Corrosion Inhibitive Behaviour of *Camellia Sinensis* on Aluminium Alloy in H\textsubscript{2}SO\textsubscript{4}

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Received: 30 October 2013 / Accepted: 6 December 2013 / Published: 5 January 2014

The effect of *Camellia Sinensis* (green tea) extract as a green inhibitor on the corrosion of aluminium alloy in 0.8M sulphuric acid was studied at room temperature. Weight loss/corrosion rate and potential measurement techniques were used for the experimental work. The results were further analysed using the two-factor ANOVA test. Potential measurement was performed using a digital voltmeter and a saturated calomel reference electrode. The tea extract was obtained from the green tea leaves. The results obtained showed effective corrosion inhibition of the extract on the aluminium alloy test specimens in the different concentrations of sulphuric acid used. The extracts gave appreciable corrosion inhibition performance of aluminium at 20 and 40\% concentrations with the weight loss of 236mg (0.236g) and 265 mg (0.265g) respectively. ANOVA test confirmed the results at 95\% confidence, and further showed that concentration of green tea extract had greater effect on potential and weight loss measurements. The value of Gibb’s free energy of adsorption obtained signified that the mechanism of adsorption of plant extract molecules on the metal surface was by physisorption.

**Keywords:** Corrosion inhibition, *Camellia sinensis*, Aluminium, ANOVA, Adsorption, Gibb’s free energy

1. INTRODUCTION

Metallic materials play a great role in the construction of equipment used in agriculture, oil and gas, petrochemical, process and allied industries as well as in medical services. In these industries, the metallic material as a result of interaction with its environment, losses integrity over a period of time [1]. Bronze, brass, most stainless steels, zinc, and pure aluminium corrode slowly in service conditions. However, corrosion of structural grades of cast iron and steel, 400 series stainless steels,
and some aluminium alloys occur rapidly unless the metal is protected against corrosion. Thus, research into corrosion; its processes and its means of prevention, are of great significance to industry and academia. The use of inhibitors is one of the best methods of preventing metals against corrosion. Most of the corrosion inhibitors are synthetic chemicals, expensive and very hazardous to environments. Therefore, it is desirable to source for environmentally safe inhibitors [2].

Plants represent a class of interesting source of compounds currently being exported for use in metal corrosion protection in most systems. More so, investigations of corrosion inhibiting abilities of tannins, alkaloids, organic, amino acids and organic dyes of plant origin are of interest. In addition to being environmentally friendly and ecologically acceptable, plant products are inexpensive, readily available and renewable sources of materials [3]. The use of phytochemicals as corrosion inhibitors can be traced back to 1960s when tannins and their derivatives were used to protect steel from corrosion [2]. Up till now, extracts of plant leaves also known as green inhibitors such as Newbouldia leavis [4]; Carica Papaya and Camellia Sinensis [5]; Euphorbia hirta [6]; Tithonia diversifolia [7]; Ocimum sanctum [8]; Cola Acuminata and Camellia Sinensis [9]; Vernonia Amygdalina [10,11]; Petersiantus macrocarpus [12]; Camellia Sinensis [13] amongst others have been studied for corrosion inhibition of metals in various media. The present study aimed at investigating the inhibitive properties of Camellia sinensis (green tea) extract on the corrosion of aluminum alloy in 0.8M H₂SO₄ solution. Tea from the leaves of camellia sinensis, a plant of the Theaceae family, is consumed by more than two thirds of the world’s population and is the most popular beverage next only to water.

As previously reviewed [13], tea leaves contain many compounds, such as polysaccharides, volatile oils, vitamins, minerals, purines, alkaloids (e.g. caffeine) and polyphenols (catechins and flavonoids). Green tea contains polyphenols which are mainly flavonoids and are subdivided into flavones, flavonones, isoflavonones, flavanols – flavandiols, anthocyanins, and phenolic acids. The other green tea polyphenols are flavonols, commonly known as catechins – the tea tannins. Green tea polyphenols include groups of compounds of different chemical structure and also possess variable biological properties. Monomeric flavanols, the major components in green tea, are precursors of condensed tannin. Tea polyphenols also have high complexation affinity to metals, alkaloids, and biologic macromolecules such as lipids, carbohydrates, proteins, and nucleic acids. Green tea has very powerful antioxidant properties.

