrosion of mild steel in 1.0 M HCl.²³ In their study, the values of ΔS^\ddagger were not only negative but higher in the inhibited systems.

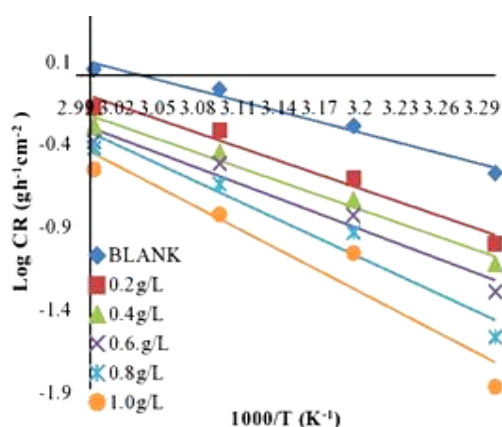


Figure 5. Arrhenius plot of the temperature -dependence of the corrosion rate of mild steel in 1.0 M HCl in the absence and presence of ET gums.

The values of ΔS^\ddagger in this study (Table 1) aligns with the findings of Shukla²³ and Refat.²⁵ The entropy of activation, ΔS^\ddagger in this present study increased in the presence of ET gums. This may be as a result of the adsorption of ET gums on the surface of the steel thereby slowing the discharge of hydrogen ions on the surface of the mild steel and the rate of corrosion of the steel.²³⁻²⁵

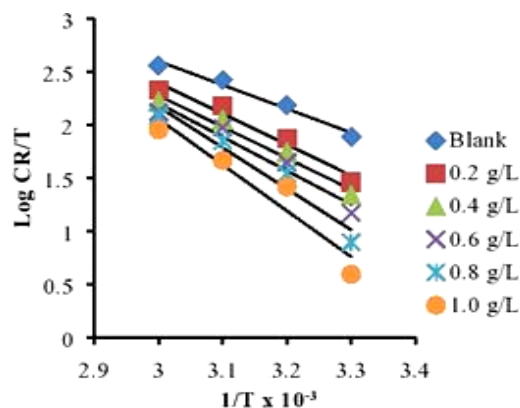


Figure 6. Eyring plot of the temperature-dependence of the corrosion rate of mild steel in 1.0 M HCl in the absence and presence of ET gums.

Table 1. Thermodynamic parameters of the corrosion of mild steel in 1.0 M HCl in the presence and absence of ET gums at 303-333 K.

Concentration, g L ⁻¹	E_a , kJ mol ⁻¹	ΔH^\ddagger , kJ mol ⁻¹	ΔS^\ddagger , J mol ⁻¹ K
Blank	16.280	-4.509	-132.0
0.2	20.104	-5.569	-113.6
0.4	20.736	-5.774	-108.1
0.6	22.457	-6.220	-66.01
0.8	25.941	-7.185	-75.45
1.0	29.018	-8.037	-68.69

Adsorption isotherms

Surface coverage (θ) values derived from weight loss measurements were adapted into various adsorption isotherms in order to know the adsorption characteristics of ET gums. The Langmuir, Freundlich and Temkin isotherms were found to give good descriptions of the adsorption characteristics of ET gums with all the plots having linear slopes and regression coefficients values $R^2 \geq 0.9$.^{5,23} The functionalized and linear forms of the Langmuir, Freundlich and Temkin isotherm models⁵ are the followings:

Isotherm	Functional form	Linear form
Langmuir	$k_{ads} C = \frac{\theta}{1-\theta}$	$\frac{C}{\theta} = C + \frac{1}{k_{ads}}$
Freundlich	$k_{ads} C^n = \theta$ ($0 < n < 1$)	$\log \theta = \log k_{ads} - n \log C$
Temkin	$k_{ads} C = \exp(f\theta)$	$-2\alpha\theta = 2.303(\log k_{ads} - \log C)$

where C is the concentration of the inhibitor in the bulk electrolyte, K_{ads} is the adsorption equilibrium constant, Θ is the degree of surface coverage of the inhibitor and α is the molecular interaction parameter.²⁰ Table 2 presents the adsorption parameters evaluated from the plots.

The adsorption equilibrium constant, K_{ads} is related to the standard free energy of adsorption, ΔG_{ads} as showed in equations 8 and 9,

$$\log K_{ads} = -1.744 - \frac{\Delta G_{ads}^0}{2.303RT} \quad (8)$$

$$\Delta G_{ads} = -2.303RT \log(55.5K_{ads}) \quad (9)$$

where ΔG_{ads} is the standard free energy of adsorption of ET gums on the mild steel surface, R is the universal gas constant, 55.5 is the concentration of water in solution in mol L⁻¹ and K_{ads} is the equilibrium constant of adsorption. The values of ΔG_{ads}^0 were negative and less than -20 kJ mol⁻¹ for all the three adsorption models.

