ASSESSMENT OF *Passiflora vitifolia* LEAVES EXTRACT AS A POTENTIAL INHIBITOR FOR MILD STEEL ACID CORROSION

R. Thilagavathi, A. Prithiba and R. Rajalakshmi*

Department of Chemistry, Avinashilingam Institute for Home Science and Higher Education for Women, Coimbatore-641043, Tamil Nadu, India

*E-mail: rajivardhan@gmail.com

ABSTRACT

The present study is focused on the application of *Passiflora vitifolia* leaves to extract as an inhibitor for corrosion of mild steel in 1M hydrochloric acid solution. Mass loss measurements reflect an inhibition efficiency of 97.1% at 12h of immersion and electrochemical measurement afforded a protection efficiency of 89.8% at 0.7% concentration of the inhibitor. The adsorption of the *Passiflora vitifolia* leaves extract onto the mild steel surface obeyed Langmuir adsorption isotherm. Thermodynamic parameters were also calculated and discussed. Potentiodynamic polarization implied that *Passiflora vitifolia* leaves extract in 1M HCl behaved as mixed type inhibitor. The Nyquist plots reflected an increase in charge transfer resistance and decreased double layer capacitance with an increase in additive dosage. Surface morphological studies further implied the effectiveness of the inhibitor under study.

Keywords: *Passiflora vitifolia* Leaves, Corrosion Inhibition, Mass Loss, Electrochemical Study, Surface Morphology.

INTRODUCTION

The serious consequences of corrosion, a naturally occurring phenomenon, tend to jeopardize the safety of metals and inhibit technology progress as metals place a vital role in the world economy. Among the acidic solutions, hydrochloric acid is the commonly used agent in industries to remove rust from the mild steel surface. This tends to considerable damage to the metal surface. Literature reports that several synthetic compounds have shown good anticorrosive activity but these inhibitors may cause reversible and irreversible damage to organ system viz., kidneys or liver, or to disturb a biochemical process or to disturb an enzyme system at some site in the body. The toxicity may manifest either during the synthesis of the compound or during its applications. These toxic effects have led to the use of natural products as anti-corrosion agents which are eco-friendly and harmless. For this reason, plant extracts have attracted the attention of researchers as ecologically acceptable corrosion inhibitors. Hence the present work is proposed to evaluate the inhibitive nature of the selected plant extracts as an inhibitor for mild steel corrosion in 1M HCl. Our research team has carried out several studies successfully in utilizing natural products for mild steel corrosion inhibitor.

*Passiflora* species have been widely investigated for the presence of bioactive compounds in all the plant organs, including leaves, flowers, fruits and seeds, and a series of flavonoids, glycosides, alkaloids, and phenols have been reported. Based on the above literature survey, the present study focuses on corrosion mitigation of MS in 1M HCl using *Passiflora vitifolia* leaves extract.

EXPERIMENTAL

Material Selection

The selected MS specimen for the present study comprised of (weight %) - carbon 0.019%, manganese 0.352%, silicon 0.049%, phosphorus 0.019%, sulphur 0.013%, chromium 0.010%, molybdenum 0.008%,...
nickel 0.010%, Copper 0.026% and iron 99.33%. The coupons having dimension of 1 cm$^2$ were used for electrochemical technique and 1 x 5 cm$^2$ was used for mass loss method. The specimens were prepared as per the procedure reported [ASTM G1-03] $^{30}$.

**Preparation of Plant Extract**
*Passiflora vitifolia* leaves (PAVL) were collected from the Maruthamalai area in Coimbatore and shade dried. The plant specimen was authenticated in Botanical Survey of India (BSI/SRC/5/23/2012-2018/Tech/1742). 500 mL of 1molar hydrochloric acid was added with 25g of powdered leaves and refluxed for 3 hours. The extract was kept overnight for cooling, then filtered and made up to 500 mL (5% extract).

**Experimental Techniques**

**Mass Loss Method**
Pre-weighed mild steel sample was immersed in 100 mL of the blank/inhibitor solution for a prearranged time period as per ASTM G 1-2$^{31}$. After a predetermined test period, the MS specimens were washed, dried and reweighed.

**Electrochemical Measurements**
Potentiodynamic measurement and EIS measurements were performed using a Frequency response analyzer (Biologic model V10.23) in a three-electrode setup. Platinum, saturated calomel electrode and mild steel were employed as an auxiliary, reference and working electrodes respectively. For polarisation studies, the measurements were undertaken by applying a potential range of -0.1v to -1mV and scan rate of 2mVs$^{-1}$. For EIS studies, a frequency range of 20 kHz to 0.1Hz at the OCP of the working electrode was applied.

**Surface Analytical Techniques**

**FT-IR Spectroscopy**
FTIR was recorded using Shimadzu IR Affinity-1S Fourier Transform Infrared Spectrometer which extended from 4000 and 400 cm$^{-1}$. It was performed to identify the functional groups present in the plant extract, while that of the corrosion products was done to ascertain the interaction between the metal and the extract.

**UV-Visible Spectrophotometer**
The investigated plants and corrosion product were subjected to UV-vis spectrophotometric characterization over 200-800 nm using AU-2701 UV-Vis Double beam spectrophotometer.

**3D Optical Profilometer**
Surface profiles and pores were studied using 3D Optical Profilometer. The MS specimen exposed in 1M hydrochloric acid solution with and without the addition of PAVL extract for 3h was examined by Zeta 3D Profiler. The specimens were mounted the sample holder and 3D images were taken with 100x magnified surface using operating program on the computer.

**SEM and EDX**
The surface topography of the metal surface in the absence and presence of PAVL extract was examined by SEM (JEOL MODEL JSM 6360). The nature of the formed protective film on MS surface was found out by EDX spectra.

**Characterization of the Investigated Inhibitor**

**Phytochemical Screening of PAVL Extract**
HPTLC, FT-IR and UV techniques were performed to characterize the PAVL extract. Standard procedures were used to determine the phytochemical constituents present in the plant extract $^{32}$.
HPTLC Technique
CAMAG HPTLC instrument with Win CATS version 1.3.4 software was performed to analyze phytochemical components and its derivatives present in the leaf extract.

