Corrosion Inhibition of Triazinedithiol for Aluminum Alloy in Hydrochloric Acid Solution

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Abstract

The corrosion inhibition property of 6-diallylamino-1,3,5-triazine-2,4-dithiol monosodium (DAN) for aluminum alloy corrosion in hydrochloric acid solution was investigated by weight loss, potentiodynamic polarization, cyclic voltammetry and scanning electron microscopy (SEM). The effect of DAN concentration, temperature, immersion time and acidic concentration and was studied. It was found that DAN presented excellent corrosion inhibition performance for aluminum alloy. Polarization curves revealed that DAN acted as the cathodic inhibitor. The inhibition efficiency was improved with the concentration of DAN inhibitor, while decreased with the improvement of corrosion temperature and immersion time. The adsorption isotherm of DAN inhibitor on aluminum alloy surface conformed to langmuir adsorption model and the value of ΔGθ inferred the interaction of DAN inhibitor with aluminum as chemical adsorption at 40°C.

Keywords: Aluminum Alloy; Weight Loss; Potentiodynamic Polarization; Adsorption Isotherm; Inhibitor

Introduction

Aluminum alloy is extensively used in many engineering applications, such as automobile, aviation, house holding appliance, container and electronic device due to its excellent properties including low cost, light weight, good appearance, mechanical strength, high thermal and electrical conductivity [1]. The thin, adherent and compact oxide film formed on aluminum surface can protect the metal in various environments [2-3]. But the oxide film is amphoteric and the metal dissolves readily in acidic or basic solutions [6]. Mineral acid, mainly hydrochloric acid is widely used to remove oxide in several industrial processes such as acid pickling, acid cleaning and acid, mainly hydrochloric acid is widely used to remove oxide in several industrial processes such as acid pickling, acid cleaning and acidic concentration and was studied. It was found that DAN presented excellent corrosion inhibition performance for aluminum alloy. Polarization curves revealed that DAN acted as the cathodic inhibitor. The inhibition efficiency was improved with the concentration of DAN inhibitor, while decreased with the improvement of corrosion temperature and immersion time. The adsorption isotherm of DAN inhibitor on aluminum alloy surface conformed to langmuir adsorption model and the value of ΔGθ inferred the interaction of DAN inhibitor with aluminum as chemical adsorption at 40°C.

Some triazine derivatives used to inhibit the corrosion of mild steel were reported previously [11,12], whereas the study on triazine inhibitor for aluminum alloy was nearly mentioned. In this work, a new triazine derivative, 6-diallylamino-1,3,5-triazine-2,4-dithiol monosodium (DAN) was synthesized by the reaction between 6-N,N-diallylamino-1,3,5-triazine-2,4-dithiol monosodium (DAN) and was used to study the corrosion inhibition for aluminum alloy. According to the previous researches about DAN and its structural features [13,14], it can be inferred that DAN could be an efficient inhibitor due to its special properties such as the presence of hetero atoms (N and S), N-containing heterocyclic conjugate system, high solubility and low toxicity. The corrosion inhibitive ability of DAN was evaluated by weight loss, potentiodynamic polarization, cyclic voltammograms and scanning electron microscope (SEM) technique.

Experimental

Materials

The substrate for this study was AA5052 aluminum alloy with the dimension of 30 mm×50 mm×0.3 mm and the chemical compositions of AA5052 are shown in Table 1. The specimen was used for weight loss and the specimens with an exposed area of 1 cm² (rest is covered by polyimide tape) were used for potentiodynamic polarization. All specimens were ultrasonically degreased with acetone for 15 min. The deoxidation treatment was carried out in alkaline solution at 60°C for 1 min, washed thoroughly with distilled water and dried at room temperature. The corrosive media 0.2 M HCl solution was prepared using AR grade HCl and double distilled water. 6-diallylamino-1,3,5-triazine-2,4-dithiol monosodium (DAN) was synthesized by the reaction between 6-N,N-diallylamino-1,3,5-triazine-2,4-dichloride and NaSH. The molecular structure of DAN is shown in Figure 1.

