


# Performance of Methanolic Extract of *Artemisia herba alba* as a Potential Green Inhibitor on Corrosion Behavior of Mild Steel in Hydrochloric Acid Solution

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**Abstract:** The influence of the methanolic extract of *Artemisia herba alba* (MEAHA) as a potential green inhibitor on the corrosion behavior of mild steel in 1M HCl solution in the presence of different concentrations (200 to 500 ppm) was explored and investigated using weight loss (WL) method, potentiodynamic polarization (PDP) technique, electrochemical impedance spectroscopy (EIS) measurement, scanning electron microscope (SEM) analysis, energy dispersive spectroscopy (EDS), temperature effect, Langmuir adsorption isotherm, and kinetic thermodynamic parameters. Obtained results suggest that the increases in MEAHA concentration lead to increases in the anti-corrosive protection efficiencies, which are 91, 89, and 90% for WL, PDP, and EIS, respectively. SEM analysis resulting after soaking in the corrosion inhibitor solution, the surface of mild steel is still relatively smooth for 24 h. Adsorption of MEAHA obeyed Langmuir adsorption isotherm model. Also, MEAHA green inhibitor is very well adsorbed by the active centers on MS area.

**Keywords:** MEAHA; green inhibitor; corrosion behavior; mild steel; MEB/EDS; Langmuir adsorption.

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## 1. Introduction

Recently, several researchers have been studying the development and investigation of new methods, materials, and technologies that can delay the corrosion of steel using essential oil and extracts of *Artemisia herba alba* as a potential green corrosion inhibitor in aggressive acidic mediums (HCl, H<sub>3</sub>PO<sub>4</sub>, H<sub>2</sub>SO<sub>4</sub>, HNO<sub>3</sub>, etc.) [1-4]. Methanolic extract of *Artemisia* is a natural substance that is not toxic for humans and environmentally friendly. It could be employed as a potential green inhibitor for mild steel protection in the hydrochloric acid solution (1 M HCl) [5]. The anti-corrosive inhibition efficiencies of natural inhibitors for mild steel include chemical conversion disposal, anodic oxidation, laser disposal, corrosion inhibitory and coatings, etc. Adding the methanolic extract of *Artemisia herba alba* has several advantages: the excellent anti-corrosive property, the low cost, and the very simple equipment, and widely employed in many industries such as chemical, cosmetic, other, etc. [6-8]. Corrosion inhibition of the mild steel surface is protected by the physical, chemical, and/or

physicochemical adsorption between the metallic surfaces and products investigated to form insoluble thin films by chemical adsorption prevent further corrosion of substrate [9-15]. The effective adsorption of corrosion inhibition requires the presence of multiple bonds and the heterocyclic compounds having heteroatoms such as nitrogen, sulfur, oxygen, phosphorus, etc. [16-22]. The development and investigation of new green inhibitors and the improvement of anti-corrosive protective efficiency are the primary objectives of green corrosion inhibitor research. Presently, the corrosion research of mild steel in 1M HCl medium mainly focuses on the influence of microstructure and phase structure on their anti-corrosive performance. However, the influence of ion Cl and concentration on the anti-corrosive protection behavior of green corrosion inhibitor is reported in several pieces of literature [23-26]. The objective of our present study is the development and application of a methanolic extract of *Artemisia herba alba* as a potential green anti-corrosive inhibitor for mild steel in 1M HCl medium. Then, the methanolic extract of *Artemisia herba alba* was studied and investigated for outgoing application as a potential green corrosion inhibitor for mild steel using weight loss (WL), potentiodynamic polarization (PDP), electrochemical impedance spectroscopy (EIS), scanning electron microscopy (SEM), energy-dispersive X-ray spectroscopy (EDS), temperature effect, Langmuir adsorption isotherm and kinetic thermodynamic parameters [27, 28].

