



Synthesis and evaluation of 5-(Phenyl)-4H-1,2,4-triazole-3-thiol as corrosion inhibitor for mild steel in 0.5M H₂SO₄ and its synergistic effect with potassium iodide

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Abstract

The corrosion inhibition by 5-(Phenyl)-4H-1,2,4-triazole-3-thiol (PTT) on mild steel in 0.5 M H₂SO₄ solution has been investigated by weight loss, electrochemical impedance spectroscopy (EIS) and potentiodynamic polarization techniques at various concentrations and temperatures. The results obtained revealed that this compound performed excellently as corrosion inhibitor for mild steel in 0.5 M H₂SO₄ solution. It was found that the inhibition efficiency increased with inhibitor concentration reached a maximum of 91.6 % at 0.5 mM. The addition of potassium iodide to PTT in solution increased the inhibition efficiency of this latter. A synergistic effect was observed between KI and inhibitor with optimum concentration of 0.5 mM/PTT and 0.2% potassium iodide. Potentiodynamic polarization studies have shown that PTT inhibitor acts as a mixed-type inhibitor retarding the anodic and cathodic corrosion reactions with predominant effect on the cathodic reaction. Adsorption of inhibitor alone or in combination with potassium iodide on the metal surface obeyed the Langmuir adsorption isotherm. The effect of temperature on the inhibition efficiency was also determined, in addition some thermodynamic parameters such as apparent activation energy and adsorption free energy have been calculated and discussed. The results obtained with different methods are in good agreement. Scanning electron microscopy (SEM) study confirmed that the inhibition of corrosion of mild steel is through adsorption of the extract molecules on the surface of metal. Quantum chemical parameters were also calculated to characterize adsorption mechanism. Acceptable correlations were obtained between experimental (inhibition efficiencies, ΔG_{ads} , E_a) and quantum calculation parameters (dipole moment, E_{HOMO} , E_{LUMO}). The high inhibition efficiency was declined in terms of strongly adsorption of protonated inhibitor molecules on the metal surface and forming a protective film. Electrochemical techniques have been used for the first time to study synergistic effect of PTT inhibitor and potassium iodide on inhibition of corrosion of mild steel in 0.5 M H₂SO₄ solution. The results suggest that the mixture (PTT + KI) could find practical applications in corrosion control in aqueous acidic environment. The effect of molecular structure on the inhibition efficiency has been investigated by quantum chemical calculations. The electronic properties of inhibitor were calculated and are discussed. The theoretical results were found to be consistent with the experimental data reported.

Keywords: Mild steel; Corrosion inhibitor; 1,2,4-triazole; H₂SO₄ solution; EIS; Tafel polarization; SEM; Density functional theory

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1. Introduction

Mild steel is one of the important materials used in various industries due to its excellent mechanical properties and low cost. The study of carbon steel corrosion phenomena has become important particularly in acidic media because of the increased industrial applications of acid solutions. One of the most common, effective and economic method to protect metals against corrosion is use Messikh et al., 2020

of organic compounds (containing heteroatoms having higher basicity and electron density like nitrogen, oxygen and sulphur) as corrosion inhibitors [1-10]. Compounds containing hetero atoms in their aromatic or long carbon chain are capable of being adsorbed on the metal surface and can protect the metal against corrosion. For this class of compounds, the presence of hetero atoms (nitrogen, sulphur, oxygen and even selenium and phosphorous) and π -

electrons in their double or triple bonds have been found to help the adsorption of the metal. The existing data reveal that most organic inhibitors act by adsorption on the metal surface [11-13]. These inhibitors are usually adsorbed on the metal surface by the formation of a coordinate covalent bond (chemical adsorption) or the electrostatic interaction between the metal and inhibitor (physical adsorption). This adsorption produces a uniform film on the metal surface, which reduces or prevents contact with the corrosive medium. Because organic inhibitors act by adsorption on the metal surface, the efficiency of these compounds depends strongly on their ability to form complexes with the metal. Both π electrons and polar groups containing sulfur, oxygen and/or nitrogen are fundamental characteristics of this type of inhibitor. The polar functional groups are usually considered the chelation center for chemical adsorption. This adsorption is influenced by the nature and surface charge of metal, the type of aggressive electrolyte and the chemical structure of inhibitors [4, 14-27]. Triazole and its derivatives have been widely studied [28-32] in experimental and theoretical researches. Triazole and triazole-type compounds containing nitrogen, sulphur, and heterocycle on the corrosion inhibition of metal in acidic media have attracted more attention because of their excellent corrosion inhibition performance [33-39]. They are not only used against corrosion but also preferred for biological activities such as antiviral, antibacterial, antifungal and antituberculous. It is known that most of them were reported as green inhibitor due to environmentally friendly effects [28]. The majority of corrosion inhibition studies of steel in H_2SO_4 media using different organic compounds showed less inhibition efficiency than in HCl medium. Interestingly, like HCl, H_2SO_4 is also used in large quantity for removing undesirable scale and rust formed on steel surface especially in boilers, heat exchangers and oil refineries. Hence in these cases the inhibition efficiency of added inhibitor enhanced to greater than 95% by the addition of halide ions especially iodide ions [40]. Synergism is an effective method to improve the inhibitive action of an inhibitor in presence of another substance in corrosive medium and to decrease amount of usage. The addition of halide salts to some inhibitor solution results in synergistic effect. Recently, several studies have been reported to explain role of synergism on corrosion inhibition mechanism of steel in sulphuric acid (H_2SO_4) medium [41-46]. This synergism is due to increased surface coverage as a result of ion-pair interactions between organic cation and halide anion. The halide ions present in an inhibiting solution adsorb on corroding electrode surface by creating oriented dipoles and facilitates the adsorption of inhibitor cations on the dipoles [42]. The synergism between an inhibitor and other ions such as iodide has been extensively studied by many authors [47-49].

In the present investigation, the synergistic influence of iodide ions in the performance of PTT as a corrosion inhibitor of carbon steel in 0.5 M H_2SO_4 has been systematically studied by weight loss measurements, potentiodynamic polarization studies and impedance measurements. The surface morphology of inhibited mild steel was analyzed by scanning electron microscope technology SEM. DFT methods have been used to analyze the characteristics of the inhibitor/surface mechanism and to describe the structural nature of the inhibitor in the corrosion process [12]. Presently, quantum chemical calculations and molecular simulation studies have been proved to be a very useful tool for studying the mechanism of corrosion inhibition. Quantum chemical parameters such as energy of molecular orbitals, highest occupied molecular orbital (HOMO) and lowest unoccupied molecular orbital (LUMO), energy gap (ΔE) between HOMO and LUMO, dipole moment, electronic chemical potential (μ) are directly related to the performance of inhibitors. Results are reported and discussed [50-54].

2. Experimental section

2.1. Materials and sample preparation

The triazole derivative namely PTT has been synthesized in our laboratory as shown in Figure 1 via aromatic carboxylic acid hydrazide intermediate [55, 56]. The aromatic carboxylic acid is first esterified in the presence of concentrated H_2SO_4 and absolute ethanol. The corresponding acid ester is then treated with 80% hydrazine hydrate in the presence of absolute ethanol as solvent that resulted in the formation of corresponding carboxylic acid hydrazide in equimolar proportions. The resulting hydrazide is treated with potassium thiocyanate in acidic medium which resulted in the formation of thiosemicarbazide and the yields were quantitative. Cyclization of the thiosemicarbazide in the presence of NaOH resulted in the formation of the corresponding carboxylic acid-1,2,4-triazole-3-thiol derivative [57]. The obtained compound was purified and analyzed by IR and NMR spectroscopies before use.

The aggressive solutions of 0.5 M H_2SO_4 were prepared by dilution of analytical reagent grade 98 % H_2SO_4 with double-distilled water. The concentration range of the inhibitor employed was 0.1 to 0.5 mM, and the solution in the absence of inhibitor was taken as blank for comparison. The blend inhibitor, PTT+KI, solutions were prepared by adding specified amount of AR grade KI to PTT- H_2SO_4 solution. All experiments were performed in aerated and stagnant solutions. Mild steel specimens containing C \leq 0.214 %, Si \leq 0.237 %, Mn \leq 1.054 %, S \leq 0.006 %, P \leq 0.015 %, Ni \leq 0.066 %, Cu \leq 0.059 % and the remainder iron was used as the working electrode for all studies. Prior to all measurements, the samples were mechanically abraded with 220, 400, 800, 1200 and 2400 emery paper,

decreased with ethanol, washed with double-distilled water and dried at room temperature and stored in vacuum desiccator before immersion in the test solution.

2.2. Methods of study

2.2.1. Weight loss measurements

The weight loss experiments were carried out using cylindrical mild steel specimens (98 %) with exposed total area of 2.58 cm². Mild steel specimens were polished, washed and dried, then weighted and suspended in a 50 ml solution of 0.5 M H₂SO₄ for 1 h in the absence and the presence of various amounts of inhibitor (PTT) and KI. After the elapsed time, the specimens were taken out, washed, dried and weighed accurately. The experimental studies were performed at temperature range of 298 K to 323 K. All weight loss measurements were performed in triplicates. The inhibition efficiency was evaluated from the calculated corrosion rate (CR) using the Eq. (1) [58, 59]:

$$IE (\%) = \frac{CR_0 - CR_{inh}}{CR_0} \times 100 \quad (1)$$

Where CR₀ and CR_{inh} are the corrosion rates of steel (mg cm⁻²h⁻¹) in absence and in presence of inhibitors, respectively.