Furthermore, the complex nature of green tea’s chemical composition and structure is expected to prove its effectiveness in corrosion inhibition of aluminium in H₂SO₄, a strong acid. This work therefore, reports the results obtained in the evaluation of the corrosion inhibitive effectiveness of the tea extract on the corrosion of aluminium specimens immersed in 0.8M dilute sulphuric acid at ambient temperature. This study has also yielded some insight into the adsorption mechanisms responsible for the inhibitive property of camellia sinensis in the same corrosive environment. The equation for corrosion rate according to [4] is given by

\[ C = \frac{k \Delta W}{\rho At} \]  

\[ \text{Where } K = \text{Rate constant equal 534 mpy; (mpy means mils per year)} \]
\[ W = \text{Weight loss in mg} \]
\[ \rho = \text{Density of material in g/cm}^3 \]
T = Exposure time in hours
A = Exposed area of coupon in in$^2$ noting that 1 in$^2$ = 6.5416cm$^2$.

2. EXPERIMENTAL DETAILS

2.1 Specimen Preparation

Aluminium sheets of composition (wt %) Al (99), Si (0.157), Mn (0.024), Cr (0.023), Mg (0.51), Cu (0.0025), Ni (0.001), Co (0.001), Ti (0.0049), V (0.0035), Sn (0.001), Zn (0.001), Ca (0.0011), Zr (0.002) and Fe (0.282) were employed for this study. The aluminium sheets were cut into an average size of 2.5cm x 2.5cm coupons for weight loss measurements and 1cm x 1cm coupons for potential measurements. A total number of 12 samples used for the weight loss experiment were descaled with a wire brush, ground with various grades of emery paper and then polished. They were further rinsed in distilled water to remove any corrosion products and then cleaned with acetone to degrease. The samples were fully immersed thereafter preventing further exposure to moisture in the atmosphere. Another set of 12 samples for the corrosion potential experiment were cleaned in the same manner as those for the weight loss experiment except that they were mounted in resin to ensure that only the surface of the samples were exposed to the corrosive medium. Before mounting, copper wire was spot welded to each of the samples.

2.2 Preparation of plant extracts and test media

The experiment was performed in sulphuric acid medium (0.8M H$_2$SO$_4$) of AnalAr grade. 0.8M H$_2$SO$_4$ was prepared by diluting 44.44cm$^3$ of concentrated H$_2$SO$_4$ in 1 liter of distilled water. 

Camellia sinensis (Green tea) was purchased at a local supermarket. The leaves were removed from the bags stored in a container for further use. The plant extract was obtained by the acid extraction method which involved boiling a weighted quantity of leaves in the concentration of sulphuric acid required for a period of two hours and then leaving the mixture to cool. After cooling, the tea leaves are filtered out leaving the solution which contains the leached out constituents of the green tea. The ratio of green tea to acid was 1:10; therefore, for every 40g of green tea leaves used, it was boiled in 400ml of 0.8M H$_2$SO$_4$ solution. From the stock solution produced, inhibitor test solutions were prepared in the percentage concentrations of 20, 40, 60, 80 and 100 respectively. 100ml of the stock inhibitor solution was used as 100% inhibitor concentration. 80% concentration of inhibitor in acid was obtained by mixing 80ml of stock with 20ml of 0.8M H$_2$SO$_4$. 60% was also obtained by mixing 60ml of stock with 40ml of 0.8M H$_2$SO$_4$. The same procedure was followed to obtain 40% and 20% inhibitor concentrations.

2.3 Weight loss experiment

Weighed test specimens were totally immersed in each of the test media contained in a 250ml beaker for 24 days. Experiments were performed with 0.8M sulphuric acid test medium in which some
had the green tea extract added. Test specimens were taken out of the test media every 2 days, washed with distilled water, rinsed in methanol, air-dried, and re-weighed. Plots of weight loss and of calculated corrosion rate versus exposure time respectively (Figs. 1 and 2) were made. Corrosion rate was calculated from the formula in equation 1. The inhibitor efficiency (%) was also calculated for all the inhibitors for every 2 days of the experiment (Fig. 6).

2.4 Potential measurements

Potential measurements were performed on the mounted specimens in turns by immersing them in each of the acid test medium with and without inhibitor. The potential was recorded at 2 – day intervals using a digital voltmeter and saturated calomel reference electrode. Plots of variation of potential (vs. SCE) with the exposure time were made, and these are presented in Fig. 4.

