

## Electrochemical Characterization of Catechol-Dimethylamine Adduct at Different pH Values

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### Abstract

The efficiency of *Commelina benghalensis* leaves extract as a mild steel corrosion inhibitor in hydrochloric acid (HCl) and sulphuric acid (H<sub>2</sub>SO<sub>4</sub>) has been evaluated. The corrosion rates were determined using the gravimetric (weight loss) and electrochemical techniques. The results obtained in the absence and presence of *Commelina benghalensis* extracts were used to calculate the inhibition efficiency (%), and to propose the inhibition mechanism and adsorption type. The obtained results showed that the inhibition efficiency increased with the increase in concentration of *Commelina benghalensis* extract. The phenomenon of chemical adsorption was proposed, and the inhibitor adsorption was found to obey Langmuir, Temkin and kinetic thermodynamic models. The polarization study reveals that, in the inhibitor presence, the corrosion current value decreases, indicating the formation of a protective film on the metal surface, which prevents the electron release from it.

**Keywords:** *Commelina benghalensis* leaves extract, corrosion inhibition, weight loss, polarization, impedance study, adsorption isotherm and green inhibitors.

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### Introduction

Corrosion is a ubiquitous problem that continues to be of great relevance in a wide range of industrial applications and products. The use of corrosion inhibitors is the most economical and practical method in reducing corrosive attack on metals. Nevertheless, the popularity and use of synthetic compounds as corrosion inhibitors are diminishing, due to the strict environmental regulations, and to the toxic effects of synthetic compounds on human and animal life. In

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order to find out non-toxic, cheap and effective green corrosion inhibitors from renewable sources, many natural products have been used by researchers.

Chraibi et al. have used *M. piperita* and *M. pulegium* essential oils for the corrosion control of mild steel in a 1.0 M hydrochloric acid solution [1]. These substances also have antibacterial activity. Evaluation of *Alpinia galanga* and its active principle, 1'-acetochavicol acetate, as an eco-friendly corrosion inhibitor on mild steel in acidic media, has been done by Ajeigbe et al. [2]. The anticorrosive activity of *Schreabera swietenioids* leaves as a green inhibitor for mild steel in an acidic solution has been reported by Sivakumar and Srikanth [3]. The corrosion inhibition of carbon steel in a 1 M H<sub>2</sub>SO<sub>4</sub> solution by *Thapsia villosa* extracts was investigated by Kalla et al. [4]. The use of *Piper longum* extract as a green corrosion inhibitor for aluminium in a NaOH solution has been studied by Singh et al. [5].

Junior et al. have evaluated the antioxidant activity and corrosion inhibition by Brazilian plant extracts and their constituents [6]. The inhibition effect of some trees cultivated in arid regions, against the corrosion of steel reinforcement in an alkaline chloride solution, has been evaluated by Etteyeb et al. [7].

Mott-Schottky analysis showed the formation of a passive layer on the metallic surface. EIS results suggest an increase in carbon steel corrosion resistance, in the plants extract presence, comparatively to the blank systems. Furthermore, SEM and EDS analyses corroborate these results.

An eco-friendly inhibitor for mild steel corrosion made from *Pennisetum purpureum* biomass and synergistic intensifiers has been formulated by Ituen et al. [8]. Kinetic data treatment indicated an increase in the energy barrier by intensifier ions. The results demonstrate that elephant grass extract, blended with halide ions, can act as an alternative eco-friendly inhibitor for mild steel, at elevated temperatures.

Dos Santos et al. have investigated carbon steel corrosion inhibition in acidic chloride by Mangiferin. The results showed that carbon steel exposed to this medium containing Mangiferin had a reduction in the mass loss process and a higher electrochemical behavior, relatively to the carbon steel samples exposed to the same medium, in Mangiferin absence. These results suggest that Mangiferin shows an inhibitory action on the carbon steel corrosion process, in the studied conditions [9].

Umoren et al. have evaluated mustard seed extract (MSE) as an eco-friendly corrosion inhibitor for X60 steel in acidic media [10]. The obtained results show that MSE inhibited steel corrosion in both media, which was more pronounced in H<sub>2</sub>SO<sub>4</sub> than in HCl environments. The inhibition efficiency increased with higher extract concentrations, but decreased with increased temperatures. The potentiodynamic polarization studies revealed that MSE functions as a mixed-type inhibitor. The corrosion inhibition is assumed to occur via adsorption of the extract components onto the steel surface, which was found to obey Langmuir adsorption isotherm model. The morphology of the corroding steel surface, in the absence and presence of the mustard seed extract, was visualized using scanning electron microscopy. Rajendran et al. have used extracts of various plant

materials, such as henna leaves, curcumin, caffeine and spirulina, to control metals corrosion [11-36].

The present work was undertaken to evaluate the inhibition efficiency of an aqueous extract of *Commelina benghalensis* leaves against mild steel corrosion in hydrochloric acid and sulphuric acid media. Weight loss and electrochemical methods, such as polarization study and AC impedance spectra, have been used.

## Materials and methods

The extract of *Commelina benghalensis* leaves was used as corrosion inhibitor in the present study. The leaves and flowers of *Commelina benghalensis* are shown in Fig. 1.



**Figure 1.** *Commelina benghalensis* plant.

The botanical details of *Commelina benghalensis* are given below: it belongs to the *Commelinaceae* family. The Tamil name is “Kanavazhiain”. In China, it is used as a medicinal herb that is said to have diuretic, febrifugal and anti-inflammatory effects, while in Pakistan it is used to cure skin swellings, leprosy, and as a laxative.

The phyto-chemical screening of *Commelina benghalensis* leaves extract is given in Table 1 [37].

**Table 1.** Phyto-chemical screening of *Commelina benghalensis* leaves extract.

Phyto-chemical constituents	Aqueous	Methanol	Hexane	CCl <sub>4</sub>
Flavonoids	+	-	-	-
Alkaloids	-	-	++++	++++
Glycosides	-	-	-	-
Steroids	-	-	-	-
Total Phenols and Tannins	+	+	-	-
Triterphenoids	-	-	-	-
Saponins	++	-	-	-
Proteins and amino acids	+++	+++	-	-
Anthraquinones	-	-	-	-

- = absent; + = present; ++ = moderately present; +++ = appreciable amount

Preliminary phyto-chemical screening mainly revealed the presence of saponins, phenols, flavonoids, alkaloids, tannins, proteins and amino acids.

**Extract preparation***Cold percolation method*

The leaves were collected, shade dried and powdered. About 1 g of the powdered leaves was macerated with 1000 mL of double distilled water, tightly covered with aluminium foil, and kept for 24 hrs. After 24 hrs, the macerated extract was filtered by a Whatmann filter paper. From the extract, the various concentrations were prepared.

*Used materials and chemicals*

Mild steel specimens (Wt %: 0.026%-S, 0.06%-P, 0.4%-Mn, 0.1%-C, and the remainder iron) with the dimensions 1 x 5 x 0.2 cm were abraded to mirror finish, degreased with acetone, and used for the weight loss method.

For the polarization study, a cylindrical mild steel rod embedded in Teflon, with an exposed area of 1 cm<sup>2</sup>, was used. The electrodes were abraded with emery papers of 0/0, 2/0, 3/0, and 4/0 grades, degreased with acetone, dried and used.