2.2.2. Electrochemical measurements

Electrochemical experiments were carried out in a glass cell (CEC/TH-Radiometer) with a capacity of 500 ml. A platinum electrode and a saturated calomel electrode (SCE) were used as a counter electrode and a reference electrode. The working electrode (WE) was in the form of a disc cut from mild steel under investigation and was embedded with epoxy steel resin to offer an exposed surface area of 1 cm². Electrochemical impedance spectroscopy (EIS) and potentiodynamic polarization were conducted in an electrochemical measurement system (VoltaLab40) which comprises a PGZ301 potentiostat, a personal computer and Voltmaster 4 and Ec-Lab software. The potentiodynamic current–potential curves were recorded by changing the electrode potential automatically from – 800 to – 200 mV with scanning rate of 0.5 mV s⁻¹. The polarization resistance measurements were performed by applying a controlled potential scan over a small range typically 15 mV with respect to E_{corr}. All experiments were carried out in freshly prepared solution at 303 K. The AC impedance measurements were performed at corrosion potentials (E_{corr}) over a frequency range of 100 kHz – 10 mHz, with a signal amplitude perturbation of 5 mV. Nyquist plots were obtained. The inhibition efficiency was evaluated from the measured I_{corr} values using the Eq. (2) [59, 60]:

$$IE (\%) = \frac{I_{corr(0)} - I_{corr(inh)}}{I_{corr(0)}} \times 100 \quad (2)$$

Where I_{corr(0)} and I_{corr(inh)} are the corrosion current densities in the absence and presence of the inhibitors, respectively.

The inhibition efficiency of the inhibitor has been found out from the charge transfer resistance values using the Eq. (3) [59, 60]:

$$IE (\%) = \frac{R_{ct(inh)} - R_{ct(0)}}{R_{ct(inh)}} \times 100 \quad (3)$$

Where R_{ct(0)} and R_{ct(inh)} are the charge transfer resistance values without and with inhibitors, respectively.

2.2.3. Scanning electron microscopy (SEM)

The mild steel specimens of dimensions, 1.15 cm diameter and 0.15 cm height were abraded with a series of emery papers (grades 220, 400, 800, 1200 and 2400) and then washed with distilled water and ethanol. After immersion in 0.5M H₂SO₄ solution in the absence and the presence of optimum concentration of inhibitor (PTT) and 0.2% KI at 303K for 2h, the specimens were cleaned with double-distilled water, dried with a cold air blaster, and then the SEM images were recorded using ZEISS scanning electron microscope.

2.2.4. Computational details

Theoretical calculations of the compound were carried out using Density Functional Theory (DFT) as implemented in the Gaussian 09 quantum chemistry program. Geometry optimization was started from the X-Ray Diffraction (XRD) experimental atomic coordinates. Initial calculations were performed using the restricted B3LYP exchange and correlation functional and the B3LYP/6-311++G(d,p) basis set for all atoms, Default SCF and geometry convergence criteria were used and no symmetry constraints were imposed, harmonic frequency analysis based on analytical second derivate was used to characterize the optimized geometry as global minimum on the potential energy surface of the title molecule [61-66]. Geometry optimization of the molecule was done at B3LYP/6-311++G(d,p) level of theory then a Single point at the same theory level was performed for the calculations of the molecular orbital energies and dipole moments. The purpose of this work was to investigate whether there is a clear relationship between experimentally determined inhibition efficiency and quantum chemical data for the inhibitors studied. Quantum chemical parameters most relevant to their potential action as corrosion inhibitors such as E_{HOMO} (highest occupied molecular orbital energy), E_{LUMO} (lowest unoccupied molecular orbital energy), energy gap (ΔE), dipole moment (μ), ionization potential (I), electron affinity (A), chemical potential (π), absolute electronegativity (χ), global hardness (η), softness (σ), fraction of electrons transferred (ΔN), electrophilicity index (ω) and such electronic data as Mulliken atomic populations have been calculated and discussed. The ionization potential (I) and the electron affinity (A) are defined as follows deduced from Koopmans' theory [67, 68]:

$$I = -E_{\text{HOMO}} \dots\dots\dots (4)$$

$$A = -E_{\text{LUMO}} \dots\dots\dots (5)$$

Then the electronegativity and the global hardness were evaluated, based on the finite difference approximation, as linear combinations of the calculated I and A [61, 69]:

$$\chi = \frac{I+A}{2} \dots\dots\dots (6)$$

$$\eta = \frac{I-A}{2} \dots\dots\dots (7)$$

Softness is the inverse of hardness:

$$\sigma = \frac{1}{\eta} \dots\dots\dots (8)$$

The obtained values of χ and η are used to calculate the fraction of the electron transferred, ΔN , this parameter evaluates the electronic flow in a reaction of two systems with different electronegativities, in particular case; a metallic surface (Fe) and an inhibitor molecule. ΔN is given as follows:

$$\Delta N = \frac{\chi_{\text{Fe}} - \chi_{\text{inh}}}{2(\eta_{\text{Fe}} + \eta_{\text{inh}})} \dots\dots\dots (9)$$

Where χ_{Fe} and χ_{inh} represent the absolute electronegativity of iron atom (Fe) and the inhibitor molecule, respectively; η_{Fe} and η_{inh} represent the absolute hardness of Fe atom and the inhibitor molecules, respectively. In order to calculate the fraction of electrons transferred, a theoretical value for the electronegativity of iron was employed $\chi_{\text{Fe}} = 7$ eV and a global hardness of $\eta_{\text{Fe}} = 0$ eV, by assuming that for a metallic bulk $I = A$ because they are softer than the neutral metallic atoms [61, 62, 67, 70-72].

Recently, a new global chemical reactivity parameter has been introduced and known as electrophilicity index (ω). It is defined as [66, 67, 70, 71]:

$$\omega = \frac{\chi^2}{2\eta} \dots\dots\dots (10)$$

3. Results and discussions

3.1. Electrochemical impedance spectroscopy (EIS)

Impedance diagrams obtained for frequencies ranging from 100 kHz to 10 mHz at open circuit potential for mild steel in 0.5 M H₂SO₄ in the presence of various concentrations of PTT are shown in Figure 2. All the plots display a single capacitive loop and they are not perfect semicircles. The difference has been attributed to frequency dispersion [73]. The fact that impedance diagrams have a semicircular appearance shows that the corrosion of steel is controlled by a charge transfer process and the presence of inhibitor does not alter the mechanism of dissolution of steel in H₂SO₄ [74-76]. Figure 2 reveals that increase in concentration of PTT results in increase in size of the semicircle, which is an indication of the inhibition of the corrosion process. The equivalent circuit diagram is suggested as in Figure 3, where, R_s represents the solution resistance, R_{ct} the charge transfer resistance, and CPE, the constant phase element. In most times, CPE is required for modeling the frequency dispersion behavior corresponding to different physical phenomena such as surface heterogeneity which results from surface roughness,

impurities, dislocations, distribution of the active sites, adsorption of inhibitor and formation of porous layers [77]. The impedance of CPE is given by the Eq. (11) [78]:

$$Z_{\text{CPE}} = Y_0^{-1} (j\omega)^{-\alpha} \dots\dots\dots (11)$$

Where Y₀ is proportionality coefficient, ω is the angular frequency ($\omega = 2\pi f$) and j is the imaginary number, α is a measure of surface irregularity. Depending on the value of the exponent α , Y₀ may be a resistance, R ($\alpha = 0$); a capacitance, C ($\alpha = 1$); Warburg impedance, W ($\alpha = 0.5$) or an inductance, L ($\alpha = -1$). According to Xiaojuan Wu et al. [79], the value of double layer capacitance, C_{dl}, can be calculated for a parallel circuit composed of a CPE (Y₀) and a resistor (R_{ct}) using the Eq. (12) [78]:

$$C_{\text{dl}} = (Y_0 R_{\text{ct}}^{(1-\alpha)})^{1/\alpha} \dots\dots\dots (12)$$

The values of the charge transfer resistance, R_{ct} double layer capacitance C_{dl} and inhibition efficiency obtained from the above plots are given in Table 1. The impedance data listed in Table 1 indicate that the values of both R_{ct} and IE (%) are found to increase by increasing the inhibitor concentration, while the values of C_{dl} are found to decrease. This behavior was the result of an increase in the surface coverage by the inhibitor molecules, which led to an increase in the inhibition efficiency. The decrease in C_{dl} values may be considered in terms of Helmholtz model [80]:

$$C_{\text{dl}} = \frac{\epsilon_0 \epsilon}{d} A \dots\dots\dots (13)$$

Where ϵ_0 is the permittivity of air, ϵ the local dielectric constant, d the thickness of the film and A is the surface area of the electrode.

In fact, the decrease in C_{dl} values can result from a decrease in local dielectric constant and/or an increase in the thickness of the electrical double layer. It could be assumed that the decrease of C_{dl} values is caused by the gradual replacement of water molecules and other ions that were originally adsorbed on the electrode surface by adsorption of inhibitor molecules to form a protective film, which decreases the extent of the metal dissolution [81]. With higher concentration of PTT, either the thickness of the protective layer or the surface coverage by PTT increased due to more PTT electrostatically adsorbed on the electrode surface [82].