3. RESULTS AND DISCUSSION

3.1 Weight loss method

The results obtained for the variation of weight loss and corrosion rate with exposure time respectively for the aluminium specimens immersed in 0.8M sulphuric acid with varied concentrations of green tea extract are presented in Figs. 1 and 2.

![Graph of weight loss vs. exposure time]

**Figure 1.** Variation of weight loss with exposure time in days for aluminium immersed in 0.8M H₂SO₄ in addition of different concentrations of green tea.

The acid test medium with 80 and 100% concentration of extract addition had the least corrosion inhibition effect of the immersed specimens by the end of the 24th day achieving a weight
loss value of 285mg and 266mg respectively. At this value it could be considered to be fairly protective. The control experiment (without added extracts) recorded a weight loss value of 400mg at the 24th day of the experiment. The test medium with added 60% concentration of green tea performed better than that of the 80%’s and close to 100%’s. It recorded a weight loss value of 267mg on the 24th day of the experiment.

The acid test medium with 20% concentration of extract addition recorded the lowest weight loss with a value of 236mg. These weight loss values showed that the green tea extract exhibited very good corrosion inhibition at the end of the experiment on the 24th day.

The corresponding corrosion rate vs. the exposure time results in Fig. 2 gave a good correlation with the results in Fig. 1. The corrosion rate decreased with time.

![Figure 2. Variation of corrosion rate with exposure time in days for aluminium immersed in 0.8M H₂SO₄ in addition of different concentrations of green tea.](image)

The corrosion rate bears the same relationship as the weight loss with the specimen immersed in the 20% inhibitor having the least corrosion rate throughout the duration of the experiment. Among all the specimens immersed in solutions containing inhibitor, the sample with 100% inhibitor had the highest corrosion rate up until the 18th day of exposure after which it reduced slightly to the rate of the specimens immersed in 60% and 40% inhibitor. The poor effect in high concentration of inhibitor when used for aluminium could be due to the environment or due to the passive nature of aluminium.

**Statistical Analysis**

Two-factor single level experiment ANOVA test (F-test) was used to evaluate the separate and combined effects of concentration of green tea (GT) extracts and exposure time on the weight loss of the aluminium in 0.8M H₂SO₄ solution. The F-test was used to examine the amount of variation within
each of the samples relative to the amount of variation between the samples. As previously used by [14], the Sum of Squares among columns (exposure time) was obtained with the equation:

$$SS_c = \frac{\sum \tau_e^2}{nr} - \frac{\tau^2}{N}$$  \hspace{1cm} (2)

Sum of Squares among rows (concentration of GT):

$$SS_r = \frac{\sum \tau_r^2}{nc} - \frac{\tau^2}{N}$$  \hspace{1cm} (3)

Total Sum of Squares:

$$SS_{total} = \sum x^2 - \frac{\tau^2}{N}$$  \hspace{1cm} (4)

The calculation using the ANOVA test is tabulated (Table 1) as shown. On the basis of the results in Table 1, it can be concluded with 95% confidence that the exposure time and concentration of GT significantly affects the corrosion rate of aluminium in 0.8M H$_2$SO$_4$ environment. This gave good correlation with the scatter plots obtained in Figs. 1 and 2. It was further observed that the test exposure time had a greater significant effect on the corrosion rate (Fig.3).

Table 1. Summary of ANOVA analysis for weight loss measurements

<table>
<thead>
<tr>
<th>Source of Variation</th>
<th>SS</th>
<th>Df</th>
<th>MS</th>
<th>F</th>
<th>Significance F</th>
</tr>
</thead>
<tbody>
<tr>
<td>Exposure Time</td>
<td>453265.78</td>
<td>11</td>
<td>41205.98</td>
<td>100.77</td>
<td>1.97</td>
</tr>
<tr>
<td>Concentration of GT</td>
<td>42549.78</td>
<td>5</td>
<td>8509.96</td>
<td>20.81</td>
<td>2.38</td>
</tr>
<tr>
<td>Residual</td>
<td>22489.56</td>
<td>55</td>
<td>408.90</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>518305.11</td>
<td>71</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Figure 3. F values showing the influence of exposure time and GT concentration on Weight Loss measurements
3.2 Potential measurement

Potential readings for the aluminium specimens were taken over a period of 24 days at an interval of 2 days. The curves obtained for the variation of potential (mV) vs. saturated calomel electrode (SCE) with the exposure time are presented in Fig. 4. The specimens were immersed separately, in 0.8M H$_2$SO$_4$ with different concentrations (20, 40, 60, 80 and 100%) of green tea.