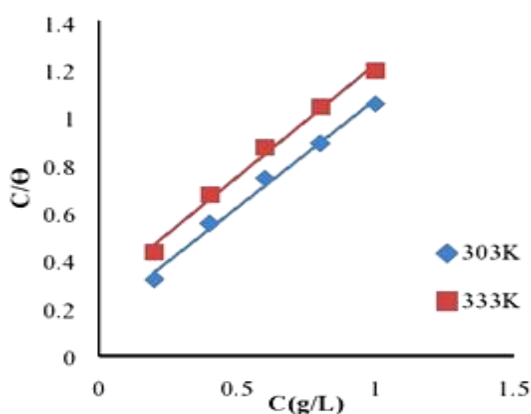


Figure 7. Langmuir isotherm for the adsorption of ET gums on mild steel surface at 303 K and 333 K respectively.

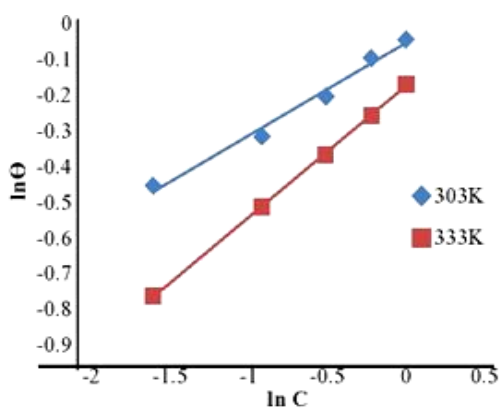


Figure 8. Freundlich isotherm for the adsorption of ET on mild steel surface at 303 K and 333 K respectively.

This implies that the adsorption of ET gums on the surface of the mild steel coupon is spontaneous and physisorptive.⁷ The values of $\Delta G_{ads}^0 \leq -20$ kJ mol⁻¹ and $\Delta G_{ads} \geq -40$ kJ mol⁻¹ have been reported in studies conducted by Momoh-Yahaya *et al.* and Oguzie *et al.* to signify the mechanisms of physisorption and chemisorption respectively.^{21,5}

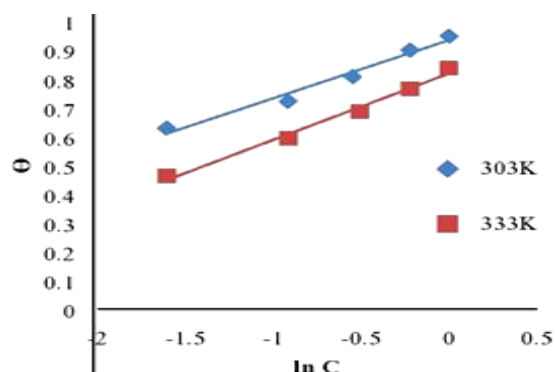


Figure 9. Temkin isotherm for the adsorption of ET on mild steel surface at 303 K and 333 K respectively.

Linear polarization

Linear polarization plots for the corrosion of mild steel in the various test solutions are shown in Figure 10 while Table 3 presents all the linear polarization parameters derived from the polarization plots and inhibition efficiency values.

The corrosion current density is a function of the reactivity of a metal in an aqueous environment. The higher the values of I_{corr} , the higher the dissolution of the metal and vice versa. Addition of ET gums is observed to reduce the anodic and cathodic current densities and the corresponding corrosion current density (I_{corr}) by shifting the corrosion potential (E_{corr}) toward more negative (cathodic) values.

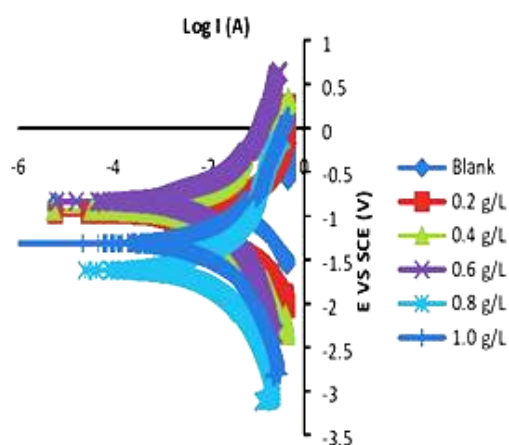


Figure 10. Linear polarisation plots of mild steel in 1.0 M HCl in the absence and presence of ET.

These shifts in the corrosion potential (E_{corr}) are more pronounced at higher ET concentrations, and the cathodic inhibiting effect becomes more significant. This implies that the gums functioned as a mixed-type inhibitor, with predominant cathodic effect.

