Opuntia ficus-indica (Nopal) extract as green corrosion inhibitor for carbon steel in HCI1M solution

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The effect of *Opuntia ficus-indica* (Nopal) as green corrosion inhibitor for carbon steel in HCl 1M solution has been investigated by weight loss measurements, potentiodynamic polarization curves and electrochemical impedance spectroscopy. Also, scanning electron microscopy (SEM), and Fourier transform infrared spectroscopy (FT-IR). The inhibitor concentrations used ranged from 0 to 300ppm at 25, 40 and 60°C. The results indicated a) the inhibition efficiency increases with increasing extract concentration and b) the inhibitor act as a mixed-type inhibitor. The inhibitor has been associated with adsorption effects. In fact, the adsorption of the inhibitor on the steel surface follows the Langmuir adsorption isotherm, indicating monolayer adsorption. On the whole, the best efficiency was obtained at 300ppm and 60°C.

Keywords: Carbon steel, Corrosion inhibition, Adsorption.

1. INTRODUCTION

Acid solutions are often used in industry for cleaning, decaling and pickling of steel structures, processes which are normally accompanied by considerable dissolution of the metal. Corrosion of metals, however, is considered to be a serious problem in most industries. The new generation of environmental regulation requires the replacement of toxic inhibitors with non-toxic inhibitors. In this context, many alternative eco-friendly corrosion inhibitors have now been developed. A number of organic compounds are known to be applicable as corrosion inhibitors for steel in acidic environments. An immense number of scientific studies have been devoted to the inhibitive action of green inhibitors on the corrosion of mild steel in acidic solutions, showing that these extracts could serve as good corrosion inhibitor; the

cited extracts include Brugmansia-suaveoles and Cassia-roxburghil¹, Musaparadisica², Zenthoxylum-alatum³, Spirulina-platensis⁴, Olea-europaea⁵, Punicagranatum⁶, Lupinous-albus⁷, Occimum-viridis, Telferia-occidentalis, Azadirachtaindica and Hibiscus-sabdariffa⁸, Murraya-koenigii⁹, Medicago-sative¹⁰, Arecacatechu¹¹, Rosmarinus-officinalis¹², Ilex-paraguariensis¹³, Gundelia-tournefortii¹⁴. A number of organic compounds represent this type of inhibition, particularly those containing elements of Group V and VI of the periodic table, such nitrogen, phosphorous, arsenic, sulphur, oxygen and selenium. The efficiency of an organic compound as an inhibitor is mainly dependent upon its ability to get adsorbed on a metal surface¹⁵. It can then retard the cathodic and/or anodic reaction, thus, reducing the corrosion rate¹⁶. The stability of the adsorbed inhibitor film on the metal surface depend on some physicochemical properties of the molecule related to their functional groups, aromaticity, the possible steric effects, electronic density of donor atoms, type of corrosive environment and the nature of the interaction between the π orbital of the inhibitors and the d orbitals of iron^{17,18}. This study aimed at investigating the inhibition effect of Opuntia ficus-indica extract on mild steel in 1M HCl solution using weight loss measurements and electrochemical techniques. The inhibitor was investigated and characterized using FT-IR, SEM and Thermodynamic analysis.

2. EXPERIMENTAL

2.1. Preparation of specimens

The mild steel specimens of composition C = 0.15%, Mn = 0.70%, P = 0.010%, S = 0.027%, Cr = 0.016%, Ni = 0.12%, Al = 0.006%, Cu = 0.044% and the balance Fe. The bar of mild steel with diameter of 1 cm was cut off in section of 1 cm for weight loss study. The specimens were polished successively by use of SiC papers of 100, 260, 400, 600 and 800 grade; and then thoroughly cleansed with distilled water and then with ethanol, being dried later on and kept in a desiccator till their use. While coupons of size 0.7850 cm² were used for electrochemical studies. It was encapsulated in commercial epoxy resin.

2.2. Inhibitor preparation

50 g of *Opuntia ficus-indica* fresh were soaked in 100 ml using double distilled water and refluxing the solution for one hour. After cooling, solutions were filtered followed by drying in vacuum oven for one night (lyophilized). The extract solid was used as a corrosion inhibitor.