![Figure 4. Variation of potential with exposure time for aluminium immersed in 0.8M H$_2$SO$_4$ in addition of different concentrations of green tea.](image)

The test medium without the inhibitor addition recorded the most negative potential in the first 4 days of the experiment with a potential value of −708mV on the 4th day.

This was a clear indication of active corrosion reactions and can thus be correlated with the results obtained in the weight loss experiments. The curves for variation of potential with exposure time in the test media for the extracts of 20, 40 and 60% concentrations showed effective corrosion protection throughout the experimental period. The extracts with the 20% concentration showed the best corrosion inhibition performance with potential values of -102mV on first day of the experiment and -216mV on the 14th day and on the 24th day, the potential value was -250mV. All these values fell within the passive corrosion reactions range with respect to SCE. The potential values for the 40% concentration extracts were slightly more negative than the former and showed less passive corrosion reactions – with a potential value of -290mV on the first day and -253mV on the 24th day of the experiment.

The effect of these variables on the corrosion potential of the aluminium alloy was further confirmed with the ANOVA test using equations (2) – (4) as stated earlier. The results are displayed in Table 2.
Table 2. Summary of ANOVA analysis for potential measurements

<table>
<thead>
<tr>
<th>Source of Variation</th>
<th>SS</th>
<th>Df</th>
<th>MS</th>
<th>F</th>
<th>Significance F</th>
</tr>
</thead>
<tbody>
<tr>
<td>Exposure Time</td>
<td>21142.49</td>
<td>11</td>
<td>1922.04</td>
<td>0.95</td>
<td>1.97</td>
</tr>
<tr>
<td>Concentration of GT</td>
<td>1930574.90</td>
<td>5</td>
<td>386114.98</td>
<td>190.87</td>
<td>2.38</td>
</tr>
<tr>
<td>Residual</td>
<td>111258.60</td>
<td>55</td>
<td>2022.88</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>2062975.99</td>
<td>71</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Figure 5. F values showing the influence of exposure time and GT concentration on potential measurements.

On the basis of the results in Table 2, it can be concluded with 95% confidence that varied concentration of GT significantly affects the corrosion potential of aluminium in 0.8M H$_2$SO$_4$, while the effect of exposure time is insignificant (Fig. 5).

3.3 Inhibitor Efficiency

The results of the inhibitor efficiency obtained by calculations are presented in Fig. 6. There was increased inhibitor efficiency with decreased per cent concentration of inhibitor and increased exposure time throughout the whole experimental period. The extracts with 20% concentration addition gave the optimal inhibition efficiency of 35% on 8th and 40% on the 20th day of the experiment. Similarly, the 40% concentration addition gave inhibition efficiency of 18 and 30% on the 8th and 20th day of the experiment respectively. The 60% concentration addition gave the inhibition efficiency of 15 and 29% on the 8th and 20th day respectively. Though negative inhibition results were obtained during the first few days of the experiments with both 100 and 80% concentrations, inhibition was nonetheless achieved for the greater part of the experiment.
After removing the samples from solution at the end of the experiment, severe corrosion was observed (Fig. 7a) by inspecting the surface of the sample without inhibitor under the microscope. This suggests that sulphuric acid can be aggressive on the aluminium alloy to induce corrosion.

A positive change was observed by using the green tea inhibitor (Fig. 7b), indicating some degree of corrosion inhibition.