## 2. Materials and Methods

### 2.1. Extraction of the methanolic extract of *Artemisia Herba Alba*.

In the Imilchil area in the Atlas Mountains of Morocco, *Artemisia herba alba* was gathered fresh during February (2016). Randomly, it is dried in the shade, protected from moisture, and kept carefully in a non-humid and ventilated area within the Laboratory of Materials, Nanotechnology, and Environment at the Faculty of Sciences, Mohammed V University in Rabat. An electric mill is used to grind the aerial part of these species until a powder was obtained. Using Soxhlet for 8 h, 30 g of this aerial part's powder is subjected to methanol extraction [29, 30]. The extract was, therefore, concentrated completely in a rotary evaporator. Until its use is needed, the residue was saved in a tightly closed vial of colored glass at 4°C.

### 2.2. Material employed and corrosive solution.

In this study, mild steel (MS) specimens was employed as a material with a chemical composition (C=0.29, Cr=0.18, Mn=0.62, P=0.05, Cu=0.27, S=0.05, Ni=0.09, Al=0.02 and rest of iron). Also, the MS samples were treated and polished with emery paper (600 to 2000), rinsed with double-distilled water and degreased with propanone and ethanol at room temperature. For WL, PDP and EIS tests, the MS sample surface was 1 cm<sup>2</sup>. 1 M HCl solution was realized using a dilution of 37% HCl with bidistilled water [31]. The concentration range of MEAHA studied was 200 to 500ppm.

### 2.3. Electrochemical investigation.

The corrosion inhibition effect of MEAHA on MS samples was performed using the potentiostat PGZ 301 (Radiometer Analytical) apparatus. Then, the electrochemical cell is composed of three electrodes. They are mild steel that operates as a working electrode. The platinum serves as a counter electrode and the saturated calomel electrode stands as a reference

electrode, respectively [32]. The PDP parameter was 0.3 mV/s of scan rate in the range of -800 to 0 mV/SCE, and the EIS parameter was 10 mV of sine wave voltage and the frequency between 100 to 10 mHz. The effect of temperatures was realized and investigated at 303, 313, 323, and 333 K, respectively.

#### 2.4. SEM and EDS analysis.

Scanning electron microscope (SEM) and the energy dispersive spectroscopy (EDS) analysis (JEOL JSM-IT 100 type) was employed to investigate the morphology of the MS substrate area immersion in 1 M HCl uninhibited and inhibited by optimal concentration (500 ppm) of the MEAHA tested as a potential green inhibitor, with an accelerating voltage of 20 kV [33].

### 3. Results and Discussion

#### 3.1. Weight loss measurements.

Weight loss (WL) measurements were employed to study the anti-corrosion inhibition efficiencies for MS in 1M HCl solution without and with different concentrations of methanolic extract of *Artemisia herba alba* investigated as a potential green inhibitor. The corrosion rate (W) and the corrosion inhibition efficiency ( $\eta$  (%)) were calculated according to equations 1 and 2, respectively [34]. The W and  $\eta$  (%) are listed in Table 1 [35, 36].

$$W = \frac{(m_1 - m_2)}{S \cdot t} \quad (1)$$

$$\eta (\%) = \frac{(W^0 - W_{inh})}{W^0} \times 100 \quad (2)$$

Where  $m_1$ ,  $m_2$ ,  $W^0$ ,  $W_{inh}$ , S, and t denote the initial weight, the final weight, the corrosion rate without inhibitor, the corrosion rate with different concentrations of methanolic extract of *Artemisia herba alba*, the surface of MS and the time, respectively.

**Table 1.** Weight loss and inhibition efficiency without and with varying concentration of MEAHA

Concentrations (ppm)	W (mg cm <sup>-2</sup> h <sup>-1</sup> )	$\eta$ (%)	$\theta$
Blank	0.670		
200	0.226	66	0.66
300	0.141	79	0.79
400	0.091	86	0.86
500	0.059	91	0.91

As shown in Table 1, the weight loss decreases with increases in the methanolic extract of *Artemisia herba alba* concentrations, owing to that the  $\eta$  (%) increases. Then, the maximum  $\eta$  (%) value at the optimum concentration of methanolic extract of *Artemisia herba alba* (500 ppm) was 91 %. The increase of the MEAHA concentrations causes by an increase considerably in the anti-corrosive inhibition efficiencies. However, suppose the MEAHA Alba concentrations increase above an optimal value. In that case, the green inhibitor adsorbs on a mild steel surface by the electrostatic interaction.