**Weight loss method**

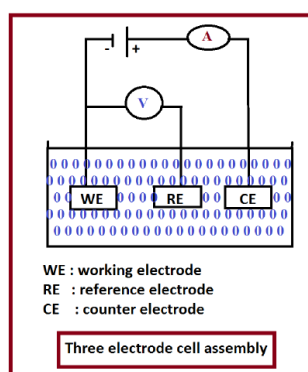
Weight loss measurements were performed in 1 M HCl and 0.5 M H<sub>2</sub>SO<sub>4</sub>, with and without the inhibitor presence. Inhibition efficiencies for various inhibitor concentrations were calculated using the relation:

$$\text{Inhibition efficiency (\%)} = \frac{W_0 - W_i}{W_0} \times 100$$

where  $W_0$  = weight loss in blank and  $W_i$  = weight loss in the inhibitor presence.

**Electrochemical studies**

The polarization study and AC impedance spectra were recorded in a potentiostat. The experiments were recorded in a three electrodes cell assembly, which is shown in Scheme 1.



**Scheme 1.** Three electrodes cell.

Mild steel was used as working electrode. A saturated calomel electrode (SCE) was used as reference electrode. A platinum foil (2 cm<sup>2</sup>) was used as counter electrode. The used working electrode was a mild steel rod of the same composition, embedded in araldite, and with an exposed area of 1 cm<sup>2</sup>.

From the potentiodynamic polarization study, the corrosion current, corrosion potential and the Tafel slopes were derived. From AC impedance spectra, the charge transfer resistance and double layer capacitance were calculated.

The counter electrode area is much larger compared to the working electrode area. This will exert a uniform potential field on the working electrode.

100 mL of the test solution were taken in a polarization cell. The working electrode was successively abraded with 0/0, 1/0, 2/0, 3/0, and 4/0 emery papers, and degreased with acetone. The working electrode, reference and auxiliary platinum electrodes were assembled, and connections were made. Stirring was provided to the test solutions, to avoid the system concentrations polarization before the experiment start. A time interval of about 30 min was given for the system to attain its state, and open circuit potential was recorded.

## Results and discussion

### Weight loss studies

The inhibition efficiency of *Commelina benghalensis* leaves extract in controlling mild steel corrosion in 0.5 M H<sub>2</sub>SO<sub>4</sub> has been evaluated by the weight loss method.

Tables 2 and 3 give the inhibition efficiency of different concentrations of *Commelina benghalensis* leaves extract in 1 N HCl and 0.5 M H<sub>2</sub>SO<sub>4</sub>, respectively. It was observed that, as the inhibitor concentration increased, the inhibition efficiency also increased. 900 ppm of inhibitor offered 93.40 % inhibition efficiency in 1 N HCl, and 85.71% in 0.5 M H<sub>2</sub>SO<sub>4</sub>.

**Table 2.** Inhibition effect of mild steel corrosion in 1 N HCl, using *Commelina benghalensis* leaves extract.

[Inhibitor] (ppm)	Rate of corrosion (g cm <sup>-2</sup> h <sup>-1</sup> )	Inhibition efficiency (%)
Blank	0.000183	-
100	0.000152	17.27
200	0.000138	24.55
300	0.000130	29.09
400	0.000102	44.55
500	0.000082	55.45
600	0.000060	67.27
700	0.000042	77.27
800	0.000028	84.55
900	0.000010	93.40

### Adsorption isotherms

Adsorption isotherms are usually used to describe the adsorption process. Adsorption isotherms are very important to determine the organo-electrochemical reaction mechanism. The most frequently used isotherms include: Langmuir, Temkin, Flory-Huggins, and the recently formulated thermodynamic/kinetic model of El-Awady et al. The use of adsorption isotherms that describe the adsorption of a corrosion inhibitor can provide important clues about the nature of the metal-inhibitor interaction. Adsorption of the organic molecules occurs when the interaction energy between the molecules

and the metal surface is higher than that between the H<sub>2</sub>O molecules and the metal surface. It is important to know the-adsorption mode, to understand this part of the study.

**Table 3.** Inhibition effect of mild steel corrosion in 0.5 M H<sub>2</sub>SO<sub>4</sub>, using *Commelina benghalensis* leaves extract.

[Inhibitor] (ppm)	Rate of corrosion (g cm <sup>-2</sup> h <sup>-1</sup> )	Inhibition efficiency (%)
Blank	0.000093	-
100	0.000085	8.92
200	0.000075	19.64
300	0.000065	30.35
400	0.000062	33.92
500	0.000042	55.35
600	0.000035	62.50
700	0.000028	69.64
800	0.000018	80.35
900	0.000013	85.71

### Langmuir isotherm

In order to obtain the adsorption isotherm, the degree of surface coverage ( $\Theta$ ) for various inhibitor concentrations has been calculated according to its equation. Langmuir isotherm was tested for its fit to the experimental data. Langmuir isotherm is given by:

$$C/\Theta = 1/K_{ads} + C$$

where  $\Theta$  is the degree of surface coverage,  $C$  is the molar inhibitor in the bulk solution and  $K_{ads}$  is the equilibrium constant of the adsorption process.

**Table 4.** Langmuir adsorption isotherm for mild steel corrosion inhibition in 1 N HCl, using *Commelina benghalensis* leaves extract.

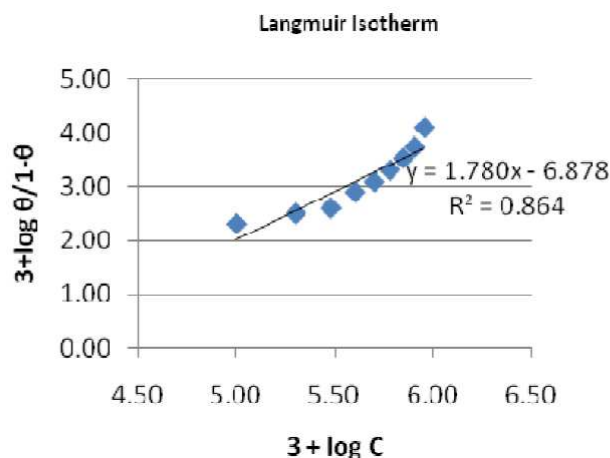
[Inhibitor] (ppm)	$\Theta/(1-\Theta)$	$3+\log\Theta/(1-\Theta)$	$3+\log C$
Blank	-	-	-
100	0.2088	2.3197	5.0000
200	0.3253	2.5123	5.3010
300	0.4103	2.6131	5.4771
400	0.8033	2.9049	5.6021
500	1.2449	3.0951	5.6990
600	2.0556	3.3129	5.7782
700	3.4000	3.5315	5.8451
800	5.4706	3.7380	5.9031
900	14.1500	4.1507	5.9542

Langmuir isotherm assumes that the metal surface contains a fixed number of adsorption sites, and that each site holds one adsorbate;  $\Delta G^0_{ads}$  is the same for all sites, and it is independent from  $\Theta$ ; the adsorbates do not interact with each other, i.e., there is no effect of the adsorbates lateral interaction on  $\Delta G^0_{ads}$ . It is important to know this part of the study. Tables 4 and 5 show that the obtained values were fitted to Langmuir isotherm, and the best fit to the experimental data was obtained.