Methods

Weight loss measurements: Weight loss measurements were performed by immering the aluminum alloy specimen in the glass beaker containing HCl solution without and with different concentration of DAN inhibitor (0.1 ~ 2.0 mM). After the stipulated immersion time, the specimen was taken out and washed with distilled water, dried and weighed accurately using digital balance (accuracy: ± 0.1 mg). Each measurement was repeated three times and an average value was calculated. Immersion experiments were carried out at temperature of 30, 40, 50, 60 and 70°C in a digital temperature controlled water bath. The immersion time of 1, 5, 10, 24 and 48 h was controlled respectively. The corrosion inhibition efficiency ηw (%) and surface coverage (θ) can be obtained by the following equations.

\[ \eta_w(\%) = \frac{W_0 - W_i}{W_0} \times 100 \]  

\[ \theta = \frac{W_i - W_0}{W_0} \]  

where \( W_0 \) is the weight of the specimen before immersion, \( W_i \) is the weight of the specimen after immersion and ΔW is the weight loss.

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Where \( W_0 \) and \( W_i \) are the weight losses of AA5052 in the absence and in the presence of DAN inhibitor, respectively. And the same experiment was carried out for three times. The data in this study was an average value from three repetitions.

**Electrochemical measurements:** The potentiodynamic polarization and cyclic voltammogram measurements were carried out in CHI 660C electrochemical work station in a three-electrode cell system with a saturated calomel electrode (SCE) as reference electrode at 30°C. The counter electrode was rectangular graphite in potentiodynamic polarizations study, while it was stainless steel in cyclic voltammogram study. The working electrode was pretreated AA5052. An immersion time of 1 h was given to allow the stabilization of the open circuit potential prior to any electrochemical measurements. The experimental solutions were 0.5 M NaCl and 0.1% H2SO4, respectively. The corrosion inhibition efficiency \( \eta_p (\%) \) was calculated from the measured corrosion current density (\( I_{corr} \)) by the following relationship.

\[
\eta_p(\%) = \left(\frac{I_{corr} - I_{corr}}{I_{corr}}\right) \times 100 \tag{3}
\]

where \( I_{corr} \) and \( I_{corr} \) are the corrosion current densities of AA5052 in the absence and in the presence of DAN inhibitor, respectively.

**Scanning electron microscopy (SEM):** The surface morphologies of AA5052 before and after the 0.2 M HCl immersion for 24 h in the absence and in the presence of 2.0 M DAN inhibitor were observed by SEM (JSM-6360LV) at accelerating voltage of 20 kV.

**Results and Discussion**

**Weight loss studies**

**Effect of DAN inhibitor concentration:** The variation of \( \eta_w (\%) \) with the concentration of DAN inhibitors (0.1~2.0 mM) for 24 h of immersion time in 0.2 M HCl solution at 30°C is presented in Figure 2. It can be seen from the figure that the corrosion inhibition efficiency is proportional to the DAN concentration up to 0.5 mM. Thereafter, a marginal increase is noticed in the range 0.5 ~ 2.0 mM concentration. The result indicated that DAN inhibitor molecules could adsorb on the surface of aluminum alloy to form a relative complete protective layer even in low concentration. The maximal inhibition efficiency was obtained at the 2.0 M DAN concentration. It was proposed that the adsorption rate of DAN inhibitor molecule on AA5052 surface was equal to desorption rate at the concentration.

**Effect of temperature:** To examine the action of inhibitors at elevated temperatures, weight loss experiments were carried out at different temperatures keeping immersion time at for 24 h in 0.2 M HCl solution with 2.0 mM DAN. The variation of \( \eta_w (\%) \) with temperature is shown in Figure 3, which indicated that \( \eta_w (\%) \) exhibited decrease with the increasing of experimental temperature. It is well known that decrease in efficiency with the increase of temperature is attributed to the physical adsorption. The time gap between the process of adsorption and desorption of inhibitor molecules over the metal surface is becoming shorter with increase in the temperature. Hence, the metal surface with the incomplete protective layer remains exposed to the acid environment for longer period, therefore the inhibition efficiency falls at elevated temperature [15].

