SEED EXTRACT OF SYZYGIUM JAVANICUM AS ECOFRIENDLY

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CORROSION INHIBITOR FOR MILD STEEL IN HYDROCHLORIC

ACID MEDIUM

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Abstract - The corrosion inhibition characteristics of the extract obtained from the seeds of Syzygium javanicum (syn. Eugenia javanica) on mild steel corrosion in hydrochloric acid were investigated by weight loss, electrochemical, IR spectral and SEM studies. The inhibition efficiency increased with increase in inhibitor concentration, but decreased with increase in temperature. Thermodynamic parameters and adsorption behavior were evaluated to reveal the spontaneity of the adsorption process. The polarization measurements indicated that the inhibitor is a mixed type. The confirmation of the passive layer formation was done by Fourier transform infrared (FTIR) spectra and scanning electron microscopy (SEM).

Key words: S. javanicum, steel, corrosion, polarization, impedance, eco-friendly

I. INTRODUCTION

S. javanicum (Fig. 1 & 2), known to be wax apple is a member of Myrtaceae family. The tree is native of Malaya and common in Thailand, Cambodia, Laos, Vietnam and Taiwan, frequently cultivated in India. Primarily the plant is grown as an ornamental, as the fruits are not much valued for food [1]. Even though the uses of seeds are not very much reported, the leaves are found to have medicinal uses.

Corrosion of metals and certain alloys of industrial use have sensed a substantial magnitude of attention of researchers. Because corrosion is a world-wide scientific as well industrial problem as it affects the metallurgical, chemical and oil industries. Mild steel, an alloy of iron finds a number application in above mentioned and other industries. Because of its ease of fabrication is widely used in making vessel, tanks, boilers, pipeline and so on. However it suffers from severe corrosion in aggressive environments, particularly in acidic conditions. In most of the above concerns HCl is used often as a cleaning agent, for acidification, descaling agent, etc. It is customary to make use of corrosion inhibitors to minimize the dissolution of metals by acidic medium. A number of synthetic organic inhibitors having hetero atoms, especially N, S and O are reported to control corrosion of mild steel in acidic solution [2-6]. As most of them are synthetic chemicals, expensive, and hazardous to environments, the use of environmentally safe inhibitors for the metals and alloys is of emerging practice. *Penicillin V Potassium* [7], *Vegetal Tannins* [8], *Parthenium Hystophrous* [9], *Ficus Exasperata* [10] *Citrus aurantiifolia* [11], *Prunus cerasus* [12], *Solanum Tuberosum* [13], *Azadirachta indica* [14], *Andrographis paniculata* [15], *Fenugreek Leaves* [16], Beet root [17] and *Acacia seyal* [18] are also evaluated for anticorrosion properties. In this study the extract obtained from the seeds of *S. javanicum* is evaluated for corrosion inhibition characteristics.

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Fig. 1. S. javanicum plant



Fig. 2. S. javanicum seeds

II. MATERIALS AND METHODS

A. Mild steel specimen:

Mild Steel (MS) specimens of composition Fe = 99.51%, P = 0.08%, Mn = 0.034%, Si = 0.6% and C = 0.16%, polished with different grades of emery paper and then degreased with trichloroethylene is used for the entire study. Weight loss and SEM studies are carried out with MS specimens of size $4.0 \times 2.0 \times 0.19$ cm are employed. MS powder for IR and specimens with an exposed area of 1 cm² are used for electrochemical studies.

B. Preparation of the extract and corrosive environment:

The extract is prepared by refluxing 100 ml of 5% HCl with 50g of dried powder of *S. javanicum* seed for one hour. The extract is cooled, filtered off and made up to 100 ml using double distilled water.

A corrosive environment of 5% (v/v) HCl solution is prepared using HCl and double distilled water. From this stock solution, 100 ml each of standard solutions are prepared with and without the addition of different concentrations of *S. javanicum* seed extract.

Reagent grade, pure HCl (Merck-61752605031730) and double distilled water are used for the entire study.

C. Weight loss study and thermodynamic studies:

Polished and degreased mild steel specimens of known weight are immersed in 100 ml of test solutions with and without different concentrations of inhibitor separately for one hour at four different temperatures viz., 303, 308, 313, and 318K. Afterwards, these specimens are washed with double distilled water, dried well and weighed using Schimadzu AUX220 balance.

D. Infra-Red Spectroscope studies:

FTIR spectra for *S. javanicum* liquid extract and the dried adsorption product formed between MS powder and concentrated solution of the extract is recorded separately using Bruker FTIR model- Tensor27.