Nyquist plots for mild steel in 0.5M H₂SO₄ in the presence of different concentrations of PTT in combination with 0.2% KI are shown in Figure 4. The high-frequency part of the impedance and phase angles describes the behavior of an inhomogeneous surface layer, while the low-frequency contribution shows the kinetic response for the charge transfer reaction [83]. The impedance spectra, in this case, were fitted by a simple Randles circuit. Table 2 gives the values of the charge transfer resistance R_{ct} obtained from above plots. It is found that the addition of KI further enhances R_{ct} values and reduces C_{dl} values. This can be attributed to the enhanced adsorption of PTT in the presence of KI because of the synergistic effect of iodide ions [84]. The inhibitive effect of iodide ions is often attributed to its

adsorption on the metal surface which depends on its large ionic radius and high hydrophobicity [85]. On the other hand, the exponential term of the CPE between 0.8 and 0.9 indicate that there is no diffusion process normally evidenced by an exponential term of ca. 0.5 [83].

3.2. Polarization measurements

Potentiodynamic anodic and cathodic polarization scans were carried out at 303 K in 0.5 M H₂SO₄ solution with different concentrations of PTT and 0.2 % KI. Figure 5 shows anodic and cathodic polarization curves of mild steel in 0.5 M H₂SO₄ blank solution and in the presence of different concentrations (0.1–0.5 mM) of PTT. The linear Tafel segments of the cathodic and anodic curves were extrapolated to the point of intersection to obtain the corrosion potential (E_{corr}) and corrosion current density (I_{corr}). The electrochemical parameters determined from polarization curve are summarized in Table 3.

It is seen that PTT inhibitor shifts both the cathodic and anodic branches of the polarization curves of the pure acid solution to lower values of current density indicating the inhibition of both the hydrogen evolution and mild steel dissolution reactions. According to Ferreira et al. [86], if the displacement $E_{\text{corr}} > 85$ mV, the inhibitor acts as a cathodic or anodic inhibitor, if the displacement $E_{\text{corr}} < 85$ mV, the inhibitor is considered as mixed type inhibitor [87]. In the present study, in Table 3, the maximum shifting in the E_{corr} value compared to the blank is 90 mV for PTT alone, indicating that the inhibitor studied are a cathodic inhibitor. Some retardation of the anodic reaction is observed but cathodic polarization is clearly dominant. PTT inhibited the hydrogen evolution more than anodic dissolution of mild steel. It can also be seen that PTT decreases the corrosion current densities at all the studied concentrations, meaning that the corrosion rate of steel is reduced significantly.

The addition of KI further reduces the I_{corr} values. Values of inhibition efficiency are found to increase with increase in the concentration of PTT reaching maximum value 91.6 % at 0.5 mM and the addition of KI improved the inhibition efficiency of PTT to 97.7 %. Moreover, the anodic and cathodic Tafel slope values are different from the ones obtained with and without the presence of PTT, respectively, suggesting that the mechanism of the reaction of mild steel in 0.5 M H₂SO₄ is influenced by the presence of PTT. The addition of KI affects slightly the values of β_c and β_a . The maximum shifting in the E_{corr} value in the presence of 0.2% KI is 35 mV. The values of corrosion potential indicate that a mixed-type control and PTT is an inhibitor of mixed-type with predominant effect on the cathodic reaction for the corrosion of mild steel in 0.5 M H₂SO₄. The inhibition efficiency obtained from polarization measurement is in good agreement with results obtained from EIS tests.

3.3. Weight loss measurements

Values of the inhibition efficiency obtained from the weight loss measurements of mild steel for different concentrations of PTT in 0.5 M H₂SO₄ at 303 K in the absence and the presence of 0.2% KI after 1 h of immersion [1-9]. PTT inhibit the corrosion of mild steel in 0.5 M H₂SO₄ solution at all concentrations used in this study and the corrosion rates (CR) is seen to decrease continuously with increasing additive concentration at 303 K. Indeed, corrosion rate values of mild steel decrease when the inhibitor concentration increases while IE (%) values of PTT increase with the increase of the concentration, the maximum IE (%) of 91.62 % is achieved at 0.5 mM [58, 59]. The inhibition of corrosion of mild steel by PTT can be explained in terms of its adsorption on the metal surface. This compound can be adsorbed on the metal surface by the interaction between lone pairs of electrons of nitrogen and sulfur atoms of the inhibitor and the metal surface. This process is facilitated by the presence of vacant orbitals of low energy in iron atom, as observed in the transition group metals [88-90]. Moreover, the formation of positively charged protonated species in acidic solutions facilitates the adsorption of the compound on the metal surface through electrostatic interactions between the organic molecules and the metal surface. On the other hand, it can be seen from Table 4 that the addition of KI in the solution improved the inhibition efficiency of PTT significantly. The synergistic effect between PTT and KI can be due to coulombic interactions between chemisorbed I⁻ and organic polycations PTT⁺. The stabilization of adsorbed PTT⁺ on the iron surface which may be caused by electrostatic interactions with I⁻ ions leads to more surface coverage and consequently greater corrosion inhibition.

3.4. Effect of immersion time on corrosion of mild steel

In order to assess the stability of inhibitive behavior of inhibitor on a time scale, weight loss measurements were performed in 0.5 M H₂SO₄ in absence and presence 0.5 mM of PTT and 0.2 % KI for different immersion time at temperature 303 K. The table 5 gives the obtained results. This table shows that the inhibition efficiency of PTT in absence and presence KI decreased with immersion time. Shriver et al. [91-93] and Ishtiaque et al. [94] explained that decrease in inhibition for long period of immersion can be attributed to the depletion of available inhibitor molecules in the solution due to chelate formation between iron and the inhibitor ligands.

3.5. Synergism parameters

Halides ions have been widely reported to synergistically increase the inhibition of the corrosion of metals in acidic medium by organic compounds [95-106]. The enhanced corrosion inhibition on addition of the halides

was attributed to promotion of the adsorption of the organic molecules on the metal surface. The effect of KI on the inhibition efficiency of PTT inhibitor was investigated by EIS measurements in the present work at 303 K. Table 6 gives values of the inhibition efficiency for the corrosion of mild steel in 0.5M H₂SO₄ in the presence of 0.5 mM PTT at 303K and different concentrations of KI. It can be seen that the maximum synergistic effect is obtained for a KI concentration of 0.2%. Figure 6 shows the plot of different values of R_{ct} for all concentrations of PTT inhibitor in the absence and presence of 0.2% KI at 303 K after one hour of immersion. Figure 6 reveals clearly the existence of synergism phenomenon between iodide ions and PTT molecules considering that R_{ct} was increased on addition of the iodide ions to PTT inhibitor in comparison to PTT alone.

To further confirm whether synergism is taking place, one has to determine the synergism parameter (S), the synergism parameters were calculated using the relationship proposed by Aramaki and Hackerman [97, 102]:

$$S = \frac{1 - IE_{1+2}}{1 - IE'_{1+2}} \dots\dots\dots (14)$$

Where, $IE_{1+2} = (IE_1 + IE_2) - (IE_1 \times IE_2)$; IE_1 is the inhibition efficiency (no exprimed in percent) of substance 1 (PTT), IE_2 is the inhibition efficiency of substance 2 (KI), IE'_{1+2} is the measured inhibition efficiency for substance 2 in combination with substance 1. S approaches 1 when no interaction between the inhibitor compounds exists, while $S > 1$ points to a synergistic effect. In the case of $S < 1$, the adsorption of each compound antagonizes the others adsorption [102, 107]. Values of S are given in Table 7.

It can be seen from this table that most of values are greater than unity; except for the lower concentration. This result indicates that the increase in inhibitor efficiency is generated by the addition of KI to PTT due to a synergistic effect and this may be related to adsorption of iodide ions on metal surface that caused the increase of adsorption of inhibitor. In addition, the synergistic effect is more pronounced for the optimal concentration. However, a careful inspection of this table suggests that a competitive adsorption appears for the lower concentration of PTT [108].

Globally, the inhibition efficiency measured for different methods was estimated at about 91.6% in the absence of 0.2% KI and 97% in the presence of KI, the inhibition efficiency obtained from weight loss and electrochemical measurements are in good agreement (Figure 7).

3.6. Effect of temperature

In order to study the effect of temperature on the inhibition characteristic of triazole derivative (PTT), weight loss measurements were performed at different temperatures from 298 to 323 K in the absence and presence of different additives for 1 h immersion time. The results are given in Table 8.

It can be seen that, the inhibition efficiencies of PTT decrease with increase in the temperature in the absence and presence of 0.2% KI. However, these values decrease more rapidly (around 15 %) with temperature in the case of PTT alone, which indicated desorption of inhibitor molecules to some extent with increasing temperature [1]. The activation energy (E_a) for the corrosion process in the absence and presence of PTT was evaluated from Arrhenius equation [59, 78, 109]:

$$\text{Log CR} = -\frac{E_a}{2.303 RT} + \text{log A} \dots\dots\dots (15)$$

Where CR is the corrosion rate, E_a is the apparent activation energy, A the pre-exponential factor, R the universal gas constant (8.314 J K⁻¹mol⁻¹), T the absolute temperature. The Arrhenius plots for mild steel immersed in 0.5 M H₂SO₄ are shown in Figure 8 for a blank solution and in the presence of PTT and (PTT + KI). Straight lines are obtained. The values of E_a obtained from the slopes of the lines ($-E_a/2.303R$) are listed in Table 9.