FTIR Spectroscopy
The inhibitor was characterized by FT-IR spectroscopy for identification of active functional groups. The FT-IR study was carried out by using Shimadzu IR Affinity-1S Fourier Transform Infrared Spectrometer.

UV-Visible Spectrophotometer
UV-Vis spectral studies of the investigated plant extract were done using AU-2701 UV-Vis Double beam spectrophotometer.

RESULTS AND DISCUSSION
Mass Loss Study
The effect of PAVL extract on the inhibition of MS corrosion was tested by mass loss measurement. Table-1, Fig.-1 indicated the variation of corrosion rate (CR) and inhibition efficiency (IE) with an increase in the concentration of PAVL extract. The corrosion rate decreased with increased concentration of the PAVL extract. The IE increased from 92.3% to 97.1% for 0.7% concentration of PAVL extract.

| Conc (v/v%) | 1/2h | 1h | 3h | 6h | 12h | 24h |
|------------|------|----|----|----|-----|-----|
| CR (mpy)   | IE (%) | CR (mpy) | IE (%) | CR (mpy) | IE (%) | CR (mpy) | IE (%) |
| Blank      | 4538 | - | 4405 | - | 2706 | - | 2625 | - | 2123 | - | 1707 | - |
| 0.1        | 623  | 86.3 | 431 | 90.2 | 269 | 90.1 | 195 | 92.6 | 148 | 93.0 | 157 | 90.8 |
| 0.2        | 537  | 88.2 | 320 | 92.7 | 191 | 93.0 | 142 | 94.6 | 96  | 95.5 | 108 | 93.7 |
| 0.3        | 469  | 89.7 | 303 | 93.1 | 159 | 94.1 | 126 | 95.2 | 78  | 96.3 | 103 | 94.0 |
| 0.4        | 409  | 91.0 | 282 | 93.6 | 137 | 95.0 | 109 | 95.9 | 72  | 96.6 | 92  | 94.6 |
| 0.5        | 384  | 91.5 | 256 | 94.2 | 131 | 95.2 | 94  | 96.4 | 64  | 97.0 | 91  | 94.7 |
| 0.6        | 375  | 91.7 | 252 | 94.3 | 117 | 95.7 | 92  | 96.5 | 64  | 97.0 | 82  | 95.2 |
| 0.7        | 350  | 92.3 | 235 | 94.7 | 112 | 95.8 | 82  | 96.9 | 63  | 97.1 | 79  | 95.4 |

An insight into the stability of PAVL extract with time may be gained by studying the effect of the extract on MS specimen for various time of immersion. The results evaluated for the variation of mass loss with

![Figure 1: Inhibition efficiency of PAVL on corrosion of MS in 1M HCl](image-url)
exposure time for the MS specimen immersed in 1M HCl with and without inhibitor are presented in Table-1. A maximum IE of 97.1% was maintained till 12 h and then a slight decline was observed. But the inhibition efficiency was found to stabilize at 24h to afford an efficiency of 95.4% demonstrating the effectiveness of the inhibitor at longer periods of immersion.

**Effect of Temperature**

The effect of temperature on inhibition efficiency of PAVL extract in an acidic environment is depicted in Fig.-2. The corrosion rate decreased by increased in PAVL extract at any given temperature. Analyzing the temperature effect of PAVL, it can be noticed that the IE increased with increase in temperature up to 333K giving rise to 94.0% and then a slight decrease was noticed from 333K to 353K which then stabilized to 90.8% at 353K. It was due to desorption of the adsorbed PAVL molecules at elevated temperatures.

Table-2: Effect of Temperature on MS Specimen in Acidic Medium with and without Addition of PAVL Extract

| Conc (%v/v) | 303K | 313K | 323K | 333K | 343K | 353K |
|------------|------|------|------|------|------|------|
| Conc (mpy) | CR   | IE   | CR   | IE   | CR   | IE   |
| Blank      | 4538 | 6585 | -    | -    | 14884| 20130| 29819|
| 0.1        | 623  | 86.3 | 904  | 86.3 | 1152 | 89.5 | 2337 | 88.7 |
| 0.2        | 537  | 88.2 | 768  | 88.3 | 1152 | 88.0 | 2712 | 86.5 |
| 0.3        | 469  | 89.7 | 631  | 90.4 | 1007 | 89.5 | 2337 | 88.4 |
| 0.4        | 409  | 91.0 | 546  | 91.7 | 904  | 90.6 | 1970 | 90.2 |
| 0.5        | 384  | 91.5 | 546  | 91.7 | 836  | 91.3 | 1732 | 91.4 |
| 0.6        | 375  | 91.7 | 478  | 92.7 | 725  | 92.4 | 1612 | 92.0 |
| 0.7        | 350  | 92.3 | 461  | 93.0 | 682  | 92.9 | 1552 | 92.3 |

| Conc (v/v%) | CR (mpy) | IE (%) | Conc (v/v%) | CR (mpy) | IE (%) |
|-------------|-----------|--------|-------------|-----------|--------|
| 0.1         | 4538      | 6585   | 0.2         | 537       | 88.2   |
| 0.3         | 469       | 89.7   | 0.4         | 409       | 91.0   |
| 0.5         | 384       | 91.5   | 0.6         | 375       | 91.7   |
| 0.7         | 350       | 92.3   | 0.1         | 623       | 86.3   |

**Energy of Activation**

The effect of corrosion rate on temperature can be regarded as an Arrhenius – type process, the rate of which is given by

\[
\text{Log CR}=\text{logA}-\frac{E_a}{2.303R}
\]

Where, CR is the corrosion rate of MS, A is Arrhenius constant, \(E_a\) is the activation energy, \(R\) is the gas constant and \(T\) is the temperature.
From the slope of Arrhenius plot (log CR vs. 1/T - Fig.-3a) $E_a$ values were obtained. Inspection of Table-3 revealed that the $E_a$ value increased with increase in PAVL extract. The $E_a$ values for the investigated inhibitor were found to be higher than the blank implying the increased energy barrier of the corrosion process due to the presence of PAVL extract. This was due to an electrostatic nature of the adsorbed inhibitor on the MS surface$^{34, 35}$. This inspection further supported the physical adsorption mechanism as reported. Several investigators observed lower the values of $E_a$ with and without the addition of an inhibitor solution were denoting the chemical adsorption mechanism, whereas the reverse statement reflected a physical adsorption mechanism$^{36, 37}$.