2.3. Solution preparation

The corrosive medium was 1 M HCl prepared with 38% analytical grade supplied by Sigma-Aldrich. Double distilled water was used for the preparation of all reagents.

2.4. Weight loss measurements

Mild steel specimens were immersed in 50 ml of 1M HCl with various extract concentrations (0, 50, 75, 100, 150, 200 and 300ppm) for time of exposition of 4.5, 6.5, 12, 24, 48 and 120 hours. After a total time of exposition specimens were taken out, washed with double distilled water, degreased with methanol, dried and weighted accurately. The weight loss (in grams), was taken as the difference in the weight of the mild steel specimens before and after immersion in different test solutions. The test was performed in triplicate to guarantee the reliability of results, and the mean value of the weight loss is reported. Tests were performed at room temperature 25, 40 and 60°C by using a hot plate. Corrosion rates, in terms of weight loss measurements, ΔW , were calculated as follows:

$$\Delta W = (m_1 - m_2) / A \tag{1}$$

were m_1 is the mass of the specimen before corrosion, m_2 the mass of the specimen after corrosion, and A the exposed area of the specimen. For the weight loss test, inhibitor efficiency (IE) was calculated as follows:

IE (%) = 100
$$(\Delta W_1 - \Delta W_2) / \Delta W_1$$
 (2)

were ΔW_1 is the weight loss without inhibitor, and ΔW_2 the weight loss with inhibitor.

2.5. Electrochemical techniques

The electrochemical experiments were performed using a typical three electrode cell (A platinum rod was used as counter electrode and saturated calomel electrode (SCE) as reference electrode) at room temperature (25, 40 and 60°C) and naturally aerated conditions. Polarization curves were recorder at a constant sweep rate of 1 mV/S at the interval from -500 to +500 mV respect to the E_{corr} value. The polarization curves were studied in 50 ml 1M HCl solutions using Potentiostat / Galvanostat Autolab-84861. The values of inhibition efficiency (%) were determined from equation (3), where, *lcorr*₁ and *lcorr*₂ are current densities with and without addition of inhibitor.

 $IE (\%) = 100 (Icorr_2 - Icorr_1) / Icorr_2$

The impedance studies were carried out using AC signals of 10mV amplitude for the frequency spectrum from 100 MHz-100Kz. The charge transfer resistance values were calculated from the diameter of the semi-circles of the Nyquist plots. The impedance studies were studied using Solartron Impendace / Gain-Phase analyzer SI 1260. The corrosion inhibition efficiency (%) was determined by equation (4), where Rct_1 and Rct_2 are the charge transfer resistances in presence and absence of inhibitor.

$$\mathsf{IE}(\%) = 100 \left(Rct_2 - Rct_1 \right) / Rct_2 \tag{4}$$

2.6. Fourier Transform Infrared Spectroscopy (FT-IR)

FT-IR spectrum was recorder with a frequency ranging from 4000 to 700 cm⁻¹ for the solution of the *Opuntia ficus-indica* extract in 1M HCI and specimen. The immersed was for 12 hours in 60°C. After solvent evaporation, the surface film was scraped carefully and its FT-IR spectra were recorded using Perkin Elmer model spectroscopy.

2.7. Scanning electron microscope studies

Mild steel specimens were immersed in a corrosive environment of 1M HCl having an optimum concentration (300ppm) of the *Opuntia ficus-indica* extract for 12 hours at 60°C. At the end of the experiment, the specimens were washed with distilled water, dried, and examined for their surface morphology using JEOL-JSM5800LV model scanning electron microscope.

3. RESULTS AND DISCUSSION

3.1. Weight loss measurements

Figure 1 shows the effect of inhibitor concentration on the inhibitor efficiency at the different tested temperatures. It can be seen that the inhibitor efficiency increased in 40 and 60°C to a maximum value at 300ppm as the inhibitor concentration was increased. The increase in inhibitor efficiency is due to the increase in the number of constituent molecules of *Opuntia ficus-indica* extract adsorbed on the metal surface at higher concentrations, so that the active sites of the metal are protected by inhibitor molecules.