E. Surface characterization studies:

Electrochemical parameters such as potentio-dynamic polarization and Impedance measurements are carried out. Tafel polarization plots are recorded potentio-dynamically using platinum electrode, calomel electrode and MS specimen as auxiliary, standard and working electrodes respectively. Polarization studies are carried out at a sweep rate of 1mV/sec. Potential (E) versus log current (log I) plots are then recorded. Impedance measurements are carried out at a frequency range of 10 KHz to 10 mHz. The electrochemical parameters are studied in HCl medium and also with different concentrations of natural inhibitor using Solartron model SI1280B electrochemical measurement unit.

Polished mild steel specimen, specimen exposed to 5% HCl corrosive environment and specimen immersed in 10% inhibitor concentration in 5% HCl are scanned at 10K and recorded using Hitachi S-3000H model Scanning Electron Microscope.

III. RESULTS AND DISCUSSION

Weight loss studies of S. javanicum extract:

The corrosion inhibiton studies are carried out at four different temperatures and the weight loss data obtained for different concentrations of inhibitor and the IE values are listed in **Table 1**. From the table, it is

observed that weight loss is not much pronounced at higher inhibitor concentrations implying the increase in IE (**Fig.3**). However, when the temperature is raised, weight loss is much pronounced, even though the inhibition efficiency is not found to decrease considerably (**Fig 4**). It is also noted that when the exposure time is increased above one hour, the IE decreases and is drastic beyond three hours.

Table 1 Weight loss data

% Conc. of	Weight loss, g				Inhibition efficiency, %			
the inhibitor	303 k	308 k	313 k	318k	303 k	308 k	313 k	318k
0	0.0666	0.0746	0.0853	0.0912				
2	0.0097	0.0109	0.0126	0.0141	85.44	85.39	85.23	84.54
4	0.0079	0.0091	0.0106	0.0120	88.14	87.80	87.57	86.84
6	0.0057	0.0072	0.0091	0.0098	91.44	90.35	89.33	89.25
8	0.0043	0.0059	0.0070	0.0077	93.54	92.09	91.79	91.56
10	0.0031	0.0038	0.0047	0.0055	95.35	94.91	94.49	93.96

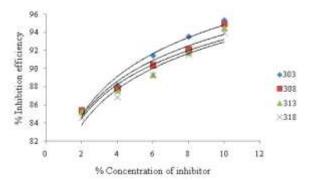


Fig.3. Plot between % IE against % concentration the S. javanicum extract

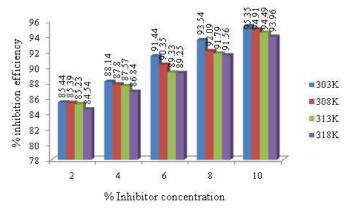


Fig. 4. Effect of temperature on inhibition efficiency of the S. javanicum extract

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Thermodynamic parameters of S. javanicum extract:

The heat of adsorption (Q) was obtained by plotting $\log \theta/1$ - θ against 1/T, where θ is the fraction of the metal surface covered by the inhibitor. Q values were calculated from the slope obtained that is equivalent to -Q/2.303R. The Q values are found to be negative (**Table 2**), which is an indication of decrease in corrosion reaction with increase in inhibitor concentrations and also the exothermic nature of the adsorption process.

A graph was plotted between log (corrosion rate/T) against 1/T to find out the entropy change (ΔS) for the adsorption process. ΔS is calculated from the intercept, which is equivalent to $[log (R/Nh) + (\Delta S/2.303R)]$. The increased adsorption of the inhibitor on the MS surface with increase in concentration is revealed by the increased negative values of ΔS . Generally adsorption results in order from disorder of the natural inhibitor on the metal surface.

The ΔG , free energy change for the adsorption process is calculated using the following formula [19] $\Delta G = -2.303 RT \log 55.5 K$, where $K = (\theta/1-\theta)/C$ ----- (1)

Chemisorption is accompanied by the change in free energy values between -49 KJ/mol and -58 KJ/mol [20]. The change in free energy (**Table 3**) for this adsorption process is in the order of -11KJ/mol, which reveals physisorption of inhibitor on MS surface. The negative free energy change values reveal the spontaneity of the adsorption of the chemical components on the metal surface.

The corrosion rate in mmpy is calculated using the formula

E, energy of activation is obtained using the formula

inhibitor concentration irrespective of the temperature.

$$\log \operatorname{CR}_2 / \operatorname{CR}_1 = \operatorname{E}/2.303 \operatorname{R} \times (1/\operatorname{T}_1 - 1/\operatorname{T}_2)$$

Where CR_1 and CR_2 are the corrosion rates at temperatures T_1 and T_2 respectively [21&22]. The values of corrosion rate and E are given in table 3. The 'E' values are not found to vary much for a particular range of temperatures (303-318 K) even though the concentration of the inhibitor is changed. The energy of activation E is very high for 10% inhibitor dose which is found to be higher than the blank value indicating the requirement of more energy for corrosion reaction to occur at higher concentrations of inhibitor [23].