**Langmuir Adsorption Model for Investigated Inhibitor**

Langmuir adsorption equation relates to the degree of surface coverage to the concentration of inhibitor according to equation (2):

$$\log \left( \frac{C}{\theta} \right) = \log C - \log K$$  \hspace{1cm} (2)

A plot of log (0/1-0) against log C from mass loss data obtained for investigated inhibitor yielded a straight line as represented in Fig.-4. The slope was found to deviate from unity and it was due to the interaction of the adsorbed molecules on the MS surface$^{38}$. Hence in the current study, the PAVL molecule can be absorbed on the anodic and cathodic area and this may be the reason for deviation from the unit gradient.
The entropy of Activation and Enthalpy of Activation
To calculate $\Delta H_a$ and $\Delta S_a$ for the corrosion process, the Arrhenius equation is used:

$$CR/T = \frac{R}{Nh} \exp \left( \frac{\Delta S_a}{R} \right) \exp \left( \frac{\Delta H_a}{RT} \right)$$ (3)

Where $h$ is the Planck’s constant, $N$ is the Avogadro’s number, $\Delta S_a$ is the entropy of activation, $T$ is the absolute temperature and $R$ is the universal gas constant. The relationship between log (CR/T) against $1/T$ for MS acid corrosion in the absence and presence of PAVL extract is depicted in Figure 3b. Straight lines were obtained from the slope of $(-\Delta H_a/2.303R)$ and an intercept of $(\log R/Nh + \Delta S_a/2.303R)$. The values of $\Delta H_a$ and $\Delta S_a$ were obtained from the slope and intercept of a straight line. From Table-3 it may be noticed that the values of $E_a$ and $\Delta H_a$ were close to each other as expected from transition state theory and were also found to vary in a similar manner with PAVL extract but however, the value of $\Delta H_a$ was greater than that of $E_a$. The positive values of $\Delta H_a$ revealed that the reaction was endothermic nature of the steel dissolution process and this confirmed that the dissolution of MS was difficult.

| S. No. | Conc (v/v) | $E_a$(KJ/mol) | $\Delta H_a$(KJ/mol) | $\Delta S_a$(J/mol) |
|-------|------------|---------------|----------------------|----------------------|
| 1.    | Blank      | 33.6          | 36.3                 | -18.1                |
| 2.    | 0.1        | 43.5          | 46.2                 | -20.3                |
| 3.    | 0.2        | 38.4          | 41.1                 | -20.5                |
| 4.    | 0.3        | 38.1          | 40.9                 | -22.5                |
| 5.    | 0.4        | 38.9          | 41.6                 | -21.2                |
| 6.    | 0.5        | 37.0          | 39.7                 | -27.8                |
| 7.    | 0.6        | 37.2          | 40.0                 | -27.7                |
| 8.    | 0.7        | 35.9          | 38.6                 | -32.6                |

Thermodynamic Adsorption Parameters
The adsorption behavior of PAVL extract on the MS surface is essential to control corrosion reaction. A plot of $\Delta G_{ads}$ versus $T$ (Figure 5) was linear for MS acid corrosion using PAVL extract in acidic medium. The slopes of the straight lines were equal to $\Delta S_{ads}$ and intercept equal to $\Delta H_{ads}$. The calculated values of $\Delta G_{ads}$, $\Delta H_{ads}$, $\Delta S_{ads}$ at all investigated temperatures (303-353K) for different concentration of PAVL extract were listed in Table-4. The $\Delta G_{ads}$ value was negative which indicated that the adsorption of the PAVL molecules on the MS surface was a spontaneous process. Generally, values of $\Delta G_{ads}$ around -20 kJmol$^{-1}$ or lower are consistent with the electrostatic interaction between the charged molecules and the
charged metal (physisorption); those around -40 kJmol\(^{-1}\) or higher involve charge sharing or transfer from organic molecules to the metal surface to form a coordinate type of bond (chemisorption)\(^{41}\). In the current study, the obtained \(\Delta G_{\text{ads}}\) values were slightly higher negative than -20 kJmol\(^{-1}\) ranging from -19 to -23 kJmol\(^{-1}\) which indicated that the adsorption process was physisorption. The obtained \(\Delta H_{\text{ads}}\) values were negative which identified that the adsorption of PAVL extract on metal surface was exothermic in nature\(^{42}\). The \(\Delta S_{\text{ads}}\) values were negative which reflected that PAVL extract was absorbed well orderly manner on the surface of MS and resulting in a decrease in entropy values. Table 4 revealed that decreased in enthalpy and entropy were the driving force for the adsorption process \(^{43}\).

Table 4: Thermodynamic Parameters for MS Corrosion in 1M HCl in the Absence and Presence of PAVL

| Conc (%v/v) | Free energy of adsorption - \(\Delta G_{\text{ads}}\) (KJ/mol) | \(\Delta S_{\text{ads}}\) J/mol | \(\Delta H_{\text{ads}}\) KJ/mol |
|------------|-------------------------------------------------|----------------------------|----------------------------|
|            | 303K 313K 323K 333K 343K 353K                   |                            |                             |
| Blank      | -     -     -     -     -     -                   | -                         | -                          |
| 0.1        | 21.8  22.5  23.0  22.8  23.6  23.5               | -33.1                     | -12.0                      |
| 0.2        | 20.5  21.2  21.8  22.7  22.8  22.8               | -49.1                     | -5.8                       |
| 0.3        | 19.9  20.7  21.1  21.8  22.1  22.3               | -48.3                     | -5.5                       |
| 0.4        | 19.5  20.4  20.7  21.1  21.8  21.7               | -44.6                     | -6.2                       |
| 0.5        | 19.1  19.8  20.3  21.1  21.6  21.4               | -50.6                     | -4.0                       |
| 0.6        | 18.7  19.7  20.2  20.7  21.3  21.1               | -49.4                     | -4.1                       |
| 0.7        | 18.5  19.4  20.0  21.1  21.0  21.0               | -52.6                     | -2.9                       |