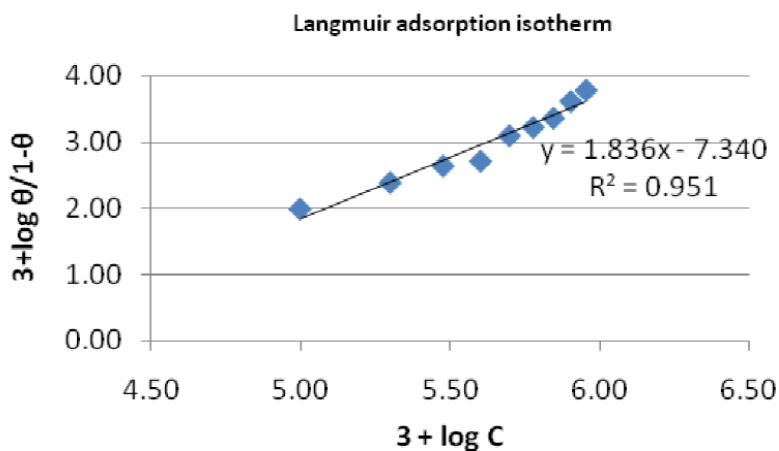
**Table 5.** Langmuir adsorption isotherm for mild steel corrosion inhibition in 0.5 M H<sub>2</sub>SO<sub>4</sub>, using *Commelina benghalensis* leaves extract.

[Inhibitor] (ppm)	$\Theta/(1-\Theta)$	$3+\log\Theta/(1-\Theta)$	$3+\log C$
Blank	-	-	-
100	0.0980	1.9914	5.0000
200	0.2444	2.3882	5.3010
300	0.4359	2.6394	5.4771
400	0.5135	2.7106	5.6021
500	1.2400	3.0934	5.6990
600	1.6667	3.2218	5.7782
700	2.2941	3.3606	5.8451
800	4.0909	3.6118	5.9031
900	6.0000	3.7782	5.9542

The plot of  $\log \log\Theta/(1-\Theta)$  vs.  $\log C$  is a straight line as shown in Figs. 2 and 3. Thus, the Langmuir isotherm is valid for the inhibitor.



**Figure 2.** Langmuir adsorption isotherm for mild steel corrosion inhibition in 1 N HCl, using *Commelina benghalensis* leaves extract.



**Figure 3.** Langmuir adsorption isotherm for mild steel corrosion inhibition in 0.5 M H<sub>2</sub>SO<sub>4</sub>, using *Commelina benghalensis* leaves extract.

**Temkin adsorption isotherm**

The Temkin adsorption isotherm is given by the expression:

$$\theta = -2.303 \log K/2a - 2.303 \log C/2a$$

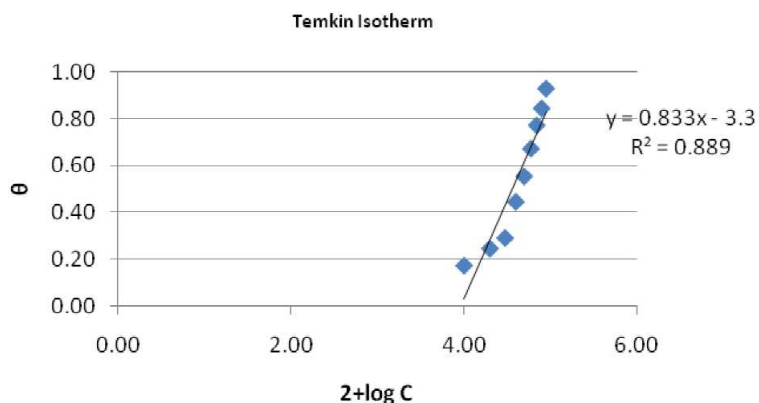
where 'K' is the adsorption equilibrium constant, and 'a' is the lateral interaction parameter. Tables 6 and 7 show the Temkin isotherm values, and Figs. 4 and 5 show the plots of (2+log C) against  $\theta$ , at various temperatures. The linear plot indicates that Temkin adsorption isotherm was obeyed, and 'a' negative value indicated repulsion in the adsorption layer.

**Table 6.** Temkin adsorption isotherm for mild steel corrosion inhibition in 1 N HCl, using *Commelina benghalensis* leaves extract.

[Inhibitor] (ppm)	2+log C	$\theta$
100	4.0000	0.1727
200	4.3010	0.2455
300	4.4771	0.2909
400	4.6021	0.4455
500	4.6990	0.5545
600	4.7782	0.6727
700	4.8451	0.7727
800	4.9031	0.8455
900	4.9542	0.9340

**Table 7.** Temkin adsorption isotherm for mild steel corrosion inhibition in 0.5 M H<sub>2</sub>SO<sub>4</sub>, using *Commelina benghalensis* leaves extract.

[Inhibitor] (ppm)	2+log C	$\theta$
100	4.0000	0.0893
200	4.3010	0.1964
300	4.4771	0.3036
400	4.6021	0.3393
500	4.6990	0.5536
600	4.7782	0.6250
700	4.8451	0.6964
800	4.9031	0.8036
900	4.9542	0.8571



**Figure 4.** Temkin adsorption isotherm for mild steel corrosion inhibition in 1 N HCl, using *Commelina benghalensis* leaves extract.

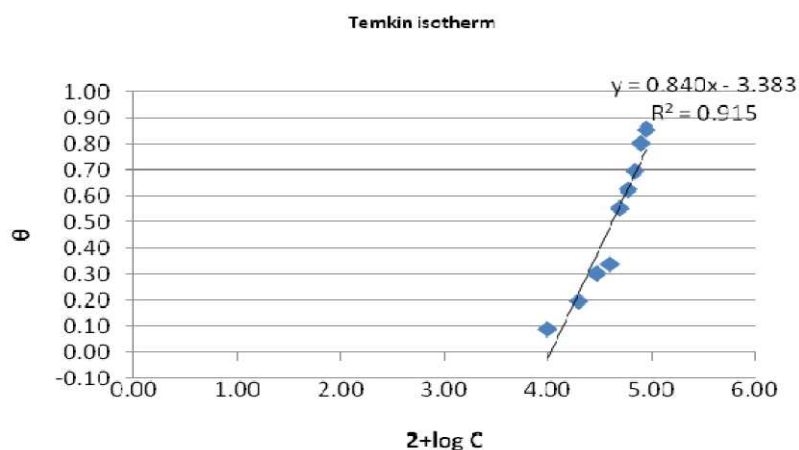


**Flory-Huggins adsorption isotherm**

Flory-Huggins adsorption isotherm is given by the expression,

$$\log \theta/c = \log K + x \log (1-\theta)$$

where 'x' is the size parameter, and it is a measure of the number of adsorbed water molecules substituted by an inhibitor molecule. Tables 8 and 9 and the linear plot of  $3+\log \Theta/C$  against  $3+\log (1-\Theta)$  (Figs. 6 and 7) show that Flory-Huggins isotherm was obeyed.



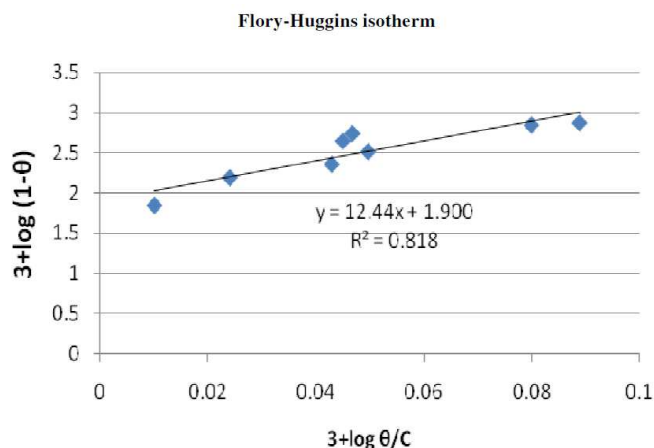
**Figure 5.** Temkin adsorption isotherm for mild steel corrosion inhibition in 0.5 M H<sub>2</sub>SO<sub>4</sub>, using *Commelina benghalensis* leaves extract.