**Effect of immersion time:** To see the effect of immersion time on \( \eta_w (\%) \), weight loss measurements were carried out for different immersion times, from 1 to 48 h. The variation of \( \eta_w (\%) \) with immersion time is shown in Figure 4. It was noted that the variation of \( \eta_w (\%) \) with immersion time became large significantly within the first 5 hours. When the immersion time was 5 hours, the inhibition efficiency of DAN molecules reached the highest value of 93.82%. In general, the adsorption of the inhibitor molecule is often a displacement reaction involving removal of adsorbed water molecules from the metal surface. It was assumed that the adsorption rate of DAN inhibitor molecules was higher than desorption rate within 5 hours and adsorption layer existed on the metal surface to make the anti-corrosion performance improved. When the immersion time was 5 hours, the adsorption of DAN inhibitor molecules became saturated. In hydrochloric acid medium, the metal surface is negatively charged due to the specifically adsorbed Cl- ions on the metal surface. The nitrogen atom of the

![Figure 1: The molecular structure of DAN.](image1)

![Figure 2: Variation of \( \eta_w(\%) \) with concentration of DAN inhibitor for an immersion time 24 h in 0.2 M HCl solution at 30°C](image2)

![Figure 3: Variation of \( \eta_w(\%) \) with temperature for an immersion time of 24 h in 0.2 M HCl solution at different temperatures. (DAN concentration is 2.0 M).](image3)
densities ($I_{corr}$) and $I_{corr}$ decreased with the increasing of DAN inhibitor. A prominent decrease of corrosion current (% was calculated from equation (3). It could be seen that the presence of DAN molecule mainly acted as the cathodic inhibitor [18]. The $\eta_p$ (%) values increased with increase in the inhibitor concentration and are in agreement with $\eta_w$ (%) obtained from weight loss measurements.

Cyclic voltammogram: The cyclic voltammograms curves can be used to study the surface coverage of inhibitor molecules. Figure 6 shows the cyclic voltammograms of aluminum alloy electrode in 0.1% H$_2$SO$_4$ solution. With the scanning potential ranged from -1.0 V to 2.0 V, the current density of the blank aluminum changed significantly from 0 to 0.88 A cm$^{-2}$, which indicated that the surface stability of the blank was poor and apt to be corroded. Compared with the blank, the aluminum alloy electrodes in the presence of DAN inhibitor had relative lower current densities and the current density became smaller gradually with the increasing of DAN concentration. This might be attributed to the adsorption of DAN molecules on aluminum alloy surface to form the protective layer. Hence, it could block corrosive electrolyte particle through the surface of aluminum electrodes and played a protective role for metal material. Table 3 presents the mathematics area and $\eta_c$ (%) of cyclic voltammogram in the absence and presence of different concentrations of DAN. As the adding amount of DAN inhibitor was improved, the mathematics area of cyclic voltammogram curve reduced from 0.08653 to 0.00366 and the $\eta_p$ (%) reached 96% at 2 mM DAN concentration, which was consisted with the results of weight loss test and potentiodynamic polarization.

| Ci(mM) | $E_{corr}$/mV(vs. SCE) | $I_{corr}$/μA·cm$^{-2}$ | $\beta_c$/V | $\eta_p$/% |
|--------|------------------------|------------------------|-------------|------------|
| 0      | -530                   | 1640                   | 0.934       | -          |
| 0.1    | -622                   | 290                    | 0.568       | 82         |
| 0.5    | -640                   | 264                    | 0.857       | 84         |
| 1.0    | -639                   | 243                    | 0.888       | 85         |
| 1.5    | -643                   | 176                    | 0.915       | 89         |
| 2.0    | -638                   | 104                    | 0.912       | 94         |

Table 1: Chemical composition for aluminum alloy AA5052 (%).

| Ci(mM) | mathematics area/[×10$^{-4}$] | $\eta_p$/% |
|--------|-------------------------------|------------|
| 0      | 865.3                         | -          |
| 0.1    | 323.9                         | 63         |
| 0.5    | 279.7                         | 68         |
| 1.0    | 151.6                         | 83         |
| 1.5    | 97.5                          | 89         |
| 2.0    | 36.6                          | 96         |

Table 3: Cyclic voltammogram of the corrosion for aluminum alloy in 0.1% H$_2$SO$_4$ in the absence and presence of different concentrations of DAN.