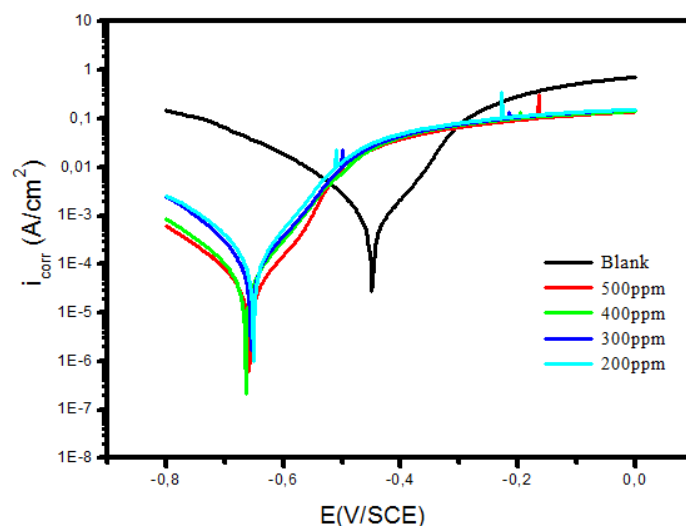
#### 3.2. Potentiodynamic polarization test.

Potentiodynamic polarization plots of mild steel in 1M HCl medium without varying concentrations (200 to 500 ppm) of MEAHA as a green inhibitor are shown in Figure 1. The

relevant electrochemical parameters are grouped in Table 2. The corrosion inhibition efficiency was calculated according to equation 3 [37-39].

$$\eta_{\text{PDP}} (\%) = \left( \frac{i_0 - i_{\text{inh}}}{i_0} \right) \times 100 \quad (3)$$

With  $i_0$  and  $i_{\text{corr}}$  denote the corrosion current density in the absence and presence of varying concentrations of methanolic extract of *Artemisia herba alba*.



**Figure 1.** Polarization plots for MS in 1M HCl medium without and with HEAHA.

**Table 2.** Varying electrochemical parameters obtained from PDP without and with different concentrations of MEAHA.

Inhibitor	Concentration ppm	$-E_{\text{corr}}$ (mV <sub>SCE</sub> )	$-\beta_c$ (mV.dec <sup>-1</sup> )	$\beta_a$ (mV.dec <sup>-1</sup> )	$i_{\text{corr}}$ (μA.cm <sup>-2</sup> )	$\eta_{\text{PDP}}$ (%)
MEAHA	1M HCl	449	92	186	570	-
	200	452	98	64	162	72
	300	656	111	71	117	79
	400	663	118	36	81	86
		659	128	52	65	89

