

# ELECTROCHEMICAL INVESTIGATION OF INHIBITORY OF NEW SYNTHESIZED TETRAZOLE DERIVATIVE ON CORROSION OF STAINLESS STEEL 316L IN ACIDIC MEDIUM

A. Ehsani<sup>1,2\*</sup>, Sh. Bodaghi<sup>2</sup>, H. Mohammad Shiri<sup>2</sup>, H. Mostaanzadeh<sup>1</sup> and M. Hadi<sup>1</sup>

\* ehsani46847@yahoo.com

Received: August 2015

Accepted: August 2016

<sup>1</sup> Department of Chemistry, Faculty of science, University of Qom, Qom, Iran.

<sup>2</sup> Department of Chemistry, Payame Noor University, Iran.

**Abstract:** In this study, an organic compound inhibitor; namely *N*-benzyl-*N*-(4-chlorophenyl)-1*H*-tetrazole-5-amine (NBTA), was synthesized and the role of this inhibitor for corrosion protection of stainless steel (SS) exposed to 0.5 M H<sub>2</sub>SO<sub>4</sub> was investigated using electrochemical, and quantum analysis. By taking advantage of potentiodynamic polarization, the inhibitory action of NBTA was found to be mainly mixed type with dominant anodic inhibition. The effectiveness of the inhibitor was also indicated using electrochemical impedance spectroscopy (EIS). Moreover, to provide further insight into the mechanism of inhibition, quantum chemical calculations of the inhibitor were performed. The adsorption of NBTA onto the SS surface followed the Langmuir adsorption model with the free energy of adsorption  $\Delta G_{ads}$  of -7.88 kJ mol<sup>-1</sup>. Quantum chemical calculations were employed to give further insight into the mechanism of inhibition action of NBTA.

**Keywords:** organic inhibitor, adsorption, stainless steel, impedance, Density functional theory.

## 1. INTRODUCTION

Since iron and its alloys are the backbone of industrial constructions, many research projects have been concerned with their stability. One of most important tasks is the retardation of the attack by acid solutions used during pickling, industrial cleaning and descaling. The use of an additive is one of the major solutions for this problem. Hence, various additives are used to protect iron and its alloy against corrosive attack. A variety of organic compounds, particularly azole derivatives containing heteroatoms such as nitrogen and sulphur, aromatic rings and  $\pi$  bonds has been already examined to inhibit the corrosion of metals [1-6]. The existing data show that organic inhibitors act by adsorption and they protect the metal by film formation. Organic compounds bearing heteroatoms with high electron density such as phosphorus, sulfur, nitrogen, oxygen or those containing multiple bonds which are considered as adsorption centers, are effective as corrosion inhibitors [6-10]. The compounds containing both nitrogen and sulfur in their molecular structure have exhibited greater inhibition compared with those possessing only one of these atoms [11-13]. In

the literature, many thiazole derivatives [14-16] have been reported as corrosion inhibitors and found to have good corrosion inhibition effect. The efficiency of an organic compound inhibitor is mainly dependent on its ability to adsorb on a metal surface, which consists of replacement of a water molecule at a corroding interface. The present work aims to evaluate the inhibitory effect of new synthesized NBTA on the steel corrosion in acidic solution. Different aspects of electrochemical measurements, including potentiodynamic polarization and electrochemical impedance spectroscopy, were used to study the inhibition mechanism of the inhibitor. Moreover, to provide further insight into the mechanism of inhibition, quantum chemical calculations of the inhibitor based on DFT method were performed.