**Fig.-5: Best fit curves of \(-\Delta G^o_{\text{ads}}\) Vs T for MS / PAVL / 1M HCl**

**Electrochemical Measurements**

**Potentiodynamic Polarisation Technique**

Electrochemical studies were used to predict the nature of the inhibitor and to determine the suitable mechanism for the inhibition process. The potentiodynamic polarisation curves reflected a slight change in the anodic (\(b_a\)) and cathodic (\(b_c\)) curves with and without the addition of PAVL extract (Fig.-6 and Table-5). This indicated that the inhibitor was able to suppress both the anodic dissolution and cathodic hydrogen evolution. The results revealed that \(I_{\text{corr}}\) values decreased with increase in the concentration of PAVL extract. Inspection of the values showed that \(I_{\text{corr}}\) values decreased from 212A/cm\(^2\) to 22A/cm\(^2\). This indicated the inhibitor was able to minimize corrosion of MS in the investigated acid medium\(^{44}\). A maximum of 89.8 percentage of inhibition was obtained at 0.7% concentration of PAVL extract. No significant shifts in the \(E_{\text{corr}}\) values were noticed in the presence of the inhibitor. In general, an inhibitor is differentiated as cathodic and anodic inhibitor if the shift is more than 85 mV with respect to that of the blank. In the current study, the \(E_{\text{corr}}\) values were lower than 85 mV than the blank value, indicating the mixed nature of the inhibitor\(^{45}\). \(R_p\) values were found to vary from 159Ohm/cm\(^2\) for the uninhibited solution to 710Ohm/cm\(^2\) for the optimum concentration of the inhibitor (0.7% PAVL). The inhibitor
Passiflora vitifolia leaves extract as a potential inhibitor

R. Thilagavathi et al.

afforded an efficiency of 77.6 percentage. This might be due to the adsorption of active compounds present in PAVL extract onto the metal surface.

![Polarization Plot for MS / PAVL/1M HCl](image)

**Fig.-6: Polarization Plot for MS / PAVL/1M HCl**

Table-5: Electrochemical Polarization Parameters for MS in 1M HCl in the Absence and Presence of Various Concentrations of PAVL Extract

| S. No. | Conc (%v/v) | \(E_{corr}\) (Amp/cm²) | \(b_a\) (mv/dec) | \(b_c\) (mv/dec) | \(I_{corr}\) (Amp/cm²) | IE % | \(R_p\) Ω/cm² | IE (%) |
|-------|-------------|-------------------------|-----------------|-----------------|------------------------|------|-------------|--------|
| 1     | Blank       | -444                    | 78.8            | 115.9           | 212                    | -    | 159         | -      |
| 2     | 0.1         | -438                    | 61.1            | 160.0           | 101                    | 52.4 | 267         | 40.4   |
| 3     | 0.2         | -447                    | 70.8            | 171.4           | 79                     | 62.7 | 276         | 42.4   |
| 4     | 0.3         | -434                    | 61.9            | 172.5           | 70                     | 67.0 | 329         | 51.7   |
| 5     | 0.4         | -430                    | 61.5            | 178.7           | 64                     | 69.8 | 335         | 52.5   |
| 6     | 0.5         | -426                    | 60.9            | 184.1           | 61                     | 71.2 | 350         | 54.6   |
| 7     | 0.6         | -418                    | 55.4            | 174.8           | 49                     | 77.0 | 379         | 58.0   |
| 8     | 0.7         | -401                    | 43.7            | 167.6           | 22                     | 89.8 | 710         | 77.6   |

**Electrochemical Impedance Measurements**

The impedance parameters for MS/1M HCl/PAVL obtained from Nyquist plots (Fig.-7) were collected in Table-6. The observed impedance spectra exhibited a single depressed semicircle whose diameter increased with increase in the concentration of the inhibitor implying a charge transfer process for the corrosion inhibition process. From Fig.-7, it can be noticed that the presence of the inhibitor did not modify the corrosion reaction of MS electrode. The impedance parameters suggested an increase in charge transfer resistance \(R_{ct}\) values with increase in the concentration of the inhibitor. A maximum of 86.3 percentage IE was obtained at 0.7% PAVL. Single capacitive semicircle in Nyquist plot corresponded to a single time constant in the Bode representation. A decrease in \(C_d\) values were noticed with increase in PAVL extract concentration. This might be due to an increase in surface coverage which increased the IE by the PAVL molecules of plant extract.

The nyquist plot was analyzed by a simple circuit model depicted in Fig.-8 which included solution resistance \(R_s\), charge transfer element \(R_{ct}\), constant phase element (CPE) and surface inhomogeneity \(n\) and the values are tabulated (Table-6). Non-linear least square fit of equivalent circuit model as shown in Figure-8 was employed for extracting impedance values from the Nyquist plot.
Table-6: Impedance Values of Mild Steel Acid Corrosion with and without Addition of PAVL in 1M HCl

| Conc (%v/v) | $R_s$ ($\Omega cm^2$) | $Y_o$ ($\mu F/cm^2$) | $n$ | $R_{ct}$ ($\Omega cm^2$) | IE (%) | CPE/Cdl ($\mu F/cm^2$) x10$^{-6}$ | $\theta$ |
|-------------|----------------------|----------------------|-----|-------------------------|--------|----------------------------------|--------|
| Blank       | 2.09                 | 8071                 | 0.84| 19.7                    | -      | 89.4                             | -      |
| 0.1         | 2.81                 | 5763                 | 0.86| 27.6                    | 28.6   | 62.7                             | 0.30   |
| 0.2         | 3.18                 | 4865                 | 0.84| 32.7                    | 39.7   | 50.0                             | 0.44   |
| 0.3         | 2.92                 | 4177                 | 0.90| 38.1                    | 48.2   | 60.7                             | 0.32   |
| 0.4         | 2.85                 | 3778                 | 0.91| 42.2                    | 53.2   | 63.4                             | 0.29   |
| 0.5         | 3.10                 | 2059                 | 0.95| 77.3                    | 74.5   | 54.3                             | 0.39   |
| 0.6         | 3.05                 | 1369                 | 0.98| 116.3                   | 83.0   | 58.8                             | 0.34   |
| 0.7         | 3.64                 | 1103                 | 0.98| 144.4                   | 86.3   | 44.1                             | 0.51   |