**Electrochemical studies**

**Potentiodynamic polarization:** Potentiodynamic polarization profiles for AA5052 in 0.5 M NaCl solution in the presence of different concentrations of DAN are presented in Figure 5. The corrosion parameters such as corrosion potential ($E_{corr}$), corrosion current density ($I_{corr}$) and cathodic Tafel slope ($\beta_c$) can be obtained from Figure 5 as shown in Table 2. The percentage inhibition efficiency $\eta_p$ (%) was calculated from equation (3). It could be seen that the presence of DAN inhibitor caused a prominent decrease of corrosion current densities ($I_{corr}$) and $I_{corr}$ decreased with the increasing of DAN inhibitor concentration. When the adding amount of DAN inhibitor was 2.0 mM, the $I_{corr}$ was the lowest with the value of 104 μA·cm$^{-2}$ and inhibition efficiency reached 94%. From Table 1, the value of $E_{corr}$ was shifted towards negative (noble) direction with the increasing of DAN inhibitor concentration while $I_{corr}$ was shifted towards lower current density region, which indicated that the cathode reaction of aluminum alloy was suppressed to some extent. Therefore, the entire corrosion process was retarded in the system with DAN inhibitor molecules. In literature, it has been reported that a compound can be classified as an anodic or cathodic-type inhibitor when the change in $E_{corr}$ is greater than 85 mV [17] with reference to blank specimen, otherwise inhibitor is treated as mixed type. In our study, maximum displacement in $E_{corr}$ value was around 113 mV, which indicated that DAN molecule mainly acted as the cathodic inhibitor [18].
The inhibition efficiency of DAN molecule mainly depends on its adsorption ability at metal-solution interface. The surface coverage \( \theta \) for different DAN concentrations is defined as \( \eta \) with the adsorption isotherm. Several adsorption isotherms like Temkin, Freundlich, Langmuir and Flory–Huggins were tested for the description of adsorption behavior of DAN molecules. Among them, the best fit was Langmuir isotherm as shown in Figure 8. It was noted that the linear correlation coefficient \( R^2=0.99417 \) of the straight line \( c/\theta = 1 + K_{ads} c \) was very close to 1 and the slope of the straight line (1.04076) was also close to 1, which indicated that the adsorption of DAN molecule on aluminum alloy surface obeyed Langmuir model at 40°C. Namely, the characteristic of Langmuir adsorption isotherm is given by following equation [19,20]:

\[
\frac{c}{\theta} = \frac{1}{K_{ads}} + c
\]

where \( c \) is the concentration of inhibitor, \( K_{ads} \) is the adsorptive equilibrium constant and \( \theta \) is the surface coverage equal to inhibition efficiency \( \eta \)/100. The \( K_{ads} \) can be calculated as 5.50 L/mmol from the intercept of fitting formula in Figure 8. For Langmuir model, the relationship of \( K_{ads} \) and standard adsorption free energy (\( \Delta G^\theta \)) can be expressed by the following Eq. (5):

\[
K = \frac{1}{c_{ads}} \exp\left(-\frac{\Delta G^\theta}{RT}\right)
\]

where \( R \) is molar gas constant, \( T \) is temperature in Kelvin and \( c_{ads} \) is the water concentration in solution. It should be noted that the unit of \( c_{ads} \) lies in that of \( K \). As can be seen from Figure 8, the unit of \( K \) is L/mmol which implies that the unit of \( c_{ads} \) is mmol/L with the value of approximate 5.55x10^-4 mol/L. Hence, the value of \( \Delta G^\theta \) was calculated to be -44.664 kJ/mol. The negative value of \( \Delta G^\theta \) suggests that inhibitor molecules are spontaneously adsorbed on metal surface. The adsorption strength is directly proportional to value of \( \Delta G^\theta \).