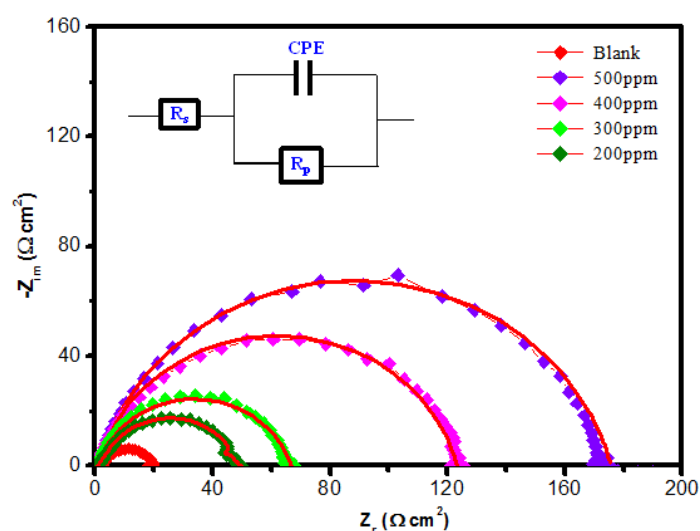
As a result, indicated in Figure 1 and Table 2, it is confirmed that the cathodic branches show a typical Tafel behavior, while the anodic branches in the presence of the methanolic extract of *Artemisia herba alba* as a green inhibitor reveal that has been related to the modification in the coverage degree of the MEAHA, which is owing to the reorientation of the heteroatoms of the MEAHA Alba formed on the mild steel surface [26]. The corrosion of mild steel is inhibited in 1M HCl medium alone using controlling both anodic dissolution and hydrogen evolution reaction, however by preventing surface blocking since both  $\beta_a$  and  $\beta_c$  are modified in the presence of a different concentration of MEAHA as a green inhibitor. Thus, there is a difference between the cathodic branch and the anodic branch. For a more detailed explanation, Figure 1 displays the PDP curves obtained without and in the presence of 200 to 500 ppm of MEAHA in 1 M HCl solution [27]. Noting that the corrosion current densities of the cathodic and anodic branches in aggressive corrosion solution decreasing with an increase in the concentration of green inhibitor studied, however the corrosion potential shifts toward the more negative values, indicating in corrosion inhibition mitigation, which is owing to the methanolic extract of *Artemisia herba alba* green inhibitor adsorbed on the metallic surface-active centers in 1 M HCl solution [14]. The MEAHA in 1M HCl solution is an cathodic-type inhibitor.

### 3.3. Electrochemical impedance spectroscopy test.

Figure 2 displays the electrochemical impedance spectroscopy plots performed on mild steel in 1M HCl medium in the absence and the presence of various concentrations of the methanolic extract of *Artemisia Herba Alba*. The different electrochemical parameters are grouped in Table 3. The reduction of corrosion inhibition of the mild steel is largely attributed to the presence of MEAHA, including the adsorption and the film formation on MS, owing to the adsorption of green inhibitor studied and the distance between the substrate surface and a corrosive solution is increased [40]. These facts could be seen through the Nyquist curves obtained from the EIS study. The protective inhibition efficiency was determined according to the following equation 4 [41]:

$$\eta_{\text{EIS}} (\%) = \left( \frac{R_{\text{inh}} - R_0}{R_{\text{inh}}} \right) \times 100 \quad (4)$$

Where  $R_0$  and  $R_{\text{inh}}$  denote the polarization resistance values in the absence and the presence of MEAHA inhibitor.



**Figure 2.** Nyquist curve for MS in 1M HCl solution without and with varying concentrations of MEAHA. Inset: Electrochemical equivalent circuits used to fit the impedance data.

**Table 3.** Varying electrochemical parameters obtained from EIS without and with different concentrations of MEAHA.

Inhibitor	C (ppm)	$R_s$ ( $\Omega \cdot \text{cm}^2$ )	$R_{\text{tc}}$ ( $\Omega \cdot \text{cm}^2$ )	$Q$ ( $\mu\text{F S}^{n-1}$ )	$n_{\text{dl}}$	$C_{\text{dl}}$ ( $\mu\text{F cm}^{-2}$ )	$\eta_{\text{EIS}}$ (%)
1M HCl	-	1.16	17	462	0.800	138	-
MEAHA	200	2.759	49	470.4	0.7113	167	65
	300	2.789	67	243.7	0.8087	122	75
	400	2.695	124	92.55	0.8398	31.08	86
	500	3.101	176	62.04	0.8381	52.31	90