Fig.-7: Nyquist and Bode Plots for MS/PAVL/1M HCl

Fig.-8: Equivalent Circuit Model
The double layer capacitance ($C_{dl}$) decreased due to increase in double layer thickness via adsorption of PAVL molecules on MS surface $^{50}$. The $C_{dl}$ values were obtained from equation (4):

$$C_{dl} = Y_0 (\omega_{max})^{n-1}$$  \hspace{1cm} \text{(4)}

Where, $Y_0$ is CPE coefficient, $n$ is CPE exponent, $\omega$ is the angular frequency.

The thickness of the protective layer ($d$) is associated with $C_{dl}$ by the equation (5)

$$C_{dl} = \varepsilon\varepsilon_0 A/d$$  \hspace{1cm} \text{(5)}

Where, $\varepsilon$ is the dielectric constant and $\varepsilon_0$ is the permittivity of the free space and $A$ is surface area of the metal.

When the inhibitor concentration was increased to 0.7% in the corrosive medium, the interface ($\tau$) parameter changes while the capacitance ($C_{dl}$) value decreased signifying that the charge and discharge rates to the metal–solution interface was greatly decreased. This showed that there was an agreement between the amounts of charge that can be stored (i.e. capacitance) and the discharge velocity in the interface ($\tau$).The double layer between the charged MS surface and the solution was considered as an electrical capacitor. The adsorption of PAVL molecules on the MS surface reduced its electrical capacity because they displaced the water molecules and other ions adsorbed on the metal surface. The results of the electrochemical studies were in good agreement with the results of gravimetric studies with slight deviations. This was due to the difference in immersion period of MS in the aggressive media $^{51}$.

Surface Analysis

**UV Spectral Analysis**

![UV-Visible Spectrum of Crude PAVL and Corrosion Product](image)

**Table-7: UV Visible Spectral Values of PAVL Extract and Corrosion Product**

| Inhibitor     | Absorption Band (nm)                  |
|---------------|---------------------------------------|
|               | Crude PAVL Extract | Corrosion Product |
| PAVL extract  | 270, 250, 232, 200, 322, 308           | 238, 242, 375, 397 |

A UV-visible spectrum is used to analyze the formation of the Fe-PAVL complex and it was recorded for 0.7% of PAVL extract in 1M HCl solution before and after immersion of MS for 3h. (Fig.-9). The absorption spectra of PAVL in 1M HCl before immersion of MS reflected two bands in the UV region. These bands were obtained due to $n \rightarrow \pi^*$ and $\pi \rightarrow \pi^*$ transitions. The change in the position of the absorption maximum was indicated the formation of a complex between the phytochemical constituents of the plant extract and MS electrode $^{52,53}$.
FT-IR Analysis

![FT-IR Spectrum](image)

**Table 8: FT-IR Spectral Values of PAVL Crude Plant Extract and Corrosion Product of MS in Acidic Medium**

| Crude Plant Extract | Corrosion Product/Mild Steel/1M HCl | Frequency Assignment |
|---------------------|------------------------------------|----------------------|
| 3977                | 3981                               | O-H stretching       |
| 3318                | 3603                               | N-H/O-H stretching   |
| 2924                | 3116                               | C-H stretching       |
| 2300                | 2314                               | C≡N stretching       |
| 1605                | 1635                               | C=C stretching       |
| 1512                | 1519                               | C=C in ring aromatic |
| 1366                | 1381                               | C-O-C stretching     |
| 1045                | 1072                               | C-O stretching       |
| 633                 | -                                  | C=C bending          |

Figure 10 depicted the IR spectrum of the PAVL extract and corrosion product. Various adsorption modes of the PAVL extract and the protective film formed on the metal surface after immersion in an acidic solution containing 0.7% PAVL extract and their corresponding frequencies were tabulated and analyzed (Table 8). Analysis of the IR spectral data reflected the following points: The band at 3977 cm⁻¹ shifted to 3981 cm⁻¹ in the corrosion product. The adsorption bands at 2314 cm⁻¹ and 1635 cm⁻¹ pertaining to C≡N stretching and C=C stretching were noticed for the corrosion product. Also, a shift from 2924 cm⁻¹ to 3116 cm⁻¹ was noted for C-H bending. A shift from 1366 cm⁻¹ to 1381 cm⁻¹ indicated C-O-C stretch. Some bands, for example, 633 cm⁻¹ corresponding to C=C bending vibration for the aromatic ring system disappeared in the corrosion product. A peak shift from 1045 cm⁻¹ to 1072 cm⁻¹ indicated C-O stretching. The interaction of the phytochemical compounds of PAVL extract with MS surface can be identified by the shift in the adsorption frequency of the inhibitor on MS surface.

**3D Optical Profilometry**

The three-dimensional optical profiler images of the MS surface with and without the addition of PAVL in 1M HCl are depicted in Fig.-11 a, b and c respectively. Surface morphology of MS showed cracks or depth on the MS surface which was due to the attack of acid on it (Fig.-11b). After the addition of optimum concentration of PAVL, (Fig.-11c) the cracks on the surface of the metal were covered homogeneously. Further, the MS surface was found to be smoother by the addition of the inhibitor.
solution. Furthermore, the average roughness on MS sample reduced from 17.28 µm for MS without inhibitor to 7.65 µm for MS with optimum concentration of PAVL. The decrease in $R_a$ value indicated the adsorption of PAVL molecules on the MS surface (Table-9).