Table 1 shows the values of inhibition efficiencies at different temperatures, indicating that inhibition efficiency increased at higher temperatures.



Figure 1. Inhibition efficiency of mild steel specimens at different immersed times in 1M HCl with and without *Opuntia ficus-indica* extract using on the weight loss methods. A is 25°C, B is 40°C and C is 60°C.

Table 1. Values of Inhibition Efficiency (%) from weight loss measurement for mild steel corrosion in 1M HCl with and without addition of different concentrations of *Opuntia ficus-indica* extract at different temperatures.

Inhibitor Concentration	Temperature (ºC)	Inhibition Efficiency (%)
0		
50		37
75	25	88
100		62
150		57
200		55
300		59
0		
50		54
75		70
100	40	74
150		77
200		85
300		89

0		
50		21
75		40
100	60	46
150		67
200		75
300		90

3.2. Polarization measurements

Polarization curves for mild steel at various concentration of Opuntia ficus-indica extract in 1M HCl at 25, 40 and 60°C are shown in Figure 2. The corrosion current density (*Icorr*), corrosion potential (*Ecorr*), and cathodic and anodic Tafel slopes (bc and ba) were obtained by extrapolation of the anodic and cathodic regions of the Tafel plot. The IE (%) was calculated using the equation 3. The electrochemical parameters obtained from the polarization measurements are listed in Table 2. As can be seen in Figure 2, the addition of *Opuntia ficus-indica* extract to the corrosive solution reduces the anodic dissolution of iron and also retards the cathodic hydrogen evolution reactions as would be expected. Both corrosion current density and corrosion rate were considerably reduced in the presence of the extract. These results are indicative of the adsorption of inhibitor molecules on the mild steel surface. The inhibition of both anodic and cathodic reactions is increasingly pronounced when increasing Opuntia ficus-indica extract concentration. These results suggest that Opuntia ficus-indica extract can be classified as the mixed type corrosion inhibitor. The highest inhibitor efficiency (90%) was obtained by adding 300 ppm of extract at 60°C.





Figure 2. Polarization plots of mild steel obtained in 1M HCl solution in absence and presence of various concentrations of *Opuntia ficus-indica* extract. A is 25°C, B is 40°C and C is 60°C.

Table 2. Determination of the electrochemical parameter for mild steel from polarization measurements.

Temperature (ºC)	Concentration (ppm)	βa (V/dec)	βc (V/dec)	Ecorr, (V)	Icorr (A/cm²)	Corr Rate (mm/year)	Inhibition Efficiency (%)
	0	0.064926	0.090826	-0.28173	0.0002077	2.4134	
	50	0.122878	0.046586	-0.25755	1.9356E-05	0.22491	90
	75	0.084406	0.152522	-0.30368	1.8258E-05	0.21216	91
25	100	0.075675	0.123373	-0.30496	3.7522E-05	0.43601	82
	150	0.044807	0.088169	-0.26157	4.0119E-05	0.46619	81
	200	0.122935	0.047867	-0.25224	4.3884E-05	0.50992	79
	300	0.085946	0.162444	-0.33769	5.1445E-05	0.59779	75
	0	0.094302	0.075496	-0.33252	0.00022423	2.6055	
	50	0.102270	0.048715	-0.34594	7.0567E-05	0.81998	69
40	75	0.034715	0.063781	-0.30228	3.1007E-05	0.36029	86
	100	0.106687	0.090766	-0.38353	2.4502E-05	0.28471	89
	150	0.066782	0.078306	-0.38698	1.5964E-05	0.18546	92
	200	0.069395	0.12349	-0.34371	1.5584E-05	0.18108	93
	300	0.036185	0.051838	-0.37397	7.2976E-06	0.08479	97
	0	0.37557	0.36803	-0.33576	0.0050078	58.194	
	50	0.28928	0.44131	-0.31354	0.0040058	46.547	20
	75	0.11158	0.06993	-0.32969	0.0002504	2.9097	95
60	100	0.12765	0.05343	-0.33041	0.0002074	2.4104	96
	150	0.12523	0.11455	-0.37001	0.0001707	1.9836	97
	200	0.08674	0.04573	-0.33536	6.6708E-05	0.7751	98
	300	0.05853	0.03505	-0.32324	4.1273E-05	0.4795	99