The adsorption isotherm was obtained between log C and θ , where 'C' is the concentration of the inhibitor (**Fig. 6**). The increased level of adsorption of the inhibitor on the metal surface with increase in inhibitor concentration is revealed from the graph [11].

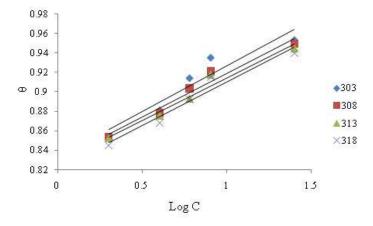


Fig. 5. Adsorption isotherm

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Table 2 Heat of corrosion reaction and change in free energy data

% Conc. of	Q in KJ/mol	ΔS in J/mol	ΔG in KJ/mol				
the inhibitor			303K	308K	313K	318K	
2	-3.52	-244.91	-12.83	-13.03	-13.21	-13.28	
4	-6.01	-249.64	-11.68	-11.79	-11.93	-11.94	
6	-14.34	-264.52	-11.57	-11.43	-11.32	-11.48	
8	-15.28	-274.03	-11.61	-11.25	-11.32	-11.43	
10	-14.76	-280.98	-11.93	-11.88	-11.86	-11.79	

Table 3 Corrosion rate and energy of activation data

% Conc. of	(Corrosion	rate, mmp	E in KJ/mol for the range (K)			
the inhibitor	303K	308K	313K	318K	303-308	308-313	313-318
0	40.66	45.54	52.07	55.67	17.58	21.47	11.07
2	5.92	6.65	7.69	8.61	18.30	23.28	18.72
4	4.82	5.56	6.47	7.33	11.57	24.29	20.67
6	3.48	4.39	5.56	5.98	36.04	37.86	12.06
8	2.62	3.60	4.27	4.70	49.29	27.35	15.89
10	1.89	2.32	2.87	3.36	31.80	34.09	26.11

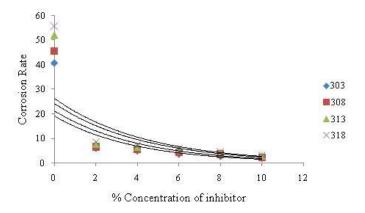


Fig. 6. Correlation between corrosion rate and % concentration of the S. javanicum extract

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IR Spectral studies of S. javanicum extract:

The IR spectra of S. javanicum extract and the adsorption product between extract and MS powder are shown in **Figs. 7 and 8.**When the spectra are compared, the hydroxyl/amine stretching frequency increases from 3421.14 to 3431.16 cm-1, but the carbonyl group frequency of amide decreases significantly from 1633.30 to 1630.01 cm-1. There is also a shift in alcohol group frequency from 1371.24 to 1370.14 cm-1. These variations in group frequency indicate the adsorption of functional groups found in the S. javanicum extract via these groups, which are responsible for corrosion prevention.

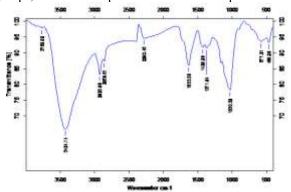


Fig. 7. IR spectra of the S. javanicum extract

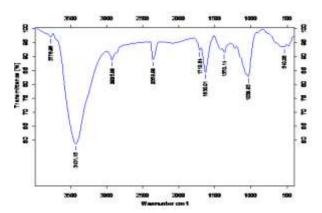


Fig. 8. IR spectra of the adsorption product between S. javanicum extract and MS powder

Electrochemical Studies:

The values of various electrochemical parameters were tabulated in **Table 4.** There is no regular change in corrosion potential (E_{corr}) values from the blank indicating that the inhibitor follows mixed type of inhibition. A steady decrease in the corrosion current (I_{corr}) values with ascending concentration of the inhibitor implies the retardation of corrosion reaction when compared to blank. In general, corrosion current is proportional to the magnitude of oxidation reaction associated with corrosion. The values of anodic and cathodic Tafel slopes (b_a and b_c) also do not increase or decrease in a regular manner inferring the mixed mode of inhibition.