Table 2. Adsorption parameters of ET gums on mild steel surface in 1.0 M HCl

	Temperature, K	Slopes	Intercepts	R^2	α	K_{ads}	$\Delta G_{ads}, \text{kJ mol}^{-1}$
Langmuir	303	0.904	0.169	0.991	-	5.921	-14.60
	333	1.076	0.276	0.990	-	3.376	-14.49
Freundlich	303	0.625	-0.2264	0.977	-	1.684	-11.43
	333	1.050	-0.608	0.976	-	4.051	-14.10
Temkin	303	0.324	0.778	0.934	0.324	11.04	-16.17
	333	0.292	0.535	0.949	0.292	6.23	-16.18

Table 4. Linear polarisation parameters for the corrosion of mild steel in 1.0 M HCl in the absence and presence of ET.

Conc., g L^{-1}	E_{corr}, mV	$I_{corr}, \mu\text{Acm}^{-2}$	$\beta_a, \text{V dec}^{-1}$	$\beta_c, \text{V dec}^{-1}$	CR, mmyr^{-1}	%IE
Blank	-843.79	299.86	0.47551	0.31335	348.44	-
0.2	-513.90	135.76	0.64010	0.54541	157.75	54.73
0.4	-844.30	266.10	0.52436	0.29977	147.12	57.78
0.6	-836.65	22.93	0.35607	0.32226	26.643	92.35
0.8	-1620.4	17.81	0.25595	0.28927	20.692	94.06
1.0	-1315.2	20.31	0.26930	0.23854	23.599	93.23

The values of b_a and b_c changed as the inhibitor concentration increased, signifying the influence of ET gum exudates on both the dissolution of the mild steel in 1.0 M HCl and the kinetics of hydrogen evolution. This is revealed in similar studies by Oguzie *et al.* and others.^{3-5,30}

Linear polarization

Linear polarization plots for the corrosion of mild steel in the various test solutions are shown in Figure 10 while Table 3 presents all the linear polarization parameters derived from the polarization plots and inhibition efficiency values.

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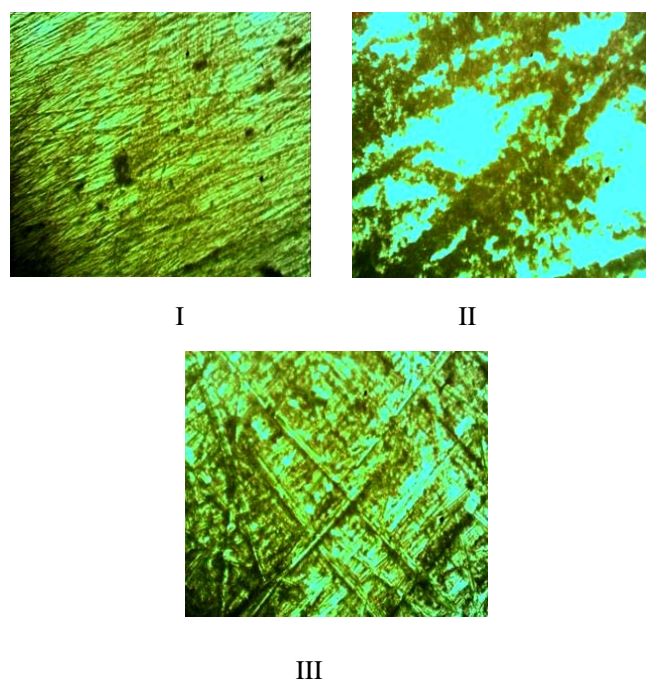
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Optical microscopy

Plates I, II and III show the micrographs of the polished mild steel, the polished steel after immersion in the uninhibited acid solution and the polished steel after

immersion in the inhibited acid solution respectively. Plate II shows a steel surface that is relatively damaged compared to Plates I and III. This is as a result of the rust formed on it after undergoing corrosion in 1.0 M solution of HCl in the absence of the inhibitor. The micrograph of mild steel surface in Plate III shows no pits and cracks except some polishing lines and the presence of an excellent protective film.

This further confirms the adsorption of the constituents of ET gums on the mild steel surface and the inhibitive property of ET gums against the corrosion of mild steel in 1.0 M solution of HCl.



Plates: I -Polished mild steel; II - Polished mild steel in 1.0 M HCl; III - Polished mild steel in 1.0 M HCl with ET gums.

Conclusion

The results obtained from this study have shown that *Euphorbia tirucalli* gum exudates acted as a useful and efficient inhibitor against the dissolution of mild steel in 1.0 M HCl. Inhibition efficiency of the gums was found to be concentration dependent. However, the inhibition efficiency of the gums decreased with immersion time and rose in temperature. Values of E_a , ΔH^\ddagger and ΔG_{ads} obtained showed that the adsorption of *Euphorbia tirucalli* gums exudates on the steel surface was physisorptive, exothermic and spontaneous. Linear polarization measurements showed that *Euphorbia tirucalli* gum exudates functioned as a mixed corrosion inhibitor by inhibiting and retarding the rates of both the anodic metal dissolution and cathode hydrogen ion reduction reactions. Optical micrographs confirm the adsorption of *Euphorbia tirucalli* gum exudates on the surface of the mild steel.

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Received: 18.01.2019.
Accepted: 27.02.2019.