In general, \( \Delta G^\theta \) value of -20 kJ/mol or lower indicates physisorption interaction while those around -40 kJ/mol or higher is generally accepted to form a coordinate bond by the charge sharing or transferring between the inhibitor molecules and the metal surface [21]. In this study, the value of \( \Delta G^\theta \) (40°C) is -44.664 kJ/mol. It was suggested that the chemical adsorption of DAN was most possible. It can be attributed to the presence of hetero atoms (N and S) and N-containing heterocyclic conjugate system. Thus, the possible reaction centres are unshared electron pairs on hetero atoms and \( \pi \) electrons of c N-containing heterocyclic conjugate system. Especially, the existence of -SH group and –S- in DAN molecule is in favor of chemisorb on the metal surface in acidic media [22].

**Conclusions**

The organic triazine compound of DAN was successfully applied as corrosion inhibitor for the aluminum alloy in HCl solution. DAN inhibitor formed the protective layer on aluminum surface to block the corrosion reaction and acted as a cathodic-type inhibitor. All of the experimental methods indicated that the inhibition efficiency of DAN compound for AA5052 in 0.2 M HCl solution is 98.5% (95.85% in absence of inhibitor) and 96.07% (93.41% in absence of inhibitor) for 2.0 mM DAN concentration whereas the inhibition efficiency of blank solution is much lower (96.20% in absence of inhibitor). This study provided a new way to find the protective layer formed on AA5052 surface in 0.2 M HCl solution.

**Scanning electron microscopy (SEM)**

The morphologies of the aluminum alloy surface were also observed before and after 24 h immersion in 0.2 M HCl and in presence of 2.0 mM DAN inhibitor, shown in Figure 7. It can be seen that the aluminum sheet before immersion seems smooth and the scratches, kinks and folds introduced by rolling process on AA5052 surface (Figure 7a). The AA5052 surface was severely infected by the presence of corrosive HCl solution as seen in Figure 7b. The cracks and grooves together with huge amount of irregular deep voids distributed along the surface, confirmed the role of HCl to make corrosion in three dimensions. Figure 7c showed the aluminum surface in presence of DAN inhibitor and corrosive HCl solution. Deep and large voids observed in Figure 7b drastically reduced due to the adsorptive layers of inhibitor that prevented HCl penetration into the aluminum surface, which further confirmed the inhibition ability of DAN. Conclusively, the regular distribution of the DAN molecules generated the formation of consistent protective layers on AA5052 surface that could protect the metal from corrosive HCl.

**Adsorption isotherm**

The inhibition efficiency of DAN molecule mainly depends on its adsorption ability at metal-solution interface. The surface coverage \( \theta \) for different DAN concentrations is defined as \( \eta \)/100 and can be obtained from weight loss measurements. Basic information about the interaction between inhibitor and AA5052 can be provided by the adsorption isotherm. Several adsorption isotherms like Temkin, Freundlich, Langmuir and Flory–Huggins were tested for the description of adsorption behavior of DAN molecules. Among them, the best fit was Langmuir isotherm as shown in Figure 8. It was noted that the linear correlation coefficient \( R^2=0.99417 \) of the straight line (1.04076) was also close to 1, which indicated that the adsorption of DAN molecule on
was highest and more than 90% at the concentration of 2.0 mM DAN. The adsorption of DAN molecules on aluminum alloy surface obeyed Langmuir adsorption isotherm. The $\Delta G^0$ value inferred that the adsorption process of DAN inhibitor was spontaneous and the interaction of inhibitor with aluminum alloy was chemical sorption at 40°C. The results of weight loss, electrochemical evaluation and SEM are in good agreement.

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