**Table 8.** Flory-Huggins adsorption isotherm for mild steel corrosion inhibition in 1 N HCl, using *Commelina benghalensis* leaves extract.

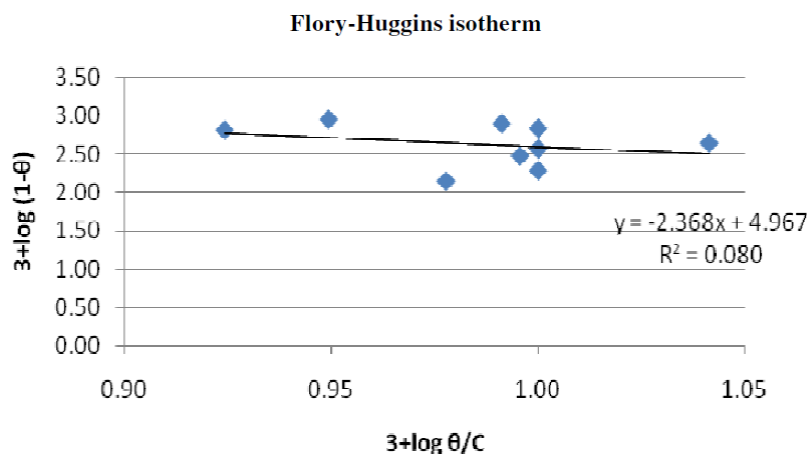
$\Theta$	[C], (ppm)	$\Theta/C$	$3+\log\Theta/C$	$1-\Theta$	$3+\log(1-\Theta)$
0.1727	100	0.0017	0.2374	0.8273	2.9176
0.2455	200	0.0012	0.0889	0.7545	2.8777
0.2909	300	0.0010	0.0734	0.7091	2.8507
0.4455	400	0.0011	0.0467	0.5545	2.7439
0.5545	500	0.0011	0.0450	0.4455	2.6488
0.6727	600	0.0011	0.0497	0.3273	2.5149
0.7727	700	0.0011	0.0429	0.2273	2.3565
0.8455	800	0.0011	0.0240	0.1545	2.1891
0.9340	900	0.0010		0.1000	2.0000

**Table 9.** Flory-Huggins adsorption isotherm for mild steel corrosion inhibition in 0.5 M H<sub>2</sub>SO<sub>4</sub>, using *Commelina benghalensis* leaves extract.

S. no	$\theta$	C	$\theta/C$	$3+\log \theta/C$	$1-\theta$	$3+\log (1-\theta)$
1	0.0893	100	0.0009	0.9493	0.9107	2.9594
2	0.1964	200	0.0010	0.9912	0.8036	2.9050
3	0.3036	300	0.0010	1.0000	0.6964	2.8429
4	0.3393	400	0.0008	0.9242	0.6607	2.8200
5	0.5536	500	0.0011	1.0413	0.4464	2.6498
6	0.6250	600	0.0010	1.0000	0.3750	2.5740
7	0.6964	700	0.0010	0.9956	0.3036	2.4823
8	0.8036	800	0.0010	1.0000	0.1964	2.2932
9	0.8571	900	0.0010	0.9777	0.1429	2.1549



**Figure 6.** Flory-Huggins adsorption isotherm for mild steel corrosion inhibition in 1 N HCl, using *Commelina benghalensis* leaves extract.



**Figure 7.** Flory-Huggins adsorption isotherm for mild steel corrosion inhibition in 0.5 M H<sub>2</sub>SO<sub>4</sub>, using *Commelina benghalensis* leaves extract.

**Table 10.** El-Awady adsorption isotherm for mild steel corrosion inhibition in 1 N HCl, using *Commelina benghalensis* leaves extract.

[C] ppm	2+log C	θ	1-θ	θ/1-θ	2+log (θ/1-θ)
100	4.0000	0.1727	0.8273	0.2088	1.3197
200	4.3010	0.2455	0.7545	0.3253	1.5123
300	4.4771	0.2909	0.7091	0.4103	1.6131
400	4.6021	0.4455	0.5545	0.8033	1.9049
500	4.6990	0.5545	0.4455	1.2449	2.0951
600	4.7782	0.6727	0.3273	2.0556	2.3129
700	4.8451	0.7727	0.2273	3.4000	2.5315
800	4.9031	0.8455	0.1545	5.4706	2.7380
900	4.9542	0.9340	0.0660	14.1515	3.1508

**El-Awady isotherm**

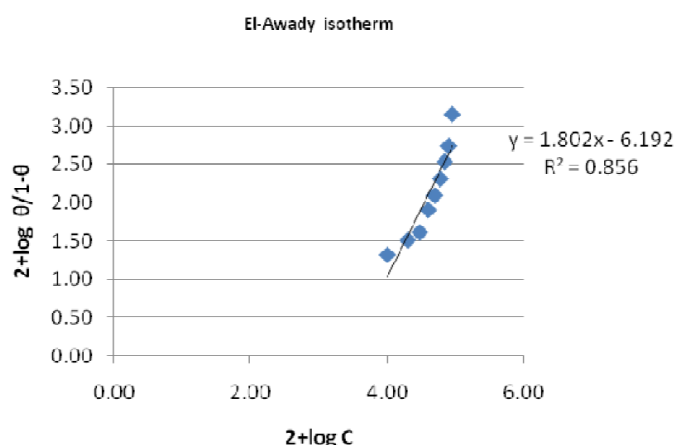
El-Awady isotherm is given by the expression:

$$\log (\theta / 1-\theta)=\log K_{\text {ads }}+y \log C$$

where  $K_{ads}$  is the equilibrium constant of the adsorption process, calculated by the relationship,  $K_{ads} = 1/k$ . Tables 10 and 11, and a linear plot of  $2+\log (\theta/1-\theta)$  against  $2+\log C$  (Figs. 8 and 9), show that El-Awady isotherm was obeyed.

**Table 11.** El-Awady adsorption isotherm for mild steel corrosion inhibition in 0.5 M  $H_2SO_4$ , using *Commelina benghalensis* leaves extract.