3.3. Electrochemical impedance spectroscopy measurements

Figure 3 shows the representative Nyquist plots of mild steel obtained in 1M HCl solution in the absence and presence of various concentrations of Opuntia ficusindica extract. The Nyquist plots of mild steel obtained in blank solution was magnified and added in Figure 3. The Nyquist plots of mild steel showed a depressed semi-circular shape. This behavior indicates that the corrosion of mild steel in 1M HCl solution was mainly controlled by a charge transfer process. Although the appearance of Nyquist plot remained the same, their diameter increased after the addition of Opuntia ficus-indica. This increase was also pronounced with increasing inhibitor concentration which indicates the adsorption of molecules on the metal surface. The impedance parameters derived from the Nyquist plots and percentage inhibition efficiency are given in Table 3. Table 3 shows that the Rct values increased and Cdl values decreased indicating a more controlled anodic and cathodic processes and the decrease in the capacitance values, which were attributed to the formation of the protective layer at the mild steel surface. To obtain the double layer capacitance (Cdl) values, the following equation was used:

$$Cdl = 1 / 2\pi f_{max}Rct$$

The inhibition efficiency was calculated using the equation 4. The highest inhibitor efficiency (99.1 %) was obtained by adding 300ppm of *Opuntia ficus-indica* extract in 60°C.



Figure 3. Nyquist plots of mild steel obtained in 1M HCl solution in the absence and presence of various concentrations of *Opuntia ficus-indica* extract in 60°C.

(5)

Table 3. Electrochemical impedance parameter values for the corrosion of mild steel in 1M HCl at 60°C in the absence and presence of *Opuntia ficus-indica* extract.

Temperature (ºC)	Concentration (ppm)	Rct (Ω cm ²)	Cdl (µF cm ⁻²)	Inhibition Efficiency (%)
		4.4	3617.2	
	50	170.4	93.6	97.4
60	75	197.3	80.8	97.7
	100	230.6	69.3	98.1
	150	260.5	61.2	98.3
	200	345.2	46.1	98.7
	300	350.8	45.5	99.1

3.4 Adsorption Behavior

To investigate adsorption behavior of *Opuntia ficus-indica* extract in 1M HCl solution, numerous isotherm models were employed, such as, Langmuir, Freundlich, Flory-Huggins, Frumkin, and Temkin, but the best fit was obtained for the Langmuir isotherm model. The Langmuir adsorption could be represented by the following equation:

$$C/\theta = (1/K) + C \tag{6}$$

Where *C* is the concentration of inhibitor, θ is surface coverage and K is the adsorption constant. The surface coverage (θ) of the inhibitor on the mild steel surface is expressed by following equation:

$$\theta = IE(\%) / 100 \tag{7}$$

The mechanism of corrosion inhibition may be explained on the basis of adsorption behavior. Basic information on the interaction between the inhibitor and the metal surface can be provided by an adsorption isotherm. The adsorption parameters, such as, regression coefficient (R^2), adsorption constant (K) and free energy of adsorption(ΔG) and slope values were obtained by straight line fitting between C/θ (y-axis) and C(x-axis). The Figure 4 shows the Langmuir isotherm of the polarization curves method.

The most important thermodynamic adsorption parameter is the free energy of adsorption (ΔG). The adsorption constant (*K*) is related to the standard free energy of adsorption, ΔG is calculated with the following equation:

$$\Delta G = -RTln(55.5K) \tag{8}$$

Where 55.5 is the water concentration of solution in mol / L. R is the ideal gas constant, T is the absolute temperature. Table 4 shows the values calculated in the

Langmuir isotherm. The negative values of ΔG indicate the stability of the adsorbed layer on the mild steel surface and spontaneity of the adsorption process. Generally, the magnitude of ΔG around -20 kJ/mol or less negative is assumed for electrostatic interaction that exist between inhibitor and the charged metal surface (physisorption). Their mechanism can be classified as mixed-type inhibitor.