As a result in Figure 2, alone one capacity loop appears in the frequency studies' impedance spectra, which suggests that the single charger transfer resistance is the main reason indicating in anti-corrosive in the absence and the presence of the methanolic extract of *Artemisia herba alba* green inhibitor. That is possibly related to the charge transfer reaction, the frequency dispersion, the metallic surface inhomogeneity, and the surface roughness [40]. As such, the higher semicircle implies that the covering film is present in the MS surface. That presence is made possible thanks to the MEAHA adsorption that works to reduce the corrosion process. Also, semicircles' diameter increases with the increase of green inhibitor concentration

up to 500 ppm [42]. At higher frequencies range, there is the one which could correspond to the metallic surface corrosion compounds. The other one is at lower frequencies, which may be attributed to the charge transfer process at the substrate/solution interface. An equivalent electrical circuit:  $R_s$  is solution resistance,  $R_p$  is polarization resistance, and CPE is a constant phase element [43, 44]. Owing to the surface roughness, the porous layer formation and the inhibitor adsorption, there is no ideal capacitive behavior in 1M HCl solution. Also, CPE is realized to replace the double layer capacity ( $C_{dl}$ ) to get the better-fitting plots with the non-ideal capacitance response [45]. The maximum inhibition efficiency in 1M HCl solution is 90%. Then, the  $R_s$  value increases with the increase of MEAHA concentration, resulting in that the chloride ions may reduce the solution's resistance. Also, the  $R_{ct}$  value elevates with the increase of MEAHA concentration, which is because that the increase of concentration of chloride ions strengthens the damage of the protective layer on the metallic surface and reduces the corrosion resistance (Table 3) [46].

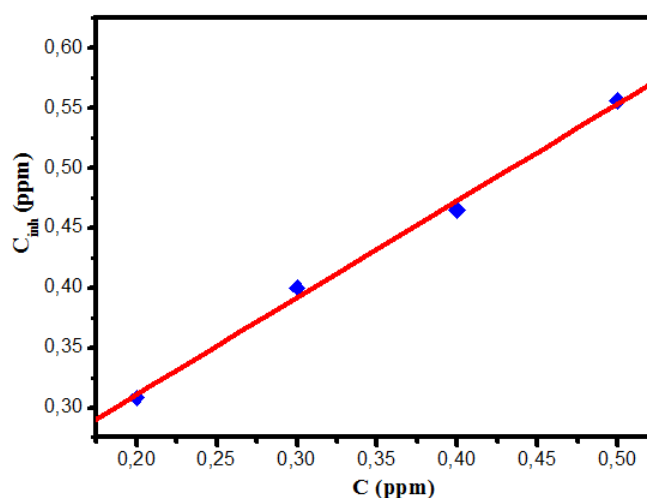
### 3.4. Adsorption isotherm.

The Langmuir adsorption isotherm of the corrosion inhibition on the metallic surface in 1M HCl solution is shown in Figure 3. As a result in Figure 3, when the  $R^2$  is close to the unit, the  $C$  is linearly related to the  $C/\theta$ . The Langmuir adsorption isotherm is defined as the following equations 5 and 6 [47, 48]:

$$\frac{C_{inh}}{\theta} = \frac{1}{K_{ads}} + C_{inh} \quad (5)$$

$$\Delta G_{ads}^0 = (-RT) \times \ln(55.5K_{ads}) \quad (6)$$

The  $\Delta G_{ads}^0$  value in 1M HCl medium investigated has to value close to  $-20 \text{ kJ.mol}^{-1}$ , resulting in that the physisorption dominates. Not alone  $K_{ads}$ , however, also  $\Delta G_{ads}^0$  of methanolic extract of *Artemisia herba alba* is the largest in 1M HCl, indicating the strongest adsorption as shown in Table 4, which is in perfect agreement with their similar electrochemical result.