[C] ppm	2+log C	$\theta$	1- $\theta$	$\theta/1-\theta$	2+log ( $\theta/1-\theta$ )
100	4.0000	0.0893	0.9107	0.0980	0.9914
200	4.3010	0.1964	0.8036	0.2444	1.3882
300	4.4771	0.3036	0.6964	0.4359	1.6394
400	4.6021	0.3393	0.6607	0.5135	1.7106
500	4.6990	0.5536	0.4464	1.2400	2.0934
600	4.7782	0.6250	0.3750	1.6667	2.2218
700	4.8451	0.6964	0.3036	2.2941	2.3606
800	4.9031	0.8036	0.1964	4.0909	2.6118
900	4.9542	0.8571	0.1429	6.0000	2.7782



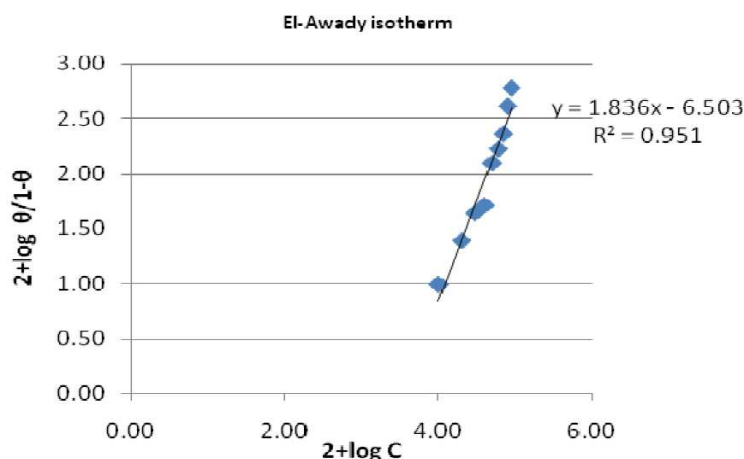
**Figure 8.** El-Awady adsorption isotherm for mild steel corrosion inhibition in 1 N HCl, using *Commelina benghalensis* leaves extract.

### Free energy change

The inhibitor's free energy change of adsorption ( $\Delta G_{ads}^0$ ) onto mild steel surfaces was related to the adsorption constant, according to the equation:

$$\Delta G_{ads}^0 = -2.303 RT \log (K_{ads} \times 55.5)$$

From the results, showed in Tables 12 and 13,  $\Delta G_{ads}^0$  values were found to be negative, and were below the threshold value of  $-40\text{kJ/mol}$ , indicating that the adsorption of *Commelina benghalensis* onto a mild steel surface is spontaneous, and that the physical adsorption mechanism is applicable.



**Figure 9.** El-Awady adsorption isotherm for mild steel corrosion inhibition in 0.5 M  $H_2SO_4$ , using *Commelina benghalensis* leaves extract.

**Table 12.** Adsorption isotherms parameter for mild steel corrosion inhibition in 1 N HCl, using *Commelina benghalensis* leaves extract.

System	Isotherm	$\Delta G^0_{ads}$	Slope	$R^2$	a	1/y
<i>Commelina benghalensis</i> leaves extract	Langmuir	-1481.35	1.780	0.864	-	0.561
	Temkin	-3087.50	0.833	0.889	-	1.200
	Flory-Huggins	-5362.50	-2.882	0.872	-	-0.109
	El-Awady	-1645.50	1.611	0.825	-	0.620

**Table 13.** Adsorption isotherms parameter for mild steel corrosion inhibition in 0.5 M  $H_2SO_4$ , using *Commelina benghalensis* leaves extract.

System	Isotherm	$\Delta G^0_{ads}$	Slope	$R^2$	a	1/y
<i>Commelina benghalensis</i> leaves extract	Langmuir	-5747.61	-7.340	0.951	-	-0.136
	Temkin	-3011.75	-3.383	0.915	-0.399	-0.295
	Flory-Huggins	-2051.29	4.967	0.080	-	0.201
	El-Awady	-1566.80	-6.503	0.951	-	-0.153

### Potentiodynamic polarization studies

The polarization study has been used to investigate the formation of a protective film during the corrosion inhibition study. Whenever there is corrosion inhibition, the corrosion current value decreases [38-42].

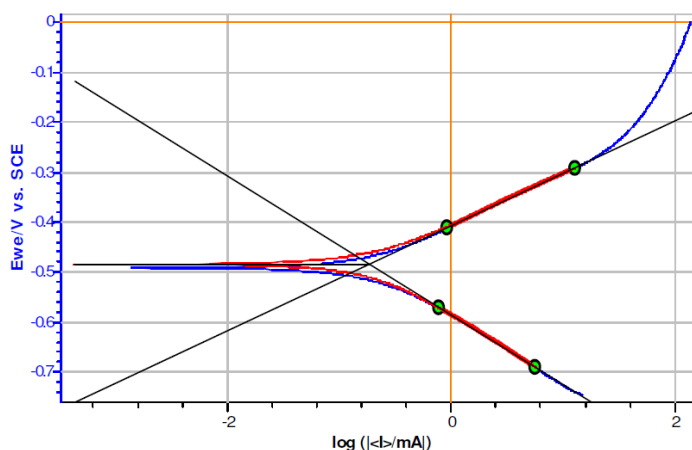
Tables 14 and 15 and Figs. 10 to 15 give the potentiodynamic parameter values, such as corrosion current ( $I_{corr}$ ), corrosion potential ( $E_{corr}$ ) and the cathodic Tafel slopes (bc and ba) for the different concentrations shown in the green inhibitor under study. It can be seen that the active principles present in the extract have been adsorbed onto the metal surface. When the inhibitor concentration is 100 ppm, the system functions as a mixed type inhibitor, because the shift in corrosion potential is within 80 mV. However, at higher concentrations (900 ppm), the system functions as a cathodic type inhibitor, because the shift in the corrosion potential is towards the cathodic side (more negative).

**Table 14.** Corrosion parameters obtained from polarization curves for mild steel in 1 N HCl, in the inhibitor presence and absence.

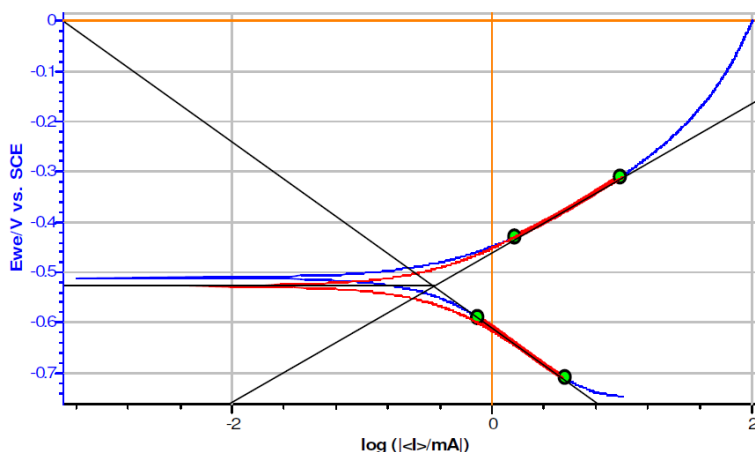
System	[Inhibitor] ppm	$I_{corr}$ mA/cm <sup>2</sup>	$-E_{corr}$ mV vs. SCE	ba. mV dec <sup>-1</sup>	bc. mV dec <sup>-1</sup>	Inhibition efficiency (%)
Blank	-	397.558	484.372	195.4	184.6	-
<i>Commelina benghalensis</i> leaves extract	100	326.066	526.778	157.1	151.5	17.98
	900	21.095	597.158	104.8	122.5	94.69

**Table 15.** Corrosion parameters obtained from polarization curves for mild steel in 0.5 M H<sub>2</sub>SO<sub>4</sub>, in the inhibitor presence and absence.