## 2. EXPERIMENTAL

### 2. 1. Materials and Apparatus

Stainless steel 316L has the composition (wt %) Fe: 67.95, Ni: 10.60, Si: 0.45, Mn: 1.75, Cr: 16.50, S: 0.025, P: 0.028, Mo: 2.10, Al: 0.008, Co: 0.16, Cu: 0.35, Nb: 0.01 and V: 0.02. The

exposed surface of the SS electrode was ground with silicon carbide abrasive paper from 400 to 1200, degreased with absolute ethanol, rinsed in distilled water, and dried in warm air. The corrosive medium was 0.5 M  $\text{H}_2\text{SO}_4$  solution prepared from analytical reagent grade 98% sulfuric acid and distilled water. For synthesizing the organic inhibitor, a mixture of N-benzyl-N-(4-chlorophenyl) cyanamide, sodium azide and  $\text{ZnCl}_2/\text{AlCl}_3/\text{Silica}$  in distilled DMF was heated up to 120 °C for 3h and stirred under ultrasonic irradiation.

The concentration range of inhibitor employed was  $10^{-4}$  to  $10^{-3}$  M in 0.5 M sulfuric acid. All electrochemical measurements were carried out in a conventional three electrode cell, powered by a potentiostat/galvanostat (EG&G 273A) and a frequency response analyzer (EG&G, 1025). The system was run by a PC through M270 and M398 software via a GPIB interface. The frequency range of 100 kHz to 15 mHz and modulation amplitude of 5 mV were employed for impedance studies. A SS electrode was employed as the working electrode. A saturated calomel electrode (SCE) and a platinum wire were used as reference and counter electrodes, respectively. Before measurement, the working electrode was immersed in test solution for approximately 1 h until a steady open circuit potential (OCP) was reached. The polarization curves were carried out from cathodic potential of -1.4 V to anodic potential of 0.10 V with respect to the open circuit potential at a sweep rate of 0.5 mV/s. The linear Tafel segments of the anodic and cathodic curves were extrapolated to corrosion potential ( $E_{\text{corr}}$ ) to obtain the corrosion current densities ( $i_{\text{corr}}$ ). In each measurement, a fresh working electrode was used. Several runs were performed for each measurement to obtain reproducible data.

## 2. 2. Computational Details

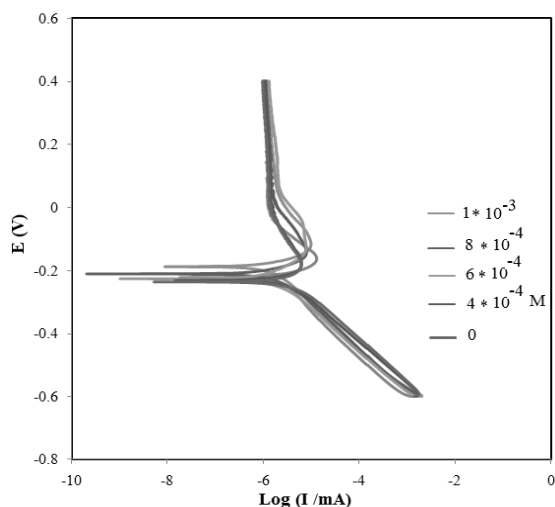
The use of quantum chemical calculations has become popular for screening new potential corrosion inhibitors [17]. Theoretical calculations were carried out at density functional theory (DFT) level using the 6-31G (d, p) basis set for all atoms with Gaussian 03 program package.

Electronic properties such as highest occupied molecular orbital (HOMO) energy, lowest unoccupied molecular orbital (LUMO) energy and frontier molecular orbital coefficients have been calculated. The molecular sketches of all compounds were drawn using Gauss View 03 [18]. The natural bond orbital (NBO) analysis, suggested by Reed et al. [19, 20], was applied to determine the atomic charges.

## 3. RESULTS AND DISCUSSION

### 3. 1. Potentiodynamic Polarization Studies

Polarization measurements were carried out to get information regarding the kinetics of anodic and cathodic reactions. The potentiodynamic polarization curves for SS in 0.5 M  $\text{H}_2\text{SO}_4$  solution in the absence and presence of different concentrations of the inhibitor molecules are shown in Fig. 1. The values of electrochemical kinetic parameters such as corrosion potential ( $E_{\text{corr}}$ ), corrosion current ( $I_{\text{corr}}$ ) and Tafel slopes, determined from these by extrapolation method, are listed in Table 1. In corrosion, quantitative information on corrosion currents and corrosion potentials can be extracted from



**Fig. 1.** Potentiodynamic polarisation curves of SS in 0.5 M  $\text{H}_2\text{SO}_4$  solution in the absence and presence of various concentrations of the NBTa.