Figure 4. Langmuir isotherm for adsorption of *Opuntia ficus-indica* extract on the mild steel surface. Polarization curves data.

Table 4	. Thermodynamic	parameters	for mild	steel in	1M HCI	in	presence	of	the
Opuntia	ficus-indica extrac	ct at different	concent	trations.					

Isotherms	Temperature (ºC)	Slope	R ²	Kads	ΔG ads KJ.mol ⁻¹
Langmuir	25	1.380	0.9986	55.50	-19.90
	40	0.014	0.9994	71.43	-21.56
	60	1.004	0.9997	250	-26.40

3.5. Fourier Transform Infrared Spectroscopy (FT-IR)

Figure 5 shows the FT-IR spectrum of the extracts at different concentrations. A careful investigation of the spectra revealed that all the extract showed almost similar peaks; however, the intensities decreased or increased with addition of the *Opuntia ficus-indica* extract. A strong and broad peak at 3000/3400 cm⁻¹ can be attributed to N-H and O-H stretching vibration. A small peak at 1720 cm⁻¹ was observed before the test. Such a peak may be attributed to C=O stretching vibration, peaks were increased with addition of the *Opuntia ficus-indica* extract in 1610 cm⁻¹. The absorptions bands at 1627/1588 cm⁻¹ were also observed due to a N-H bending vibration. A peak at 1394 cm⁻¹ was attributed to C-N bending vibration.

before and after test. A small peak at 1222 cm⁻¹ was observed before and after the test, attributed to C-N stretching vibration. Peaks before and after in 1074 cm⁻¹ attributed to C-O stretching vibration. Thus, results showed that *Opuntia ficus-indica* extract contain organic molecules that are rich in oxygen and nitrogen atoms as well as aromatic rings, which meet with the fundamental requirements of good inhibitor. A summary of these results is given in Table 5.



Figure 5. FT-IR spectrum of *Opuntia ficus-indica* extract before and after corrosion test.

Table 5. Peaks from FT-IR spectra of *Opuntia ficus-indica* before and after corrosion test, and their identification.

Acid + 300 ppm inhibitor before test		Acid + 300 ppm inhibitor after test		
Frequency (cm ⁻¹)	Functional group and vibration type	Frequency (cm ⁻¹)	Functional group and vibration type	
3118	N-H stretch	3368	N-H stretch	
3018	O-H stretch	3189	O-H stretch	
1720	C=O stretch	1610	C=O stretch	
1627	N-H bending	1588	N-H bending	
1394	C-H bending	1394	C-H bending	
1222	C-N stretch	1222	C-N stretch	
1074	C-O stretch	1074	C-O stretch	

3.6 SEM images

The scanning electron microscope images were recorded to establish the interaction of inhibitor with the metal surface. Figure 6A indicates the finely polished characteristic surface of mild steel. Figure 6B revealed that the surface was severely corroded due to the aggressive attack by 1M HCI. Figure 6C reveal the formation of a protective film by the inhibitor on the mild steel surface in acid medium. Images 6B and 6C show specimens immersed 12 hours at 60°C. The image 6C containing 300ppm of *Opuntia ficus-indica* extract in 1M HCI.



Figure 6. SEM images of (A) polished mild steel specimens. (B) mild steel specimens in HCI 1M. (C) mild steel specimens with *Opuntia ficus-indica* extract in HCI 1M at 60°C.

4. CONCLUSIONS

Opuntia ficus-indica extract act as a good corrosion inhibitor for mild steel in 1M HCl solution. The inhibition efficiency increased with increasing temperature. The change in free energy carries negative values around -20kJ / mole which indicate that the adsorption process is spontaneous and physical adsorption, respectively. The inhibitor efficiency increased (40 and $60^{\circ}C$) with concentration of *Opuntia ficus-indica* extract. The inhibitor altered both anodic and cathodic Tafel slopes, which showed the mixed mode of action of the inhibitor molecule. The increase in *Rct* values and decrease in *Cdl* values confirm the formation of an insulated protective layer over the mild steel surface, which was supported by SEM images. The corrosion inhibitive effect shown by *Opuntia ficus-indica* extract can be correlated to the presence of organic compounds in its chemical structure.

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