*Thermodynamics and Adsorption Studies*
Corrosion inhibition of aluminium alloy in 0.8M H\textsubscript{2}SO\textsubscript{4} by green tea can be further explained based on molecular adsorption. The adsorption process is influenced by the chemical structures of organic compounds, the distribution of charge in molecule, the nature and surface charge of metal and the type of aggressive media [12]. The value of \( k \) which is the adsorption equilibrium constant and the standard free energy of adsorption \( \Delta G_{\text{ads}}^{\circ} \) were evaluated based on the relation given by [4]:

\[
k = \frac{1}{55.5} \exp \left( \frac{-\Delta G_{\text{ads}}^{\circ}}{RT} \right)
\]

(5)

Where \( \Delta G_{\text{ads}}^{\circ} \) is the standard free energy of adsorption

\( R \) is the molar gas constant and

\( T \) is the absolute temperature

**Table 3.** Thermodynamic parameters for aluminium in 0.8M H\textsubscript{2}SO\textsubscript{4} with green tea extracts

<table>
<thead>
<tr>
<th>Inhibitor concentration</th>
<th>( \Theta )</th>
<th>( K )</th>
<th>( \log K )</th>
<th>( 1-\Theta )</th>
<th>( \Theta/1-\Theta )</th>
<th>( \Delta G ) (kJ/mol-1)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>0.00</td>
<td>0.00</td>
<td>0.00</td>
<td>1.00</td>
<td>0.00</td>
<td>0.00</td>
</tr>
<tr>
<td>20% GT</td>
<td>0.41</td>
<td>3474.58</td>
<td>3.54</td>
<td>0.59</td>
<td>0.69</td>
<td>-20.20</td>
</tr>
<tr>
<td>40% GT</td>
<td>0.34</td>
<td>1273.58</td>
<td>3.11</td>
<td>0.66</td>
<td>0.51</td>
<td>-17.72</td>
</tr>
<tr>
<td>60% GT</td>
<td>0.33</td>
<td>830.21</td>
<td>2.92</td>
<td>0.67</td>
<td>0.50</td>
<td>-16.66</td>
</tr>
<tr>
<td>80% GT</td>
<td>0.29</td>
<td>504.39</td>
<td>2.70</td>
<td>0.71</td>
<td>0.40</td>
<td>-15.42</td>
</tr>
<tr>
<td>100% GT</td>
<td>0.26</td>
<td>342.28</td>
<td>2.53</td>
<td>0.75</td>
<td>0.34</td>
<td>-14.46</td>
</tr>
</tbody>
</table>

The negative values of \( \Delta G_{\text{ads}}^{\circ} \) obtained implies that the adsorption process was spontaneous and there was stability of the absorbed inhibitor layer on the metal surface. The values of the standard free energy of adsorption for 20% to 100% plant extracts is within the range of -15 to -20kJ/mol, characterizing the physiosorption mode of adsorption. Studies have shown that \( \Delta G_{\text{ads}}^{\circ} \) values that are -20kJ/mol and above, i.e. less negative are consistent with physical adsorption (physiosorption) which involves electrostatic interaction between charged atoms and the charged metal, while those around -40kJ/mol and more negative are generally associated with chemical adsorption[4,7].

**3.4 Summary**

The overall corrosion and inhibition profile showed that good corrosion inhibition was achieved with the use of these extracts. The potential values obtained as presented in the curves bear correlation with the results obtained gravimetrically. The potential values obtained for the green tea extracts of different concentrations fell within the accepted range for fairly good protection for aluminium with reference to saturated calomel electrode.

In general, the effective corrosion inhibition performance of green tea extract could be associated with their complex chemical compounds which include tannin and polyphenols. The action/reaction of this compound on the surface of the aluminium could hinder the sulphate ion
species, promote more stable passive film formation and hence inhibit and stifle corrosion reactions at the steel / environment interface.

4. CONCLUSION

Green tea was found to be an effective natural corrosion inhibitor for aluminium in 0.8M H₂SO₄ by using potential and weight loss methods. The results obtained from both methods were in good agreement. ANOVA test confirmed at 95% confidence that the exposure time and concentration of Green Tea significantly affects the corrosion rate of aluminium in 0.8M H₂SO₄ environment. For the potential measurements, ANOVA test confirmed at 95% confidence that varied concentration of green tea had significant effect on corrosion potential but none on exposure time. Furthermore, the inhibition efficiency of the plant extract was found to increase with decreasing inhibitor concentration. In addition, the value of Gibb’s free energy of adsorption obtained signified that the mechanism of adsorption of plant extract molecules on the metal surface was by physisorption.

ACKNOWLEDGEMENT

The authors acknowledge the laboratory investigation of Mr. Akintomide A. Akintoye, and also, the Department of Mechanical Engineering, Covenant University, Ota for the provision of research facilities for this work.

References


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