**Figure 3.** Langmuir adsorption isotherm curve for the MEAHA in 1M HCl medium fitted by EIS data.

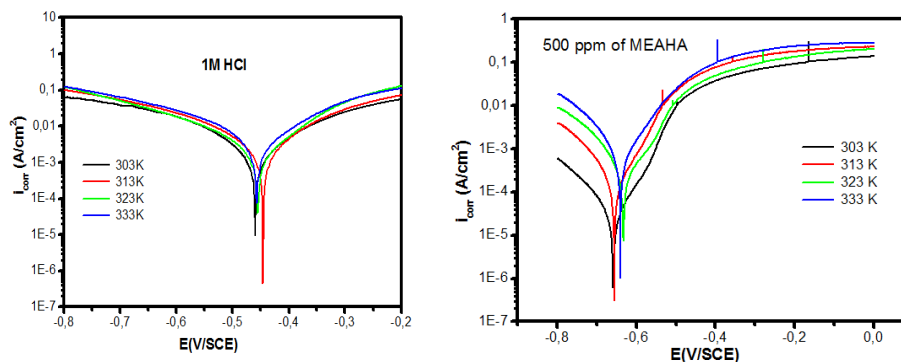
**Table 4.** Vary in parameters for Langmuir adsorption isotherm of MS with MEAHA.

Inhibitor	$R^2$	Slope	$K_{ads} \text{ (L/g)}$	$\Delta G_{ads}^0 \text{ (KJ.mol}^{-1}\text{)}$
MEAHA	0.99999	0.809	6.71	-22.2



### 3.5. Temperature effect and kinetic parameters.

The temperature effect was realized to investigate the resistance of the film adsorbed on the mild steel in 1M HCl without and with an optimal concentration of the MEAHA (500 ppm) as a potential green inhibitor (we investigated the potentiodynamic polarization in the temperature range 303 to 333 K) [49-51]. The PDP curves of the MS surface in 1M HCl only and with MEAHA (500 ppm) at varying temperatures are presented in Figure 4. The different electrochemical parameters are listed in Table 5.



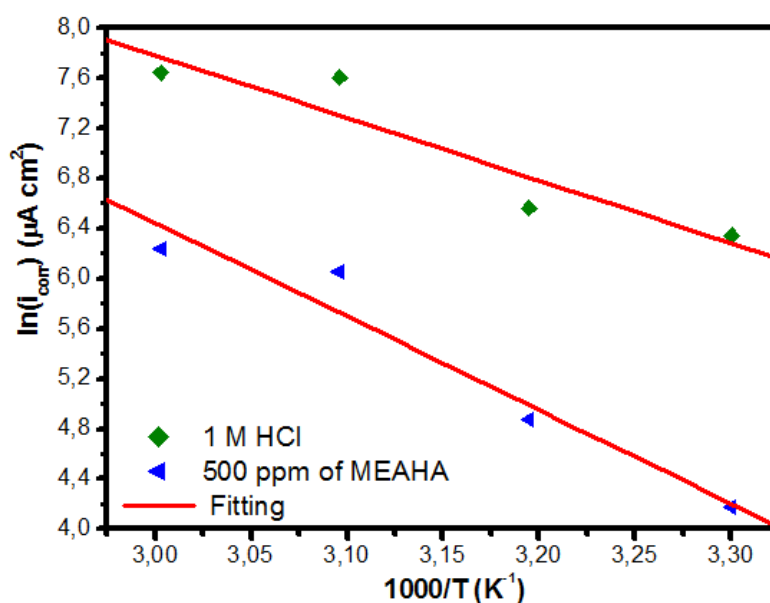
**Figure 4.** Temperature effect on the polarization plots of MS in 1 M HCl without and with 500 ppm of MEAHA at varying temperatures.

As a result, in Figure 4 and Table 5, we remarked that the corrosion current density for MS uninhibited and inhibited increases with an increase in temperature, which is owing to the diminishing of the anti-corrosive inhibition efficiencies. Several parameters were investigated to understand the adsorption mechanism of the anti-corrosion protection between the MEAHA and MS arias, such as activation energy ( $E_a$ ), standard activation enthalpy  $\Delta H_a^\circ$ , and standard activation entropy  $\Delta S_a^\circ$ . These parameters were calculated for MS uninhibited and inhibited with optimum concentration of the MEAHA according to the Arrhenius equation and alternative formula of the Arrhenius (equations 7 and 8) [14].