It is evident that the value of E_a is higher for inhibited solution than that for uninhibited solution and the value of E_a for the mixture (PTT + KI) is the highest one. The increase in E_a values confirms stronger physisorption of the inhibitor on the mild steel surface, which is facilitated by the presence of KI. Physisorption is small but important because it is the preceding stage of chemisorption. The triazole molecules create a barrier to charge and mass transfer. The increase in E_a values might also be correlated with the increased thickness of double layer [110]. To calculate enthalpy and entropy of activation for corrosion process transition state the Eq. (16) was used [59, 78, 111]:

$$\text{CR} = \frac{RT}{Nh} \exp\left(\frac{\Delta S^*}{R}\right) \exp\left(-\frac{\Delta H^*}{RT}\right) \dots\dots\dots (16)$$

Where h is the Planck's constant, N is the Avogadro's number, ΔS^* is the entropy of activation and ΔH^* is the enthalpy of activation. Figure 8b, showed the plot of log (CR/T) against 1/T. The plots obtained are straight lines and the values of ΔH^* are calculated from their gradient ($\Delta H^* = -\text{slope} \times 2.303R$) and ΔS^* from intercept [$\text{log}(R/Nh) + \Delta S^*/2.303R$]. The calculated data listed in Table 9.

The positive signs of ΔH^* reflect the endothermic nature of mild steel dissolution process in the presence of PTT. The value of ΔS^* is higher for inhibited solution than that for the uninhibited solution, suggesting an increase in randomness on going from reactants to the activated complex. The increase in values of entropy by the adsorption of PTT molecules on metal surface from the acid solution could be regarded as quasi-substitution between the inhibitor in the aqueous phase and H₂O molecules on electrode surface. In such condition, the adsorption of PTT molecules was followed by desorption of H₂O molecules

from the electrode surface. Thus increase in entropy of activation is attributed to solvent (H₂O) entropy [86, 112]. In the case of the mixture (PTT + KI) the value of ΔS^* is the highest one. This can be attributed to the gradual replacement of water molecules by the iodide ions and by the adsorption of the PTT molecules on the metal surface, decreasing the extent of dissolution reaction.

3.7. Adsorption isotherm

Basic information on the interaction between the inhibitor and the mild steel surface is obtained from various isotherms. Adsorption of this organic inhibitor has displaced the water molecules from the metal surface. The most commonly used adsorption isotherms are Langmuir, Temkin, and Frumkin. Langmuir adsorption isotherm was found to be best fit. The degree of surface coverage (θ) for different concentrations of inhibitor in 0.5 M H₂SO₄ solution has been evaluated by the polarization measurements. According to the Langmuir adsorption isotherm, the surface coverage (θ) is related to the inhibition concentration C_{inh} by the Eq. (17) [59, 60]:

$$\frac{C_{inh}}{\theta} = \frac{1}{K_{ads}} + C_{inh} \dots\dots\dots (17)$$

Where C_{inh} is the inhibitor concentration and K_{ads} is the equilibrium constant for adsorption-desorption process.

The plots of C_{inh}/θ versus C_{inh} (Figure 9) gives straight lines for all temperatures with a slope of unity is observed and the linear correlation coefficient is 0.999 which suggests that the adsorption obeys Langmuir adsorption isotherm. The adsorption of PTT inhibitor on the metal surface can occur on the basis of donor-acceptor interactions between the π -electrons of the heterocyclic compound and the lone pairs of the hetero-atoms and vacant d-orbitals of iron surface atoms [113-115]. From the intercept, K_{ads} is calculated, and by using its values, the values of ΔG_{ads}^0 were calculated by the Eq. (18) [59, 60]:

$$\Delta G_{ads}^0 = -RT \ln (55.5 K_{ads}) \dots\dots\dots (18)$$

Where K_{ads} is the adsorptive equilibrium constant, ΔG_{ads}^0 is the free energy of adsorption, R is the gas constant and T is the absolute temperature. The value of 55.5 is the molar concentration of water in solution in mol. L⁻¹ [116].

The addition of KI did not change the adsorption behavior of PTT. It can be concluded that the inhibition actions of (PTT+ KI) are mainly attributed to the adsorption of PTT. The PTT molecules may have stronger adsorption abilities than iodide ions, as indicated by their inhibition efficiencies.

On the other hand, the values of the free energy of adsorption as calculated from the Eq. (18) in the absence and the presence of KI are -36.2 and -38.1 kJ.mol⁻¹, respectively. The largest negative values of ΔG_{ads}^0 ensure the spontaneity of the adsorption process and stability of the adsorbed layer on the steel surface and also indicate that PTT is strongly adsorbed on the mild steel surface and this adsorption is greater in the presence of KI [117]. Generally,

values of ΔG_{ads}^0 , around -20 kJ.mol⁻¹ or lower are consistent with the electrostatic interaction between the charged molecules and charge metal, such as physisorption. When it is around -40 kJ.mol⁻¹ or higher values it involve charge sharing or charge transfer from organic molecules to the metal surface to form a coordinate type of bond that is chemisorption [118, 119]. The calculated values range from -30 to -40 kJ.mol⁻¹. This indicates that PTT is adsorbed chemically and physically on steel surface in 0.5M H₂SO₄ solution. The unshared electron pairs of heteroatom interact with d-orbital of iron atom of steel to provide a protective chemisorbed film [120].

PTT may adsorbed on a metal surface in the form of a neutral molecule via the chemisorption mechanism [88] involving the sharing of electrons between the nitrogen, sulphur atom and iron. The second mode (physisorption) is possible if one examines also the activation energy (E_a) that increases in the presence of PTT. Adsorption of PTT can also occur through π electron interactions between the triazole group structure of molecules and the metal surface. Therefore, we may suggest that the adsorption may occur through the lone pairs of heteroatoms and π electrons of the PTT molecules which outweigh the adsorption due to the cationic form of the PTT molecule on the metal surface. We can conclude that adsorption acts simultaneously by chemisorption and physical adsorption [121].

3.8. Scanning electron microscopy (SEM)

Figure 10 represents the SEM images of mild steel samples under various conditions studied in this work. Figure 10(a) is the image of newly polished sample before placing in 0.5 M H₂SO₄. Figure 10(b) shows the same sample after immersing in the blank solution for 2h at 303 K. The appearance of corrosion products is clearly observed. Figure 10(c) was taken from a sample in acid solution containing PTT alone and finally Figure 10(d) represents the metal sample after immersing in solution containing both PTT (0.5 mM) and potassium iodide (0.2%). It can be seen from Figure 10(b) that the mild steel surface was strongly damaged with deep cavities in the absence of the inhibitor due to metal dissolution in corrosive solution. The image of steel surface exposed to PTT containing solution (Figure 10c) exhibits good surface and no damage, the sample has smooth surface compared with that of the sample immersed in 0.5 M H₂SO₄ with few small notches. No damages expect polishing lines are observed in the micrograph after the addition of iodide ions to the corrosive solution (Figure 10d). The steel surface in the presence of KI + PTT mixture (Figure 10d) is the smoothest surface and thus the mixture is a more effective inhibitor. The resemblance of the Figure 10(d) to that of (a) is appreciable and proves the excellent corrosion prevention of PTT and iodide ions combined. The later effect is due to synergistic interactions between PTT and iodide ions resulting in higher inhibition efficiency. This

indicates that the combination of PTT and KI hinders the dissolution of iron and thereby reduces the rate of corrosion, and it reveals good protection against corrosion.

3.9. DFT calculations

The inhibitory effect of the inhibitors regularly depends on the adsorption of these molecules on the metal surface, this adsorption depending on the molecular structures. Several quantum chemical methods and molecular modeling techniques are very often used in corrosion inhibition studies in order to ascertain the relationship between corrosion inhibition efficiency and molecular properties, and consequently, to estimate the trends in the binding energy between inhibitors chemisorbed and/or physisorbed on metal surfaces [52, 61, 62, 66, 122, 123]. To investigate the relationship between the molecular structure of this triazole derivative (PTT) and its inhibition effect, quantum chemical calculations were performed. DFT is a very powerful technique to probe the inhibitor/surface interaction and to analyze experimental data. DFT methods have become very popular in the last decade due to their accuracy that is similar to other methods in less time and with a smaller investment from the computational point of view [124]. The optimized structures of the triazole derivative in the neutral form including their HOMO and LUMO distributions density at the B3LYP/6-31G(d,p) level are presented in Figure 11, the graphics was done thanks to GABEDIT. Positive and negative phase is represented in red and blue color, respectively.