![3D Optical Profiler Images for MS/PAVL/1M HCl](image)

**Fig.-11:** 3D Optical Profiler Images for MS/PAVL/1M HCl

**Table-9: $R_a$ and $R_q$ Values for MS/PAVL/1M HCl**

| S. No. | Samples                        | Average Roughness Values, $R_a$ (µm) | Root Mean Square Roughness $R_q$ (µm) |
|--------|--------------------------------|--------------------------------------|--------------------------------------|
| 1.     | Plain MS                       | 2.558                                | 3.140                                |
| 2.     | MS in 1M HCl                   | 17.28                                | 22.35                                |
| 3.     | MS in 1M HCl + 0.7% PAVL Extract | 7.65                                 | 10.16                                |

**SEM Analysis**

Surface morphology of the MS sample before immersion in the test solution and of the MS sample in 1M HCl with and without the addition of PAVL extract after 3h immersion time was recorded and presented in Fig.-12a to 12c. The MS surface immersed in an acid medium in the absence of the extract (Fig.-12b) showed defined corrosion pits reflecting a high degree of localized corrosion attack by the acid medium. The observed corrosion pit was absent in the MS surface in which the PAVL extract was present in the acid solution (Fig.-12c). The surface morphology of the samples immersed in an acidic solution containing the extract (Fig.-12c) was observed to be very much alike in topography to that of the MS before immersion in the acidic solution (Fig.-12b). This suggested the formation of a protective layer.
formed by the extract on the mild steel that tends to serve as a barrier limiting the extent of corrosion attack by the acid solution.

![Fig.-12: SEM Pictures of MS/PAVL/1M HCl](image)

**EDX Analysis**

The chemical composition of the surface of the MS sample with and without the addition of PAVL extract in 1M HCl solution was analyzed by EDX. The EDX spectra are shown in Figures-13a to 13c. The EDX spectra of inhibited samples showed suppressed Fe peaks, when compared with the polished and uninhibited mild steel sample. This suppression of Fe lines was due to the protective film formed on the MS surface. The EDX spectra of inhibited MS sample contained the peaks corresponding to all the elements present in the inhibitor molecules indicating adsorption of PAVL molecules on metal surface.

**Mechanism**

The adsorption of the inhibitor depends on the factors like the chemical composition of the inhibitor, the temperature and electrochemical potential on the metal/solution interface. In fact, the solvent water molecules could also adsorb at a metal/solution interface. Adsorption process can occur through the replacement of solvent molecules from metal surfaces by ions and molecules accumulated near the metal/solution interface. The anions get adsorbed when the metal surface has an excess positive charge in an amount greater than that required to balance the charge corresponding to the applied potential. The phytochemicals present in the PAVL extract are listed in Table 10. The results indicated that the presence of phytoconstituents like steroids, flavonoids, alkaloids, terpenoids, saponins, phenols, glycosides, cholesterol and polyphenols in the leaves and a moderate amount of flavonoids, terpenoids and anthraquinones in the flowers extract. These were corroborated by literature survey.
**Element** | **Wt%** | **At%**  
---|---|---
Si | 0.35 | 0.69  
P | 0.10 | 0.18  
Mn | 2.38 | 2.41  
Fe | 97.17 | 96.72

**Element** | **Wt %** | **At %**  
---|---|---
O | 19.79 | 45.36  
Fe | 79.00 | 53.41  
C | 0.73 | 1.23  
Si | 0.48 | 0.73

---

**S.No.** | **PAVL Extract** | **Phytochemical Constituents**
---|---|---
1. | Carbohydrates | -
2. | Steroids | +
3. | Flavonoids | +
4. | Alkaloids | +
5. | Terpenoids | +
6. | Saponins | +
7. | Phenols | +
8. | Glycosides | +
9. | Cholesterol | +
10. | Proteins | -
11. | Polyphenols | +
12. | Anthraquinones | -

(+) = indicate presence; (-) = indicate absence

---

**Fig.-13: EDX images of MS/PAVL/1M HCl**

**Table-10: Preliminary Phytochemical Screening of the Crude Extract**
Passiflora vitifolia LEAVES EXTRACT AS A POTENTIAL INHIBITOR

R. Thilagavathi et al.

UV-vis Spectrophotometer

The UV-visible spectrum of PAVL crude extract is shown in Fig.-14 and their corresponding values were listed in Table-11. UV-visible spectroscopy is an essential technique to analyze the nature of phytochemical constituents present in the leaves extract. The results revealed that the major peak appeared at 232nm and 250nm. Evidently, a peak at 232 nm was obtained by the electron transition of $n \rightarrow \sigma^*$ in N and O atoms. The result inferred that the most important constitution compounds in PAVL extract were flavonoids, alkaloid and other compounds containing N or O atoms.

FT-IR Analysis

The FTIR spectrum of PAVL is depicted in Fig.-15 and the observed frequencies are tabulated (Table-12). A strong band at 3318 cm$^{-1}$ was attributed to N-H/OH stretching. An absorption band related to –CH$_2$ stretching was noticed at 2924 cm$^{-1}$. The strong band at 1605 cm$^{-1}$ corresponds to C=C stretching. Peaks at 1512 cm$^{-1}$, 1045 cm$^{-1}$ indicated C=C stretching and C-O aromatic system. The peak at 633 cm$^{-1}$ denoted C=C bending vibration of the aromatic ring system. FTIR spectral values revealed that PAVL extract contained the functional groups such as C-N, C-O, O-H, N-H, C=N linkages and aromatic rings.
Table-12: FT-IR Spectral Details of PAVL Crude Plant Extract

| Observed IR Frequency (cm⁻¹) and Peak Assignment |
|-----------------------------------------------|
| Crude Plant Extract                           |
| Frequency Assignment                          |
| 3977 O-H stretching                            |
| 3318 N-H/O-H stretching                        |
| 2924 C-H stretching                            |
| 2300 -C≡N stretching                           |
| 1605 C≡C stretching                            |
| 1512 C-C aromatic ring                         |
| 1366 C-O-C stretching                          |
| 1045 C-O stretching                            |
| 633 C=C bending                                |