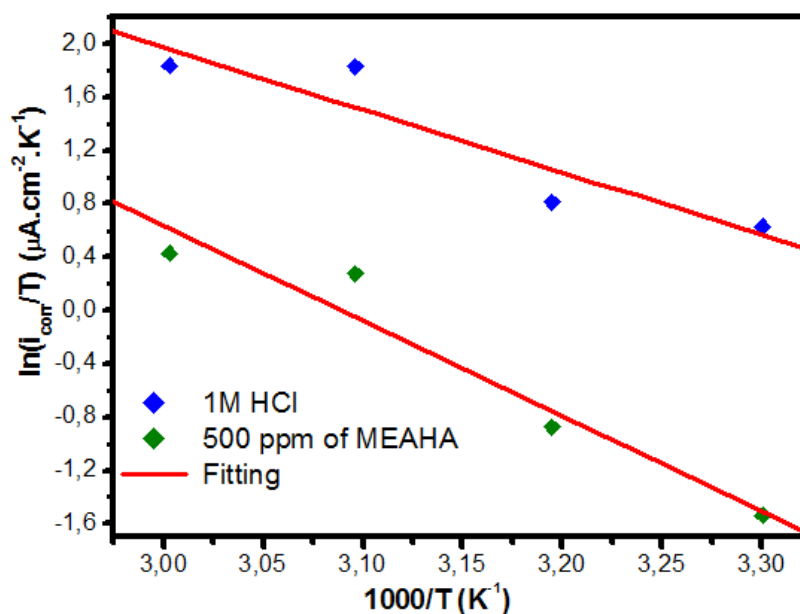
$$i_{\text{corr}} = K \exp\left(\frac{-E_a}{RT}\right) \quad (7)$$

$$i_{\text{corr}} = \frac{RT}{Nh} \exp\left(\frac{\Delta S_a^\circ}{RT}\right) \exp\left(-\frac{\Delta H_a^\circ}{RT}\right) \quad (8)$$

The  $\ln(i_{\text{corr}})$  as a function of  $1000/T$  for MS in HCl medium alone and with MEAHA at an optimal concentration (500 ppm) was displayed in Figure 5. Then, the  $\ln(i_{\text{corr}}/T)$  as a function of  $1000/T$  uninhibited and inhibited with MEAHA (500 ppm) was presented in Figure 6. Also, varying kinetic parameters such as  $E_a$ ,  $\Delta H_a^\circ$  and  $\Delta S_a^\circ$  calculated are grouped in Table 5. The activation energy value in the presence of the MEAHA is more than that of the HCl only, which owing that the higher energy was procured with MEAHA employed as a potential green inhibitor [52]. Also, the standard activation enthalpy positive value of methanolic extract of *Artemisia herba alba* suggests that the MS dissolution process is an endothermic reaction.



**Figure 5.**  $\ln(i_{\text{corr}})$  as function of  $1000/T$  for MS in HCl without and with 500 ppm of MEAHA.



**Figure 6.**  $\ln(i_{\text{corr}}/T)$  as a function of  $1000/T$  for MS in HCl without and with 500 ppm of MEAHA.

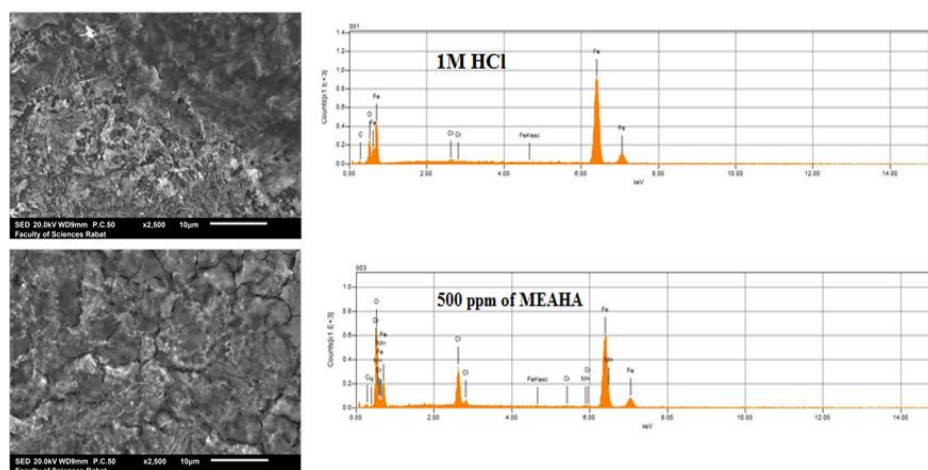
**Table 5.** PDP and kinetic parameters of MS in 1 M HCl solution in the absence and présence of 500 ppm of MEAHA at various temperatures.