Quantum chemical parameters such as: E_{HOMO} , E_{LUMO} , ΔE , μ , χ , η , σ , ω and ΔN affect the inhibitory effectiveness [71]. These quantum chemical characteristics for the estimated structure are calculated from equations 4 to 10 and presented in Table 10. The frontier molecular orbitals play an important role in the electric and optical properties [125]. The HOMO represents the ability to donate an electron, LUMO as an electron acceptor. Inspection of Table 10 reveals that the calculated HOMO and LUMO energies are, respectively, -7.731, -0.260, the gap is 7.471, The negative values of E_{HOMO} indicate that adsorption of this inhibitor on the steel surface is important [70]. Generally, the higher values of E_{HOMO} indicate an increase for the electron donor and this means a better inhibitory activity with increasing adsorption of the inhibitor on a metal surface, whereas E_{LUMO} indicates the ability to accept electron of the molecule and strong electronic interactions between inhibitor and metal surface, this later is -0.260 eV indicates the easiness for this molecule to receipt electrons from the d orbital of the metal. The adsorption ability of the inhibitor to the metal surface increases with increasing of E_{HOMO} and decreasing of E_{LUMO} . Therefore high E_{LUMO} and low E_{LUMO} values are associated with high electron donating ability of inhibitor and thereby increasing the corrosion Inhibition efficiency [126, 127]. Large values of the energy

gap (ΔE) imply high electronic stability and then low reactivity; whereas low values imply that it will be easier to remove an electron from the HOMO to the LOMO, which can result in highly efficient inhibition. These results showed that the mechanism of inhibition is based primarily on an adsorption process performed by charge transfer from the inhibitor to the metal [70]. Furthermore, less negative HOMO energy and the smaller energy gap ($\Delta E_{\text{HOMO-LUMO}}$) are reflected in a stronger chemisorption bond and perhaps greater inhibition efficiency [90, 128]. This can be possible because removing electrons from HOMO needs a small energy. This confirms the obtained results and it is in a good agreement with the values of ΔN [129]. Low values of the energy gap (ΔE) will provide good inhibition efficiencies because the excitation energy to remove an electron from the last occupied orbital will be low. A molecule with a low energy gap is more polarizable and is generally associated with a high chemical reactivity. The calculation in Table 10 shows that PTT has smaller energy gap value indicating that PTT can easily adsorb on the metal surface causing higher protection. According to literature, there is a lack of agreement on the correlation between dipole moment and inhibition mechanism. The high value of dipole moment probably increases the adsorption between a chemical compound and metal surface [130]. Meanwhile, a survey of the literature reveals that several irregularities have appeared in the case of correlation of the dipole moment with inhibitor efficiency [131, 132]. However, increasing values of E_{HOMO} and μ may facilitate adsorption by influencing the transport process through the adsorbed layer and lead to better inhibition efficiency [90]. From Table 10, the high compound polarity (dipole moment equal to 4.381 Debye) which is higher than the dipole moment H_2O (1.88 Debye). The high dipole moment value of this compound probably indicates strong dipole-dipole interactions between them and the metal surface and facilitates electrostatic interaction between the electric field, due to the metal charge and the electric moment of this inhibitor, and contributes to their better adsorption [133]. Therefore, a quasi-substitution process occurs between molecules of PTT and water ones on the steel area, with desorption of water molecules from the metal. Thus, the inhibition process can be done [134].

High ionization energy ($I = 7.731$ eV) indicates high stability, the number of electrons transferred (ΔN) was also calculated and tabulated in Table 10. The fraction of electrons transferred (ΔN) values describes the inhibition achieved from electron donations. The values (ΔN) are correlated with the inhibition efficiency resulting from the electron donation. According to Lukovits [66], if (ΔN) < 3.6, the inhibitory efficiency is increased by increasing the ability inhibitor to give electrons to metal surface, while it decreased if $\Delta N > 3.6$ (electron). It is apparent from Table 10 that ΔN value is positive and is below 3.6, this means that the inhibitor (PTT) has the tendency to donate electrons to

the metal surface by chemisorption process [62]. The inhibitory power is also influenced by the value of the electrophilicity index (ω). The inhibitor effectiveness increases with the decrease of the ω value. The value of ω (2.137) agrees with the high value of the protection rate [134]. The electronegativity value of the inhibitor molecule is equal to 3.996. This electronegativity value ($\chi=3.996$) being inferior to that of iron suggests an exchange of electron between the high occupied molecular orbital of the inhibitor (HOMO) and the unoccupied 3d orbitals of iron. This electron transfer is more probable than that from the occupied 4s orbital of Fe to the low unoccupied molecular orbital (LUMO) of the inhibitor [71]. Absolute hardness (η), and softness (σ), is important properties to measure the molecular stability and reactivity. It is apparent that the chemical hardness fundamentally signifies the resistance towards the deformation or polarization of the electron cloud of the atoms, ions or molecules under small perturbation of chemical reaction. A hard molecule has a large energy gap and a soft molecule has a small energy gap [135]. In our present study PTT with low hardness value +3.736 eV compared with other compound have a low energy gap. Normally, the inhibitor with the least value of global hardness (hence the highest value of global softness) is expected to have the highest inhibition efficiency. For the simplest transfer of electron, adsorption could occur at the part of the molecule where softness (σ), which is a local property, has a highest value. PTT with the softness value of +0.268 eV has a highest inhibition efficiency [136].

The HOMO and LUMO electronic density distributions of PTT are plotted in Figure 11. It can be observed that for the molecule of PTT, HOMO and LUMO are distributed around heteroatoms and aromatic rings, indicating that these heteroatoms and aromatic rings on which negative charge density is high, are main adsorption centers of the triazole derivative by the metal surface, by

formation of donor–acceptor coordination bonds with vacant "d" orbitals of the iron [124]. Thus, the bond with the metal from these active sites will be easily formed, rather than the others atoms. The inhibition efficiency of the compounds depends on many major factors, for example the number of adsorption active centres in the molecule and their charge density, molecule size, mode of adsorption, and formation of metallic complexes. The effect of substituents in the compounds on inhibition of corrosion of mild steel by PTT will be rationalized by use of global reactivity indexes, in the form of the Mulliken charges. Theoretical calculation of the charge density on the heteroatoms in the molecular structure of PTT reinforces the suggestion that the inhibitory effect results from the groups constituting the rigid centre of the molecule and their adsorption by the surface of the mild steel. This distribution thus ensures strong adsorption by the metal surface [70]. Mulliken charge analysis is used to estimate the adsorption centers of inhibitors, the results were presented on the atoms in the optimized structure (Figure 11). It is possible to observe that the heteroatoms afford a noticeable excess of negative charges could act as a nucleophilic reagent [62, 137]. The effective atomic charges from Mulliken populations presented in table 11 show that the highest negative charges were located on S (-0.484), N1 (-0.301) and N4 (-0.008) atoms as well as some carbon atoms of the aromatic rings C8 (-0.533), C10 (-0.311) and C11 (-0.595). These atoms are the more negative charge centers that could give electrons to the Fe atoms in order to coordinate it, and the positive charge centers that can accept electrons from 3d orbital of the Fe atoms to form feedback bond, thus further strengthening the interaction of inhibitor and Fe surface. This implies that PTT has potential heteroatoms that can be adsorbed in positive centers on the metallic surface through a donor-acceptor type reaction [138].

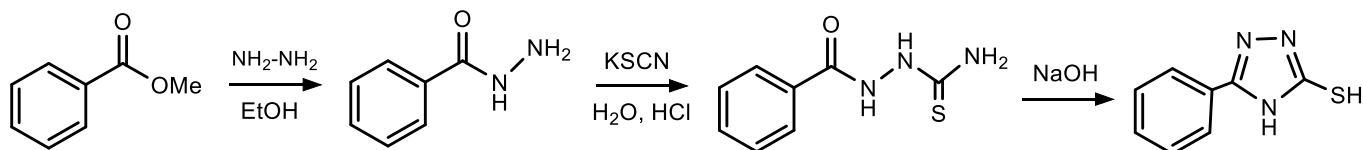


Figure 1: Scheme of synthesis of 5-Phenyl-4H-1,2,4-triazole-3-thiol (PTT)

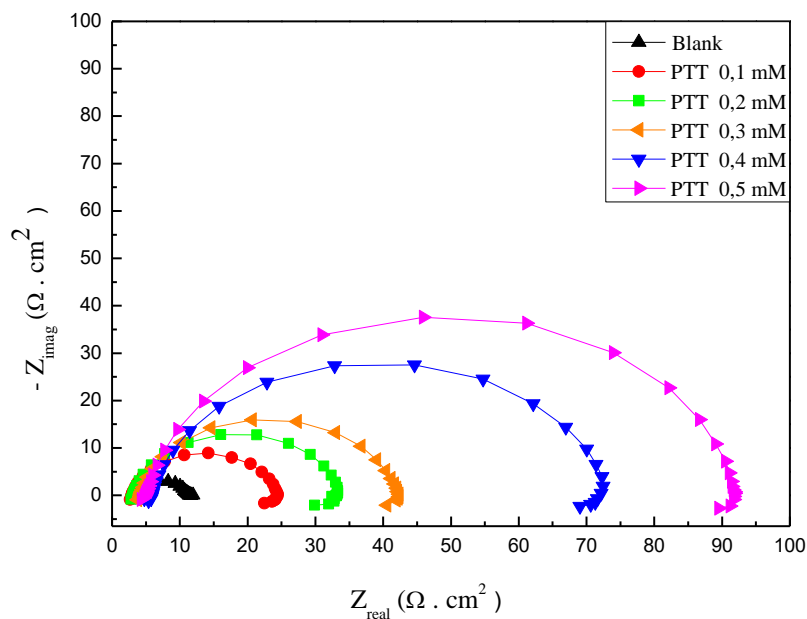


Figure 2. Nyquist plots for mild steel in 0.5 M H₂SO₄ containing different concentrations of PTT

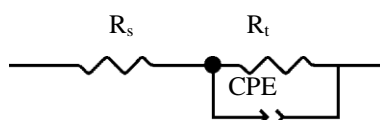


Figure 3. The equivalent circuit of the impedance spectra obtained for PTT in 0.5M H₂SO₄