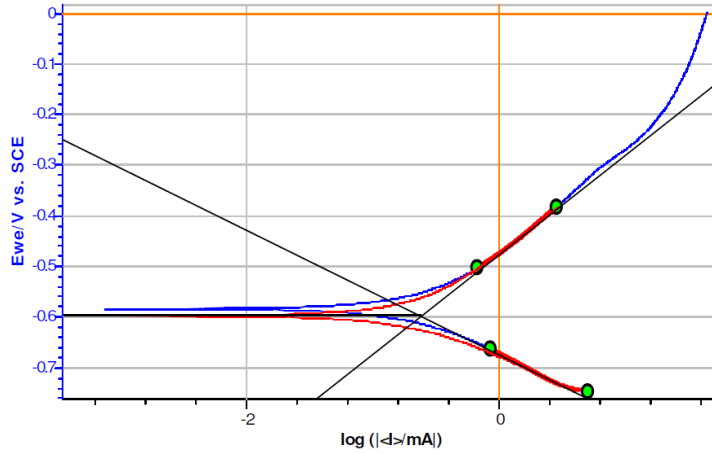
System	[Inhibitor] ppm	$I_{corr}$ mA/cm <sup>2</sup>	$-E_{corr}$ mV vs. SCE	ba mV dec <sup>-1</sup>	bc mV dec <sup>-1</sup>	Inhibition efficiency (%)
Blank	-	249.310	490.236	185.3	198.2	-
<i>Commelina benghalensis</i>	100	218.979	515.727	167.0	186.2	8.15
	900	33.39	575.930	134.1	173.2	86.60



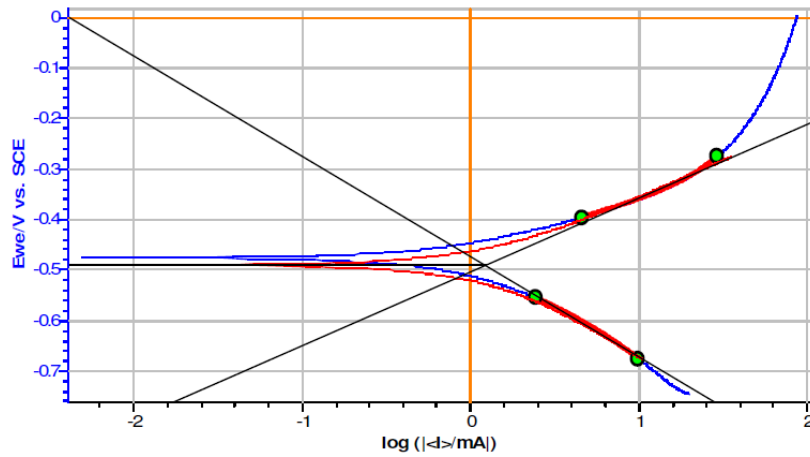
**Figure 10.** Potentiometric polarization curve for 1 N HCl medium.



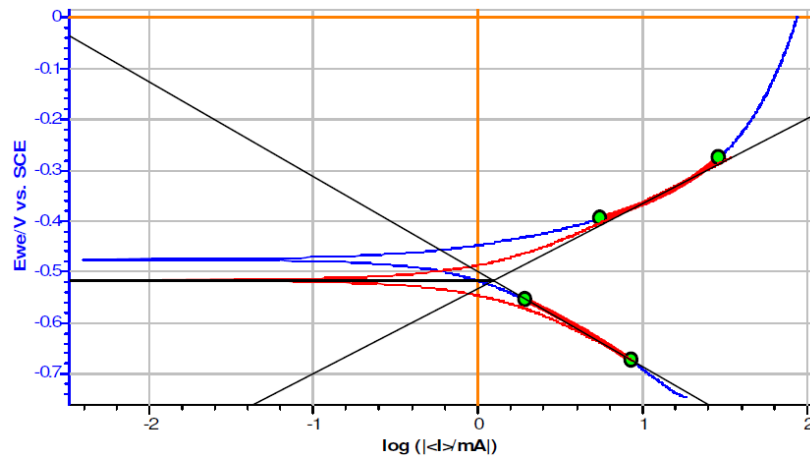
**Figure 11.** Potentiometric polarization curve for 1 N HCl medium, in the presence of 100 ppm concentration of *Commelina benghalensis* leaves extract.



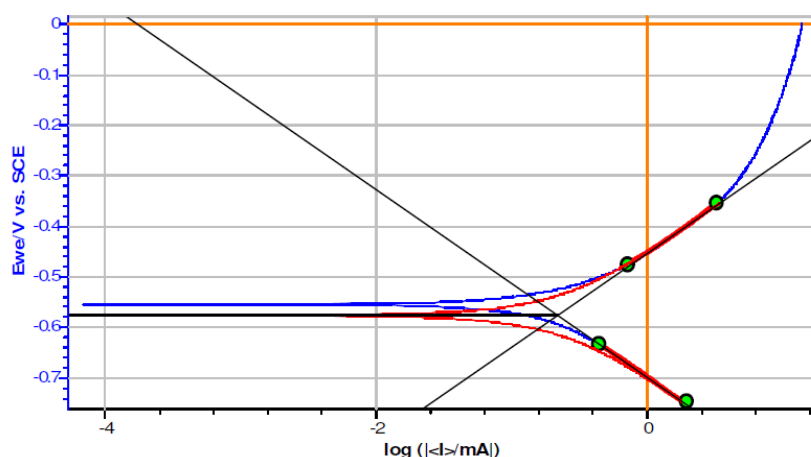
**Figure 12.** Potentiometric polarization curve for 1 N HCl medium, in the presence of 900 ppm concentration of *Commelina benghalensis* leaves extract.



**Figure 13.** Potentiometric polarization curve for mild steel in 0.5 M H<sub>2</sub>SO<sub>4</sub>.



**Figure 14.** Potentiometric polarization curve for 0.5 M H<sub>2</sub>SO<sub>4</sub> medium, in the presence of 100 ppm concentration of *Commelina benghalensis* leaves extract.



**Figure 15.** Potentiometric polarization curve for 0.5 M H<sub>2</sub>SO<sub>4</sub> medium, in the presence of 900 ppm concentration of *Commelina benghalensis* leaves extract.

### AC impedance study

AC impedance spectra have been used to investigate the formation of a protective film on the metal surface, during the corrosion inhibition process. When there is corrosion inhibition, charge transfer resistance value increases, and double layer capacitance decreases [43-48].

Tables 16 and 18 and Figs. 16 to 21 indicate AC impedance curves and values for the inhibitor concentration. The  $R_t$  value increases, and  $C_{dl}$  value decreases with the inhibitor concentration. This implies the formation of a protective film on the metal surface. It can be also seen from Tables 17 and 19 that there is a close agreement between the inhibition efficiencies values obtained from weight loss measurements, polarization and impedance studies. In the inhibitor presence, a protective film is formed on the metal surface. It prevents electrons transfer from the metal surface to the solution's bulk. That is why there is an increase in the resistance to the electron transfer, and, in the inhibitor presence, the charge transfer resistance increases.

**Table 16.** Corrosion parameters obtained from the impedance study for mild steel in 1 N HCl, in the presence and absence of *Commelina benghalensis* leaves extract.

[Inhibitor] ppm	$R_t$ Ohm cm <sup>2</sup>	$C_{dl}$ , F/cm <sup>2</sup>	Inhibition efficiency (%)
Blank	16.3	0.150	-----
100	17.1	0.123	18.00
900	39.0	0.008	94.61

**Table 17.** Comparison of the inhibition efficiency measured by weight loss, polarization and impedance studies for 1 N HCl.