Temperatures K	- E <sub>corr</sub> mV <sub>SCE</sub>	i <sub>corr</sub> μA cm <sup>-2</sup>	η <sub>PDP</sub> (%)	E <sub>a</sub> kJ mol <sup>-1</sup>	ΔH <sub>a</sub> kJ mol <sup>-1</sup>	- ΔS <sub>a</sub> J mol <sup>-1</sup> K <sup>-1</sup>
1M HCl						
303	449	570	-	41.52	38.88	64.46
313	445	708	-			
323	454	2013	-			
333	456	2090	-			
500 ppm of MEAHA						
303	659	65	89	62.04	59.40	14.04
313	655	131	81			
323	632	427	79			
333	642	511	75			



### 3.6. SEM/EDS characterization.

Using scanning electron microscopy (SEM), it has become possible to document the micrographs (Figure 7) of the mild steel strips' surfaces. The objective is to examine the transformations that occur during the corrosion process when 500ppm of AHAME is present and absent after 24h of immersion. So as to get enough information about the properties investigated in the sample of the surface of the steel both in the absence and presence of inhibitor in 1 M HCl, EDX analyses were done. Due to the absence of the extracts (Figure 7), the steel composition becomes hard and gets damaged after being immersed in 1 M hydrochloric acid solution because of the force of the rapid corrosion. As a matter of fact, there is a presence of a layer of corrosion product able to absorb fluids [53]. On the contrary, when we add to the aggressive environment 500ppm of plant extract, this leads to a remarkable decrease in mild steel's corrosive attack (Figure 7). As a result, the constitution of corrosion products gets hindered. This blockage leads to a process of changing to an inferior state of the mild steel. Indeed, as the SEM images demonstrate, there is a sort of improvement at the surface coverage level, and the protective film on the steel surface starts to take shape. The prime reason behind this is associated with the downward change in the interaction between the steel of the surface and the aggressive medium. It is evident from Figure 7 that the EDX spectrum contained in aggressive solution (1M HCl) introduces the characteristic and most extreme possible values of particular elements included in the surface of the sample's chemical properties. As such, when the plant extract exists, both the feature peak of nitrogen and enhancement in the intensity level of the peaks of oxygen and carbon become visibly apparent. These are present in the tested inhibitor's chemical constitution [1].



**Figure 7.** SEM/EDS images of the MS without and with 500 ppm of MEAHA.

## 4. Conclusions

The corrosion resistance of the corrosion inhibition of the methanolic extract of *Artemisia herba alba* used as a green inhibitor in 1M HCl solution was investigated, and the following conclusions were obtained: As the concentration of MEAHA increases, the  $R_s$  of the solutions and the  $R_{ct}$  of corrosion reaction was increased, and the optimal corrosion inhibition efficiencies are 91, 89, and 90% for WL, PDP and EIS, respectively; The resistance of corrosion inhibition is due to corrosion inhibitors' coverage to reduce mild steel's exposure surface; The small insoluble molecules react with corrosion products owing to the hydrophobic effect of corrosion inhibitors; SEM analysis indicates that the green inhibitor investigation

leads to smooth film formation on the metal surface in 1M HCl solution; These studies are very interesting and motivate researchers to improve the methanolic extract of *Artemisia herba alba* as a green inhibitor, as a corrosion inhibitor, and as an anti-corrosive coating.

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## Conflicts of Interest

The authors declare no conflict of interest.

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