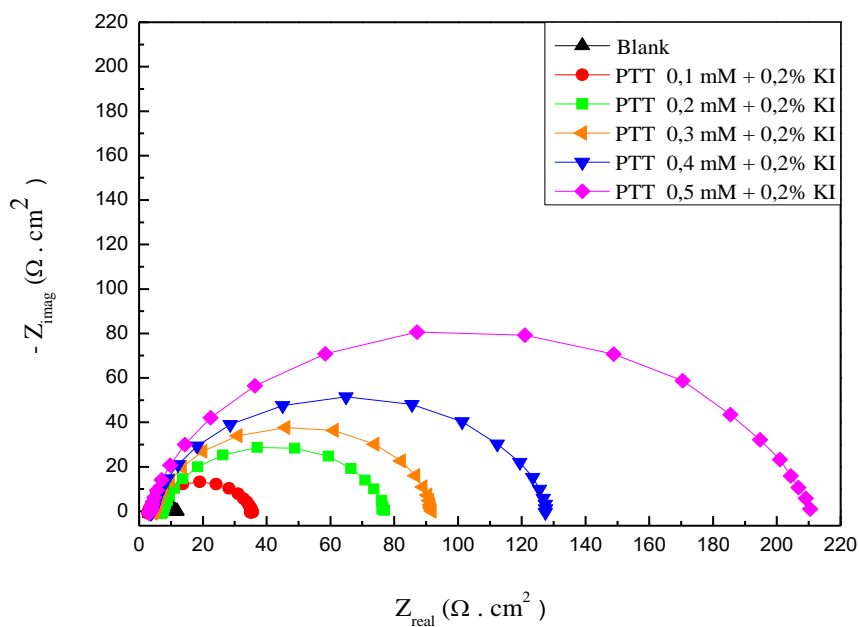


Figure 4. Nyquist plots for mild steel in 0.5M H₂SO₄ containing different concentrations of PTT in the presence of KI 0.2 %

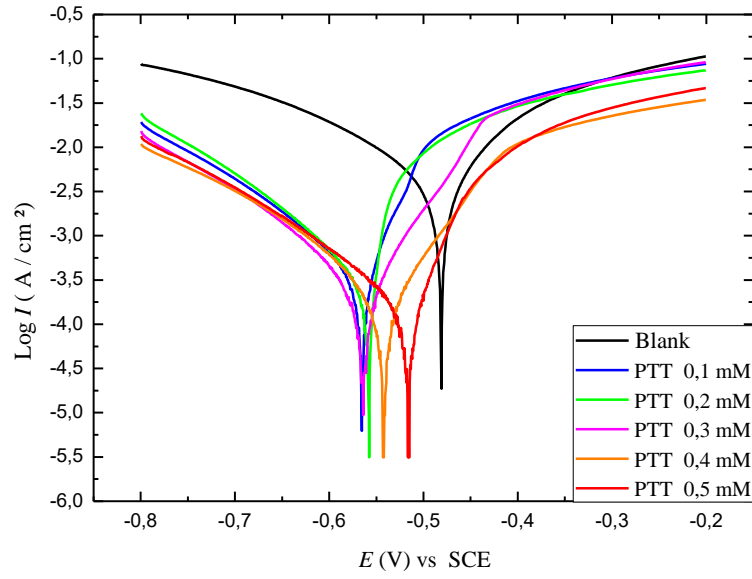


Figure 5. Potentiodynamic polarization curves of mild steel in 0.5M H₂SO₄ for different concentrations of PTT at 303 K.

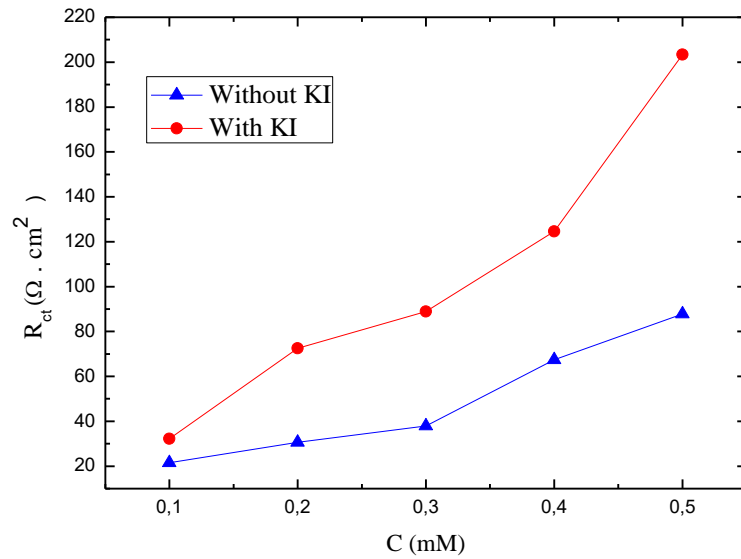


Figure 6. R_{ct} values in the absence and presence of 0.2% KI for different concentrations of PTT

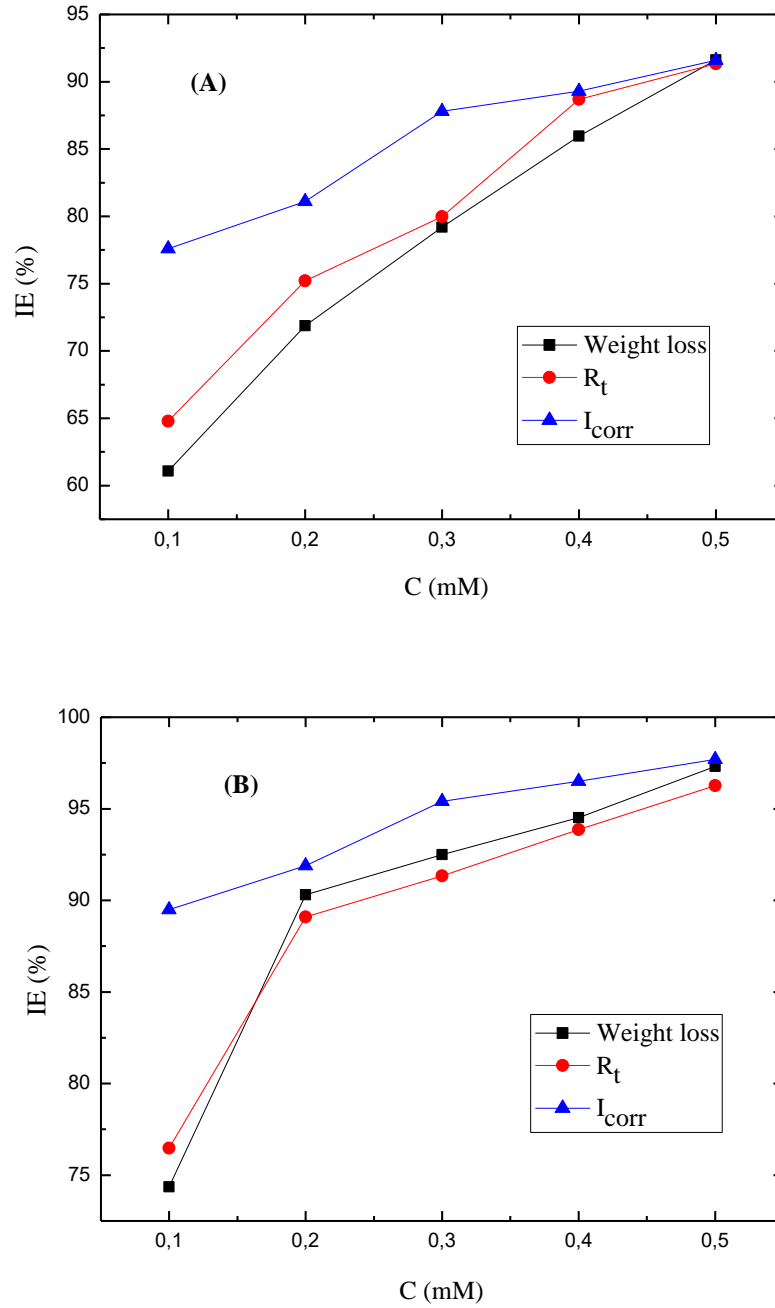


Figure 7. Inhibition efficiency vs inhibitor concentrations calculated by different methods: (A) in the absence of 0.2% KI, (B) in the presence of 0.2% KI