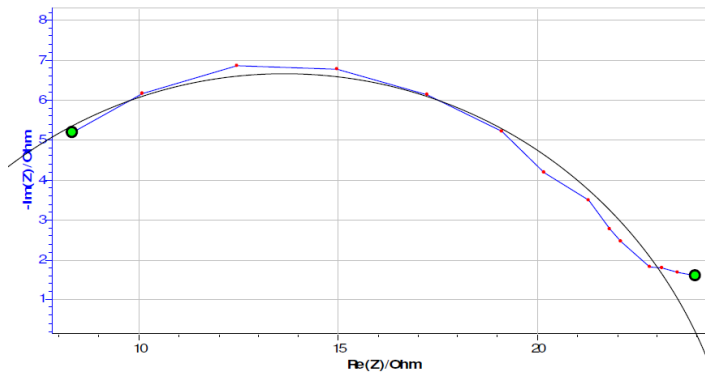
[Inhibitor] ppm	Inhibition efficiency (%)		
	Weight loss study	Polarization study	Impedance study
100	17.27	17.98	18.00
900	93.40	94.69	94.61

**Table 18.** Corrosion parameters obtained from the impedance study for mild steel in 0.5 M H<sub>2</sub>SO<sub>4</sub>, in the presence and absence of *Commelina benghalensis* leaves extract.

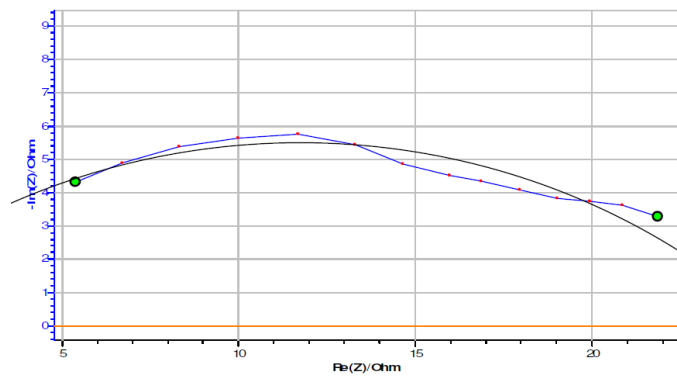
[Inhibitor] ppm	R <sub>t</sub> , Ohm cm <sup>2</sup>	C <sub>dl</sub> , F/cm <sup>2</sup>	Inhibition efficiency (%)
Blank	10.7	0.137	-----
100	11.1	0.126	8.02
900	110.0	0.018	86.86

**Table 19.** Comparison of the inhibition efficiency measured by weight loss, polarization and impedance studies for 0.5 M H<sub>2</sub>SO<sub>4</sub>.

[Inhibitor] ppm	Inhibition efficiency (%)		
	Weight loss study	Polarization study	Impedance study
100	8.92	8.15	8.02
900	85.71	86.60	86.86

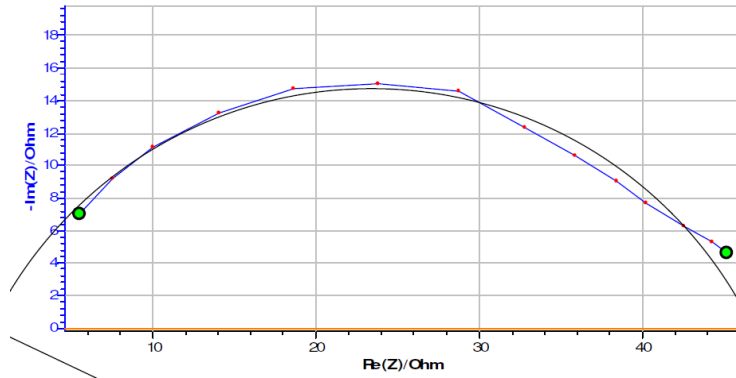


**Figure 16.** Impedance diagram for mild steel in 1 N HCl.

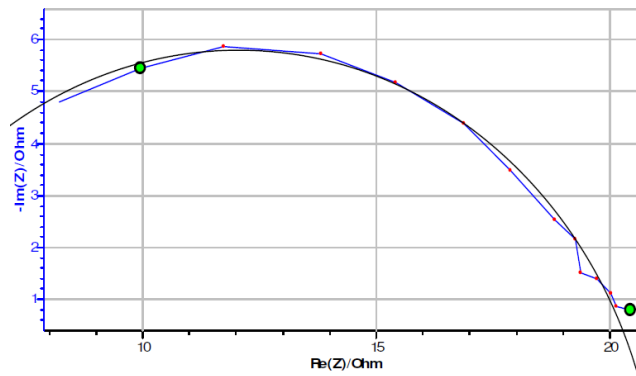


**Figure 17.** Impedance diagram for mild steel in 1 N HCl medium, in the presence of 100 ppm concentration of *Commelina benghalensis* leaves extract.

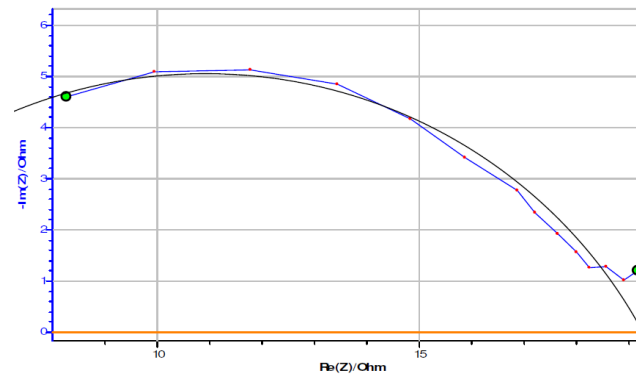




**Figure 18.** Impedance diagram for mild steel in 1 N HCl medium, in the presence of 900 ppm concentration of *Commelina benghalensis* leaves extract.

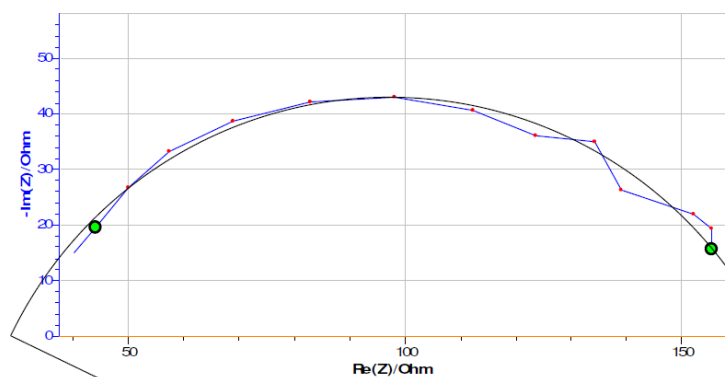


**Figure 19.** Impedance diagram for mild steel in 0.5 M  $H_2SO_4$ .

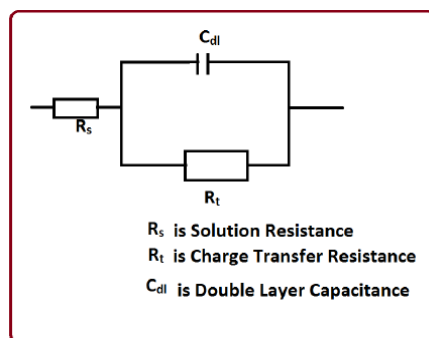


**Figure 20.** Impedance diagram for mild steel in 0.5 M  $H_2SO_4$  medium, in the presence of 100 ppm concentration of *Commelina benghalensis* leaves extract.

It is inferred from the Nyquist plots that the process represents simple corrosion and corrosion resistance. The equivalent circuit diagram for such a system is shown in Fig. 22.