The decrease in double layer capacitance ($C_{\rm dl}$) values (**Table 4**) from the blank with increase in concentration of the inhibitor confirms the increased density of adsorption of the inhibitor on the electropositive metal surface. The adsorption is due to the electronegative hetero atoms present in the constituents of the inhibitor. The better charge transfer resistance (R_{ct}) values when inhibitor concentration is increased highlights the resistance towards the charge transfer reaction viz., corrosion reaction. It should be also noted that the IE values do not vary appreciably when compared with the conventional weight loss method. All the electrochemical parameters show that the corrosion control proceeds through the adsorption of electronegative hetero atoms present in the organic constituents of the extract on the electropositive metal surface. The

inhibition property greatly depends on the concentration of the inhibitor that is evident from the Fig. 9 and 10.

Table 4 Electrochemical parameters of corrosion inhibition by S. javanicum extract

% Conc. of the inhibitor	OCP (V)	E _{corr} (V)	I _{corr} (A)	b _a (V/dec)	b _c (V/dec)	$\begin{array}{c} R_{ct} \\ (Ohm/\\ cm^2) \end{array}$	C _{dl} (A/cm ²)	%IE
Blank	-0.5151	-0.4939	0.002678	150.91	226.11	5.80	6.08 X 10 ⁻⁵	
2	-0.5325	-0.5143	3.42 X 10 ⁻⁴	99.22	156.28	55.93	2.52 X 10 ⁻⁵	87.26
4	-0.5210	-0.5115	2.95 X 10 ⁻⁴	94.87	169.04	70.94	3.95 X 10 ⁻⁵	89.00
6	-0.5231	-0.5116	1.85 X 10 ⁻⁴	94.48	155.72	102.45	2.66 X 10 ⁻⁵	93.08
8	-0.5184	-0.5037	1.84 X 10 ⁻⁴	72.25	158.24	88.73	4.39 X 10 ⁻⁵	93.14
10	-0.5078	-0.5112	1.73 X 10 ⁻⁴	106.44	173.47	145.11	4.79 X 10 ⁻⁵	93.54

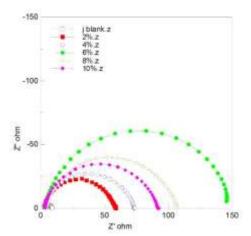


Fig. 9. Impedance spectra

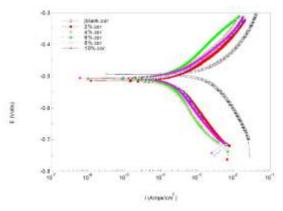
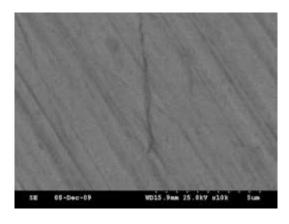


Fig. 10. Tafel polarization plot

The surface morphology of mild steel surface (**Fig. 11**), MS exposed corrosive medium (**Fig. 12**) and MS immersed in 10% inhibitor in corrosive medium (**Fig. 13**) are studied using scanning electron microscopy. The SEM photographs show that in the presence of inhibitor, the pits and corrosion products are not formed on the mild steel surface the mild steel surface. It indicates the formation of passive layer on the metal surface which [24] is clearly seen in **Fig.13**.



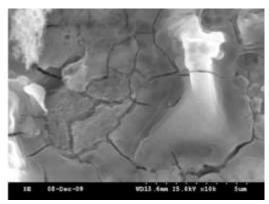


Fig. 11. Polished mild steel surface

Fig. 12. Mild steel exposed to 5% HCl alone

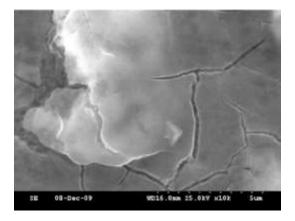


Fig. 13. Mild steel sample exposed to 5% HCl having 10% inhibitor

IV. CONCLUSION

The corrosion inhibition efficiency of S. javanicum extract increases along with the increase in inhibitor concentration and decreases as the temperature is raised as substantiated from the Weight loss studies. Thermodynamic analysis highlights the spontaneity of the adsorption reaction of the inhibitor on the metal surface. The enriched adsorption of the inhibitor on the metal surface with increase in concentration of the inhibitor is evolved from the adsorption isotherm plotted between log C and θ . The adsorption is assigned to the lone pair of electrons present in the hetero atoms of the S. javanicum extract as is revealed from IR spectral studies. The electrochemical parameters such as E_{corr} , b_a and b_c indicate the mixed mode of inhibition. The SEM photographs noticeably emphasize the protective nature of the mild steel by the extract. All these results acquaint with the fact that the seed extract of S. javanicum can be utilized as a nontoxic, alternate, green corrosion inhibitor for mild steel in HCl medium.

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