HPTLC Analysis

The ethanolic extract of PAVL extract indicated the presence of ten different types of flavonoids with $R_f$ values 0.06, 0.16, 0.29, 0.39, 0.56, 0.83, 0.91, 0.96, 0.98 and 0.80. It was confirmed that the presence of eleven different types of alkaloids with $R_f$ values 0.02, 0.07, 0.17, 0.29, 0.34, 0.42, 0.45, 0.53, 0.87, 0.93 and 0.41. The PAVL extract showed nine different types of steroids with $R_f$ values 0.09, 0.17, 0.30, 0.42, 0.70, 0.88, 0.94, 0.98 and 0.41. These results inferred that the extract showed flavonoids, alkaloids and steroids in it (Fig.-16). These analytical techniques indicated the presence of several functional groups namely O-H, C-H, -C≡N, C=C, C-O-C, C-O etc that can act as an inhibitor. Aromatic compounds (which contain the p electrons) undergo particularly strong adsorption on many electrode surfaces. The bonding can occur between the metal surface and the aromatic ring.
The exact nature of the interactions between a metal surface and an aromatic molecule depends on the relative coordinating strength towards the given metal of the particular group present\textsuperscript{61}. In aqueous acidic solution, PAVL exists either as a neutral molecule or as protonated molecules (cation). PAVL molecules might be adsorbed on the metal and acid solution interface in the following ways: (a) Electrostatic interaction of protonated molecules with already adsorbed chloride ions in the surface of MS (b) donor-acceptor interactions between the vacant d orbital of surface iron atoms and p-electron of the aromatic ring (c) interaction of unshared electron pairs of heteroatom with vacant d-orbital of iron surface atoms.

**CONCLUSION**

PAVL acted as an effective inhibitor for MS in 1M HCl solution and the extent of inhibition efficiency was directly proportional to the concentration of the inhibitor. The inhibition efficiency increased with increased inhibitor concentration and the maximum IE of 97.1\% was observed at 0.7 \% concentration of PAVL extract. The adsorption of the inhibitor followed Langmuir adsorption isotherm. From $I_{corr}$ values maximum inhibition efficiency, 89.8\% was obtained at 0.7 \% concentration of PAVL extract. Nyquist plots reflected that the charge transfer resistance values increased with increase in the concentration of PAVL extract. The inhibition efficiency (IE \%) measured from $R_{ct}$ values were found to be maximum at 0.7 \% concentration of PAVL extract. In the current investigation, the PAVL extract played a major role in reducing the metal dissolution as well as hydrogen evolution and protected the MS surface from corrosion.

**ACKNOWLEDGMENT**

The authors would like to thank the authorities of Avinashilingam Institute for Home Science and Higher Education for Women, Coimbatore-641043 and Tamilnadu, India for providing necessary facilities for carrying out this study.

**REFERENCES**

1. A.K. Mishra, N. Ebrahimi, D.W. Shoesmith and P.E. Manning, *NACE Corrosion International Conference and Expo*, 7680(2016)
2. A. Ostovari, S. M. Hoseinieh, M. Peikari, S. R. Shadizadeh and S. J. Hashemi, *Corros. Sci.*, 51, 1935(2009), DOI: 10.1016/j.corsci.2009.05.024
3. M. Abdullah, A.Y. El-Etre, M.G. Soliman and E.M. Mabrouk, *Anti-Corrosion Methods and Materials*, 53, 118(2006), DOI: 10.1108/00035590610650820
4. P.R. Vijayalakshmi, R. Rajalakshmi and S. Subhashini, *E-Journal of Chemistry*, 7, 1055(2010)
5. R. Rajalakshmi, S. Subhashini, S. Leelavathi and R. Femina Mary, *Oriental Journal of Chemistry*, 24, 1085 (2008)
6. P.R. Vijayalakshmi, R. Rajalakshmi and S. Subhashini, *Asian Journal of Chemistry*, 22, 4537 (2010)
7. S. Subhashini, R. Rajalakshmi, A. Prithiba and A. Mathina, *E-Journal of Chemistry*, 7, 1133 (2010)
8. P.R. Vijalakshmi, R. Rajalakshmi and S. Subhashini, *Portugaliae Electrochimica Acta*, 29, 9 (2011), DOI: 10.4152/pea.201101009
9. R. Rajalakshmi and S. Safina, *E-Journal of Chemistry*, 9, 1632 (2012)
10. R. Rajalakshmi and S. Safina, *Asian Journal of Chemistry*, 24, 4401 (2012)
11. A. Mathina and R. Rajalakshmi, *Rasayan Journal of Chemistry*, 9(1), 56 (2016).
12. S. Leelavathi and R. Rajalakshmi, *NACE Corrosion Conference and Expo*, 1-15(2013)
13. S. Leelavathi and R. Rajalakshmi, *Journal of Materials and Environmental Science*, 4, 625 (2013)
14. A. Prithiba and R. Rajalakshmi, *Hindawi Publishing Corporation*, *International Journal of Metals*, 9 (2016), DOI: 10.1155/2016/8579429
15. A. Prithiba and R. Rajalakshmi, *Chem .Sci. Rev. Lett.*, 5, 83 (2016)
16. D. Chassagne, J. Crouzet, C. L Bayonove and R.L. Baumes, *J. Agric. Food Chem.*, 46, 4352 (1998), DOI: 10.1021/jf980416k
17. A. Bendini, L. Cerretani, L. Pizzolante, T. Gallina-Toschi, F. Guzzo, S. Ceoldo, A. M. Marconi, F. Andreatta and M. Levi, *Eur. Food Res. Technol.*, 223, 102 (2006), DOI: 10.1007/s00217-005-0150-7
18. A.S. Patil, *J. Medic. Plants Res.*, 4, 1496 (2010), DOI: 10.5897/JMPR10.061
19. M. L. Zeraik and J. H. Yariwake, *Microchemical Journal*, 96, 86 (2010), DOI: 10.1016/j.micr.2010.02.003
20. S. S. Patel, H. Soni, K. Mishra and A. K. Singhai, *Int. J. Res. Phytochem. Pharmacol.*, 1, 1 (2011)
21. S. M. Zucolotto, C. Fagundes, F. H. Reginatto, F. A. Ramos, L. Castellanos, C. Duque and E.P. Schenkel, *Phytochem. Anal.*, 23, 232 (2012), DOI: 10.1002/pca.1348
22. K. H. Engel and R. Tressel, *J. Agric. Food Chem.*, 39, 2249 (1991)
23. K. C. Dos Santos, T. V. Borges, G. Olescowicz, F. K. Ludka, C. A. Santos and S. Molz, *J. Pharm. Pharmacol.*, 68, 282 (2016), DOI: 10.1111/jphp.12512
24. M. A. Farag, A. Otify, A.O. Porzel, C. G. Michel, A. Elsayed and L. A. Wessjohann *Anal. Bioanal. Chem.*, 408, 3125 (2016), DOI: 10.1007/s00216-016-9376-4
25. N. Conde-Martinez, D. C. Sinuco and C. Osorio, *Food Chem.*, 157, 356 (2014), DOI: 10.1016/j.foodchem.2014.02.056
26. J. J. Fu, S. N. Li, Y. Wang and L. H. Cao, *Journal of Materials Science*, 45, 6255 (2010), DOI: 10.1007/s10853-010-4720-0
37. L. Ahamad, R. Prasad, and M. A. Quraishi, *Corrosion Science*, **52**, 1472(2010), [DOI: 10.1016/j.corsci.2010.01.015]