**Figure 21.** Impedance diagram for mild steel in 0.5 M H<sub>2</sub>SO<sub>4</sub> medium, in the presence of 900 ppm concentration of *Commelina benghalensis* leaves extract.



**Figure 22.** Equivalent circuit diagram for a simple corrosion and corrosion inhibition process.

## Conclusion

The inhibitive influence of *Commelina benghalensis* leaves extract on mild steel corrosion in 1 N HCl and 0.5 M H<sub>2</sub>SO<sub>4</sub> was studied by the weight loss method, polarization and impedance measurements. The inhibition efficiency values determined by these techniques showed close agreement. The corrosion decreased with the increasing addition of *Commelina benghalensis* leaves extract, probably due to the progressive adsorption of the inhibitor onto the metal surface. The maximum inhibition efficiency was found to be 93.40% in 1 N HCl, and 85.71% in 0.5 M H<sub>2</sub>SO<sub>4</sub>.

1. Mild steel corrosion in 1 N HCl and 0.5 M H<sub>2</sub>SO<sub>4</sub> solutions was inhibited by the addition of *Commelina benghalensis* leaves extract.
2. The percentage of inhibition efficiency increased with higher inhibitor concentrations.
3. The corrosion inhibition of *Commelina benghalensis* leaves extract is attributed to the adsorption of any of the phyto-chemical components present in the inhibitor onto the mild steel surface. The adsorption was assumed to arise from the  $\pi$ -bond of the components onto the mild steel surface.
4. The values obtained from the weight loss technique for the studied inhibitor fit into the Langmuir and Temkin adsorption isotherms and the kinetic thermodynamic model. The free energy values for the adsorption

processes indicate both physisorption and chemisorption (comprehensive adsorption) of the studied *Commelina benghalensis* leaves extract onto the mild steel surface.

5. AC impedance studies reveal that a protective film was formed on the metal surface.
6. The results suggest that *Commelina benghalensis* leaves extract is a great eco-friendly inhibitor.
7. The corrosion inhibition mechanism occurred by adsorption of the active principles of the extract ingredients onto the metal surface.

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### References

1. Chraibi M, Benbrahim KF, Elmsellem H, et al. J Mater Environ Sci. 2017;8:972.
2. Ajeigbe SO, Basar N, Maarof H, et al. J Mater Environ Sci. 2017;8:2040.
3. Sivakumar PR, Srikanth AP. Asian J. Chem. 2017;29:274.
4. Kalla A, Benahmed M, Djeddi N, et al. Int . Ind Chem. 2016;7:419.
5. Singh A, Ahamad I, Quraishi MA. Arab J Chem. 2016;9:S1584.
6. Júnior JMF, Silva MG, Monteiro JA, et al. Int J Electrochem Sci. 2016;11:3862.
7. Etteyeb N, Nóvoa XR. Corros Sci. 2016;112:471.
8. Ituen E, James A, Akaranta O, et al. Chin J Chem Eng. 2016;24:1442.
9. dos Santos JEP, Barroso APRN, Parente MMV, et al. Rev Materia. 2016;21:1045.
10. Umoren SA. J Adhes Sci Technol. 2016;30:1858.
11. Anthony N, Malarvizhi E, Maheshwari P, et al. Indian J Chem Technol. 2014;11:346.
12. Rajendran S, Amalraj AJ, Joice MJ, et al. Corros Rev. 2014;22:233.
13. Rajendran S, Ganga Sri V, Arockiaselvi J, et al. Bull Electrochem. 2005;21:367.
14. Rajendran S, Muthulakshmi S, Rajeswari R, et al. J Electrochem Soc. 2005;54:50.
15. Priya SL, Chitra A, Rajendran S, et al. Surf Eng. 2005;21:229.
16. Shanmugapriya S, Rajalakshmi T, Amal Raj AJ. Corrosion. 2005;61:685.
17. Anuradha K, Vimala R, Narayanasamy B, et al. Chem Eng Comm. 2008;195:352.
18. Selvi JA, Gangasree V, Rajendran S. Port Electrochim Acta. 2008;27:1.
19. Rajendran S, Jeyasundari J, Selvi JA, et al. Port Electrochim Acta. 2009;27:153.
20. Rajendran S, Paulraj J, Rengan P, et al. J Dent Oral Hyg. 2009;1:1.
21. Rajendran S, Manivannan M. Zastit Mater. 2009;50:131.
22. Rajendran S, Agasta M, Devi RB, et al. Zastit Mater. 2009;50:77.
23. Rajendran S, Sumithira P, Devi BS, et al. Zastit Mater. 2009;50:223.

24. Rajendran S, Uma V, Krishnaveni A, et al. Arab J Sci Eng. 2009;34:47.
25. Anthony N, Sherina HB, Rajendran S. Int J Eng Sci Tech. 2009;2:2774.
26. Rajendran S, Sumithra P, Devi BS, et al. Zastit Mater. 2009;5:223.
27. Sangeetha M, Rajendran S, Megala TSM, et al. Zastit Mater. 2009;52:35.
28. Devi BS, Rajendran S. Int J Electrochem Sci Tech. 2009;1:79.
29. Sirbharathy V, Rajendran S, Sathyabama J. Int J Chem Sci Tech. 2011;1:108.
30. Sangeetha M, Rajendran S, Bama JS, et al. Port Electrochim Acta. 2011;29:429.
31. Sangeetha M, Sathyabama J, Rajendran S, et al. Nat Prod Plant Res. 2012;2:601.
32. Florence JF, Rajendran S, Srinivasan KV. Electroplating Finishing. 2012;31:1.
33. Sirbharathy V, Rajendran S. Chem Sci Rev Lett. 2012;1:25.
34. Rajam K, Rajendran S, Banu NN. J Chem. 2012;11: 521951.
35. Johnsirani V, Sathiyabama J, Rajendran S, et al. ISRN Corros. 2012;9:574321.
36. Sriharathy V, and Rajendran S. ISRN Corros, Article ID 370802, 2012, 7.
37. Tadesse S, Ganesan K, Nair SKP, et al. Int J Pharm, Chem Biol Sci. 2016;6:103.
38. Shanmugapriya S, Rajendran S, Prabakar P, et al. Int J Nano Corr Sci Eng. 2016;3:144.
39. Anusuya S, Meenakshi SD. Int J Nano Corr Sci Eng. 2016;3:204.
40. Abinaya M, Meenakshi SD. Int J Nano Corr Sci Eng. 2016;3:216.
41. Rajendran S, Krishnaveni A, Muthukumar G. Int J Nano Corr Sci Eng. 2016;3:254.
42. Thangakani JA, Rajendran S, Sathiyabama J. Int J Nano Corr Sci Eng. 2016;3:280.
43. Lavanyaa M, Suganya M. Int J Nano Corr Sci Eng. 2016;3:88.
44. Vanitha AJ, Raja AS, Rajendran S. Int J Nano Corr Sci Eng. 2016;3:223.
45. Anandan A, Rajendran S, Sathiyabama J, et al. Int J Nano Corr Sci Eng. 2016;3:9.
46. Nithya K, Meenakshi SD. Int J Nano Corr Sci Eng. 2016;3:44.
47. Devi PN, Sathiyabama J, Rajendran S, et al. Int J Nano Corr Sci Eng. 2016;3:56.
48. Madhumitha S, Priyadarshini V, Sheela A, et al. Int J Nano Corr Sci Eng. 2016;3:80.