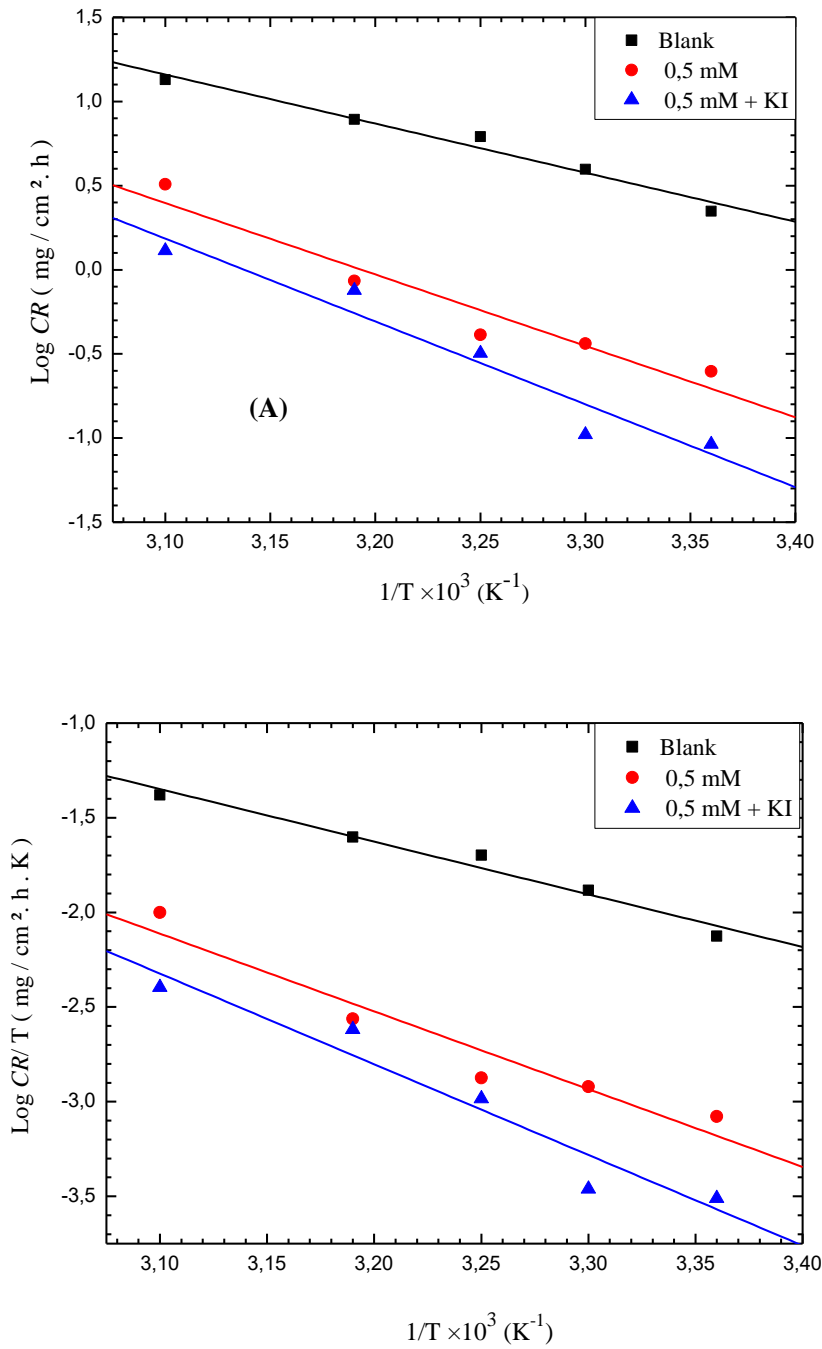


Figure 8. Arrhenius plots of: (a) log CR vs. 1 / T; (b) log (CR / T) vs. 1 / T for the mild steel in 0.5 M H₂SO₄ solution in the absence and presence of 0.5 mM PTT and 0.5 mM PTT + 0.2% KI

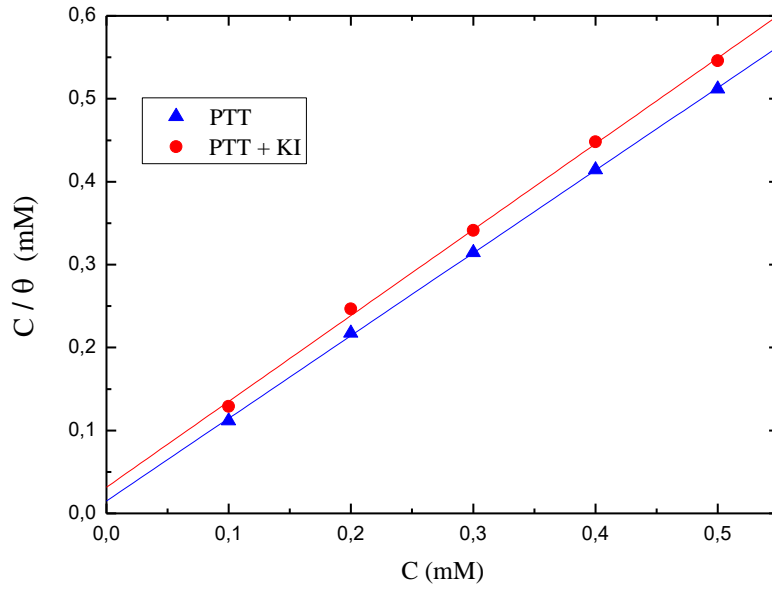


Figure 9. Langmuir's adsorption isotherm of PTT and (PTT + KI) on the mild steel surface in 0.5M H₂SO₄ from polarization measurements

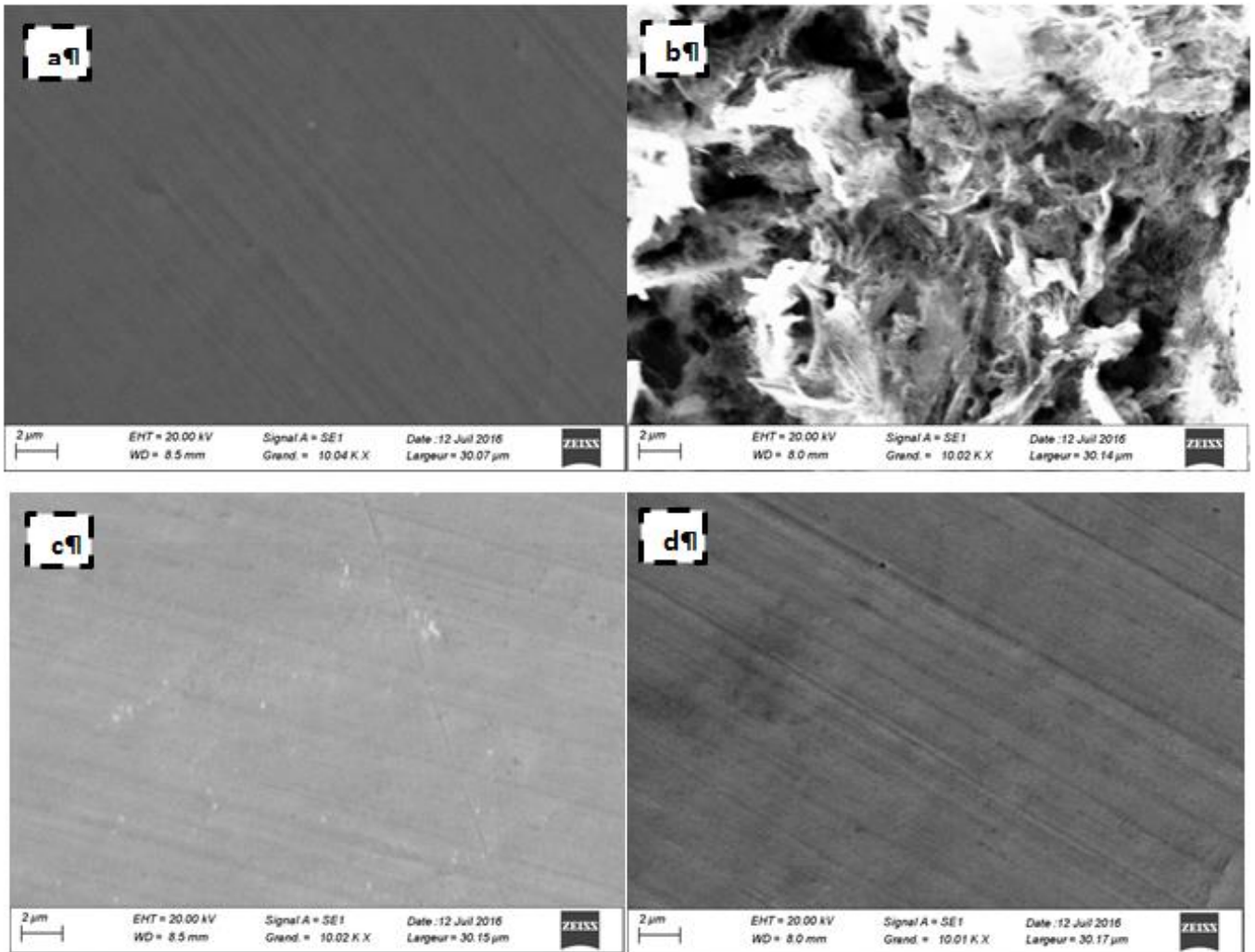


Figure 10. SEM image of the surface of mild steel after immersion for 2 h in 0.5 M H₂SO₄ solution at 303K: (a) Before immersion (polished) (b) After immersion without inhibitor (c) With inhibitor (PTT) (d) With inhibitor and in presence of KI