38. Messaouda Allaoui, Oumelkheir Rahim and Lakhdarsekhri, *Oriental Journal of Chemistry*, **33**, 637 (2017), [DOI: 10.13005/ijjc/330211]

39. U. F. Ekanem, S. A. Umoren, I. I. Udousoro and A. P. Udoh, *Journal of Materials Science*, **45**, 5558 (2010), [DOI: 10.1007/s10853-010-4617-y]

40. A. Zerga, A. Attayibat, M. Sfaira, M. Taleb, B. Hammouti, M. E. Touhami and Z. Rais, *Journal of Applied Electrochemistry*, **40**, 1575(2010), [DOI: 10.1007/s10800-010-0164-0]

41. F. M. Donahue and K. Nobe, *Journal of the Electrochemical Society*, **112**, 886(1965)

42. G. Moretti, F. Guidi and G. Grion, *Corrosion Science*, **46**, 387(2004), [DOI: 10.1016/S0010-938X(03)00150-1]

43. N. O. Obi-Egbedi and I. B. Obot, *Arabian Journal of Chemistry*, **6**, 211(2013), [DOI: 10.1016/j.arabjc.2010.10.004]

44. A. Doner and G. Kardaş, *Corrosion Science*, **53**, 4223(2011), [DOI: 10.1016/j.corsci.2011.08.032]

45. H. Bourazmi, M. Tabyaoui, L. EL Hattabi, Y. El Aouf ir, M. Taleb, *J. Mater. Environ. Sci.*, **9**, 928(2018), [DOI: 10.26872/jmes.2018.9.3.103]

46. H. Gerengi and H. I. Sahin, *Industrial & Engineering Chemistry Research*, **51**, 780 (2012), [DOI: 10.1021/ie201776q]

47. S. S. Abd El Rehim, M. A. M. Ibrahim, and K. F. Khalid, *Materials Chemistry and Physics*, **70**, 268(2001)

48. M. H. Hussin and M. J. Kassim, *Materials Chemistry and Physics*, **125**, 461 (2011), [DOI: 10.1016/j.matchemphys.2010.10.032]

49. E. E. Oguzie, D. I. Njoku, M. A. Chidebere , C. E. Ogukwe, and G. N. Onuoha, *Industrial & Engineering Chemistry Research*, **53**, 5886(2014), [DOI: 10.1021/ie404273f]

50. A. Rodriguez-Torres, O. Olivares-Xometl, M. G. Valladares-Cisneros, J. G. Gonzalez-Rodriguez, *Int. J. Electrochem.Sci.*, **13**, 3023 (2018), [DOI: 10.20964/2018.03.40]

51. A.Y. El-Etre, M. Abdullah and Z. El-Tantaury, *Corros. Sci.*, **47**, 385(2005), [DOI: 10.1016/j.corsci.2004.06.006]

52. A. Singh, E. E. Ebenso and M. A. Quraishi, *Int. J. Electrochem. Sci.*, **7**, 3409(2012)

53. M. Mobin, M. Rizvi, L. O. Olasunkanmi, and E. E. Ebenso, *ACS Omega*, **2**, 3997-4008 (2017), [DOI: 10.1021/acsomega.7b00436]

54. A. O. Okeke and A. Olaitan, *International Journal of Materials and Chemistry*, **7**, 5(2017), [DOI: 10.5923/j.ijmc.20170701.02]

55. K. K. Alaneme, S. J. Olusegun and O. T. Adelowo, *Alexandria Engineering Journal*, **55**, 673(2016), [DOI: 10.1016/j.aej.2015.10.009]

56. T. K. Bhuvaneswari, V. S. Vasantha, C. Jayaprabha, *Silicon*, **10**, 1793(2018), [DOI: 10.1007/s12633-017-9673-3]

57. G. E. Ugbabe, A. E. Ayodele, G. A. Ajoku, O. F. Kunle, I. Kolo and J. I. Okogun, *Glob Res J.*, **1**, 1(2010)

58. X. Li , S. Deng and H. Fu, *Corrosion Science*, **62**, 163(2012), [DOI: 10.1016/j.corsci.2012.05.008]

59. A. Gislaine, Silva and Carla and B. G. Bottoli, *Critical Reviews in Analytical Chemistry*, **45**, 76(2015), [DOI: 10.1080/10408347.2014.886937]

60. Y. Yetri, Gunawarman, Emriadi and J. Novesar, *ARPN Journal of Engineering and Applied Sciences*, **12**, 18, 5325-5332(2017).

61. I.M. Ritchie, S. Bailey and R. Woods, *Adv. Colloid Interface Sci.*, **80**, 183(1999)