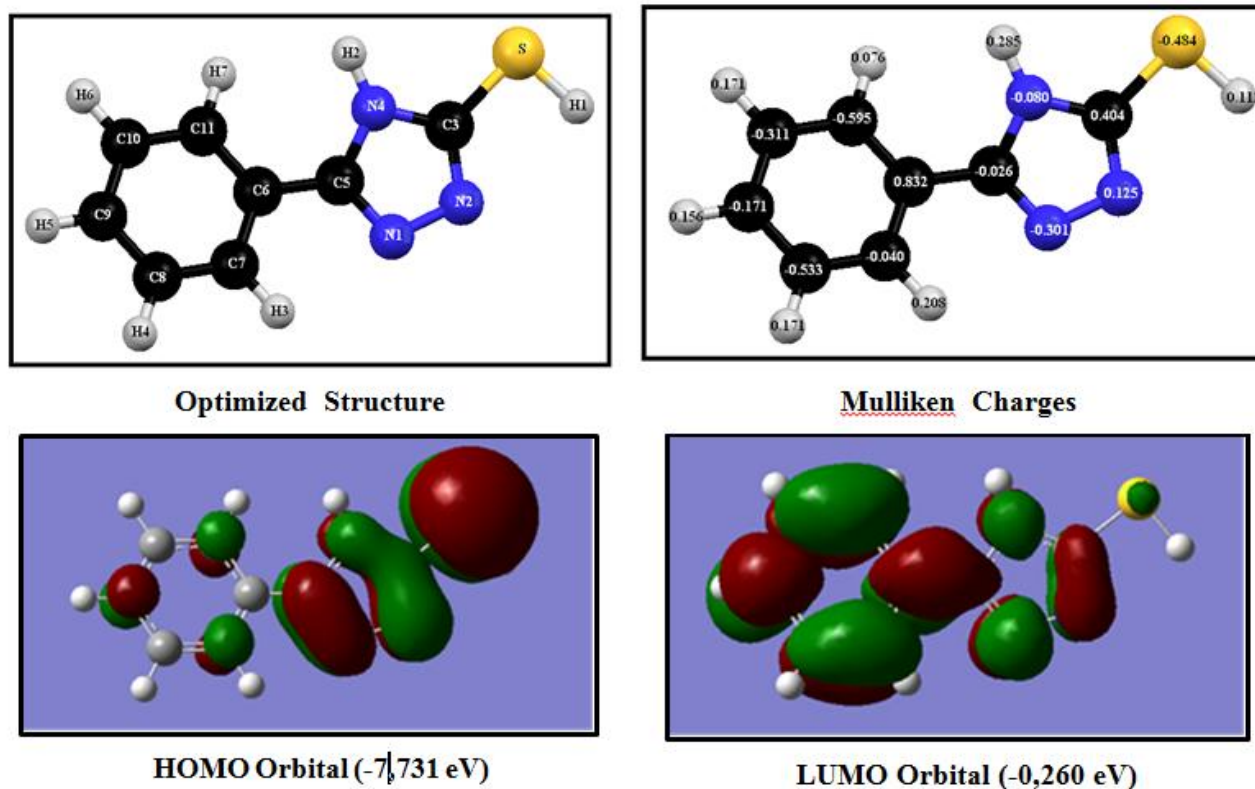


Figure 11. Optimized structure, Mulliken charges density and Frontier molecule orbital density distributions: HOMO (left) and LUMO (right) of PTT given by the B3LYP/6-31(d,p)

Table 1. Electrochemical parameters of mild steel in the presence of PTT in 0.5M H₂SO₄ solution deduced from EIS data

Conc. (mM)	R _s (Ω cm ²)	R _{ct} (Ω cm ²)	A	Y ₀ × 10 ⁴ (S ⁿ Ω ⁻¹ cm ⁻²)	C _{dl} , (μ F cm ⁻²)	IE (%)
0.5M H ₂ SO ₄	3.659	7.584	0.866	5.804	251.2	–
0.1	2.932	21.54	0.881	3.349	172.3	64.79
0.2	3.109	30.61	0.892	1.892	101.4	75.22
0.3	4.223	37.86	0.902	1.397	79.04	79.97
0.4	5.887	67.06	0.892	1.357	76.86	88.69
0.5	5.046	87.49	0.898	1.317	69.5	91.33

Table 2. Electrochemical parameters of mild steel in the presence of PTT in combination with 0.2% of KI in 0.5M H₂SO₄ solution deduced from EIS data

Conc. (mM)	R _s (Ω cm ²)	R _{ct} (Ω cm ²)	α	Y ₀ × 10 ⁴ (S ⁿ Ω ⁻¹ cm ⁻²)	C _{dl} , (μ F cm ⁻²)	IE (%)
0.1	3.031	32.23	0.874	2.531	126.4	76.47
0.2	5.531	69.51	0.888	1.376	88.44	89.09
0.3	4.698	87.47	0.899	1.316	79.56	91.33
0.4	4.222	123.8	0.883	1.228	73.57	93.87
0.5	3.340	203.6	0.874	1.092	63.02	96.27

Table 3. Potentiodynamic polarization parameters for corrosion of mild steel in 0.5M H₂SO₄ with various concentrations of PTT in the absence and presence of 0.2% KI at 303 K

Conc. (mM)	Conc. KI (%)	E _{corr} (mV vs SCE)	I _{corr} (μA/cm ²)	β _c (mV/dec)	β _a (mV/dec)	IE (%)
Blank	–	-482	3463	261.5	152.2	----
0.1	–	-572	776	205.2	117.3	77.6
0.1	0.2	-517	363	180.9	93.9	89.5
0.2	–	-558	653	167.7	102.6	81.1
0.2	0.2	-549	280	148.6	85.2	91.9
0.3	–	-564	422	147.0	91.3	87.8
0.3	0.2	-539	157	133.3	76.5	95.4
0.4	–	-540	369	167.5	101.0	89.3
0.4	0.2	-506	122	111.6	104.7	96.5
0.5	–	-516	292	181.9	95.5	91.6
0.5	0.2	-497	80	142.6	41.2	97.7

Table 4. Corrosion parameters obtained from weight loss measurements for mild steel in 0.5 M H₂SO₄ containing various concentrations of PTT in the presence of 0.2 % KI at 303K

Conc. PTT (mM)	Conc. KI (%)	CR (mg cm ⁻² h ⁻¹)	IE (%)
0.1	–	1.5421	61.07
0.1	0.2	1.0158	74.36
0.2	–	1.1141	71.88
0.2	0.2	0.3837	90.31
0.3	–	0.8239	79.20
0.3	0.2	0.2970	92.50
0.4	–	0.4772	85.95
0.4	0.2	0.2169	94.52
0.5	–	0.3318	91.62
0.5	0.2	0.1047	97.36

Table 5. Effect of immersion time on inhibitor performance for mild steel in 0.5M H₂SO₄ at 303 K for 0.4 mM and 0.5 mM PTT

Immersion time (min)	IE (%)	
	0.5 mM PTT	0.5 mM PTT + 0.2% KI
60	91.62	97.36
120	89.94	95.21
180	86.97	92.53
240	85.29	90.96
360	81.42	86.37
1440	61.77	65.71

Table 6. Inhibition efficiency for different concentrations of KI for mild steel in 0.5 M H₂SO₄ in the presence of 0.5 mM PTT at 303 K

KI (%)	IE (%)
0.05	94.78
0.1	95.24
0.15	96.24
0.2	97.36
0.25	97.30
0.3	95.52

Table 7. Values of synergism parameters (S) for different concentrations of PTT

Conc. PTT (mM)	S
0.1	0.76
0.2	1.45
0.3	1.39
0.4	1.10
0.5	1.73

Table 8. Effect of temperature on the inhibition efficiency *IE* (%) obtained from weight loss measurements

Temperature (K)	IE (%)	
	PTT (0.5 mM)	PTT (0.5 mM) + KI 0.2 %
298	88.84	95.88
303	91.62	97.36
308	90.82	94.83
313	89.05	90.37
323	76.04	90.35

Table 9. Thermodynamic activation parameters for mild steel in 0.5 M H₂SO₄ solution and in the presence of PTT and (PTT+KI) obtained from weight loss measurements

	E _a (kJ mol ⁻¹)	ΔH* (kJ mol ⁻¹)	ΔS*(J mol ⁻¹ K ⁻¹)
0.5 M H ₂ SO ₄ (Blank)	55.83	53.4	-57.92
0.5 mM PTT	81.23	79.35	7.97
0.5 mM PTT + 0.2% KI	94.28	91.84	42.53

Table 10. Molecular properties of PTT obtained from the optimized structure using DFT at the B3LYP/6-31G

inhibitor	E _{HOMO} (eV)	E _{LUMO} (eV)	ΔE (eV)	μ (Debye)	I (eV)	A (eV)	χ (eV)	η (eV)	σ (eV ⁻¹)	ω (eV)	ΔN (eV)
PTT	-7.731	-0.260	7.471	4.381	7.731	0.260	3.996	3.736	0.268	2.137	0.402

Table 11. Molecular properties of PTT obtained from the optimized structure using DFT at the B3LYP/6-31G

Atoms	Mulliken charges
N1	-0.301
N2	0.125
C3	0.404
N4	-0.080
C5	-0.026
C6	0.832
C7	-0.040
C8	-0.533
C9	-0.171
C10	-0.311
C11	-0.595
S	-0.484
H1	0.112
H2	0.285
H3	0.208
H4	0.171
H5	0.156
H6	0.171
H7	0.076

4. Conclusions

The inhibition behavior of PTT and its synergistic effect with KI for mild steel in 0.5 M H₂SO₄ has been studied. The following conclusions may be drawn:

- PTT showed excellent inhibition properties for the corrosion of mild steel in 0.5 M H₂SO₄ solution. The inhibition efficiency of PTT increased with the concentration and reached a maximum of 91.6% at 0.5 mM.
- Changes in impedance parameters were indicative of adsorption of PTT on the metal surface.
- PTT acts as a mixed-type inhibitor retarding the

anodic and cathodic corrosion reactions with predominant effect on the cathodic reaction.

- The inhibition efficiency IE% of PTT decreased with increasing temperature and its addition led to increase of the activation corrosion energy.
- Adsorption of PTT alone and in combination with KI on the mild steel surface obeys Langmuir's adsorption isotherm. The calculated value of ΔG^o_{ads} obtained from the study indicates that PTT is adsorbed chemically and physically on steel surface in 0.5M H₂SO₄ solution. The compound is adsorbed with the heteroatoms forming donor-acceptor bonds between unpaired electrons of the

heteroatoms and the active centers of the metal surface.

- Synergistic effects between PTT and KI were observed. The addition of KI to the solution enhanced the inhibition efficiency of PTT significantly. The adsorption of PTT is stabilized by the presence of iodide ions in 0.5 M H₂SO₄.
- The inhibitor efficiencies determined by polarization, EIS, and weight loss methods are in good agreement.
- SEM micrographs of mild steel specimens showed that the inhibitor molecules form a good protective film on the metal surface.
- The values of free energy of adsorption and the calculated quantum chemical suggest that the inhibition behavior of this triazole derivative involves two types of interaction, chemisorption and physisorption.
- Theoretical calculations provide good support to the experimental results.

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