**Table 1.** Corrosion parameters obtained from Tafel polarisation curves of SS in 0.5M H<sub>2</sub>SO<sub>4</sub> in the absence and presence of different concentrations of NBTA at 298 K.

No.	Inhi. Con.	Ba(v/decade)	Bc(v/decade)	I(uA)	E(v)	CR(mpy)
SS1	0	0.61	0.14	12.76	-0.22	5.25
SS2	0.0001	0.16	0.14	5.78	-0.24	2.38
SS3	0.0002	0.16	0.14	3.71	-0.23	1.53
SS4	0.0004	0.17	0.14	2.5	-0.22	1.03
SS5	0.0006	0.09	0.14	2.22	-0.21	0.913
SS6	0.0008	0.04	0.16	2.21	-0.22	0.912
SS7	0.001	0.06	0.15	1.73	-0.18	0.711

the slope of the curves, using the Stern-Geary equation, as follows [21]:

$$i_{corr} = \frac{1}{2.303R_p} \left( \frac{\beta_a \times \beta_c}{\beta_a + \beta_c} \right) \quad (1)$$

$i_{corr}$  is the corrosion current density in Amps/cm<sup>2</sup>;  $R_p$  is the corrosion resistance in ohms cm<sup>2</sup>;  $\beta_a$  is the anodic Tafel slope in Volts/decade or mV/decade of current density;  $\beta_c$  is the cathodic Tafel slope in Volts/decade or mV/decade of current density; the quantity,  $\frac{\beta_a \times \beta_c}{\beta_a + \beta_c}$ , is referred to as the Tafel constant. The corrosion inhibition efficiency was calculated using the relation:

$$\eta(\%) = 100 \left( \frac{i_{corr}^* - i_{corr}}{i_{corr}^*} \right) \quad (2)$$

where  $i_{corr}^*$  and  $i_{corr}$  are uninhibited and inhibited corrosion current densities, respectively, determined by extrapolation of Tafel lines in the corrosion potential. The corrosion rates  $v$  (mm year<sup>-1</sup>) from polarization were calculated using the following Equation:

$$v = \frac{i_{corr} \times t \times M}{F \times S \times d} \times 10 \quad (3)$$

where  $t$  is the time (s),  $M$  is the equivalent molar weight of working electrode (g mol<sup>-1</sup>),  $F$  is Faraday constant (96500 °C. mol<sup>-1</sup>),  $S$  is the surface area of electrode,  $d$  is the density of iron and the constant 10 is used to convert the unit cm to mm. The results are presented in table 1. The inhibitor molecule first adsorbs on the SS surface and blocks the available reaction sites. As the concentration of the inhibitor increases, the linear polarization resistance increases and corrosion Rate (CR) decreases. The surface coverage increases with the inhibitor concentration and the formation of inhibitor film on the SS surface reduces the active surface area available for the attack of the corrosive medium and delays hydrogen evolution and metal dissolution [22]. In the cathodic domain, as seen in table 1, the values of  $\beta_c$  show small changes with increasing inhibitor concentration, which indicates that the NBTA is adsorbed on the metal surface and the addition of the inhibitor hinders the acid attack on the SS electrode. In anodic domain, the value of  $\beta_a$  decreases with the presence of NBTA. The shift in the anodic Tafel slope  $\beta_a$  might be attributed to the modification of anodic dissolution process due to the inhibitor modules adsorption on the active sites. Compared to the absence of NBTA, the anodic curves of the working electrode in the acidic solution containing the NBTA clearly shifted to the direction of current reduction, as it could be observed from these polarization results; the

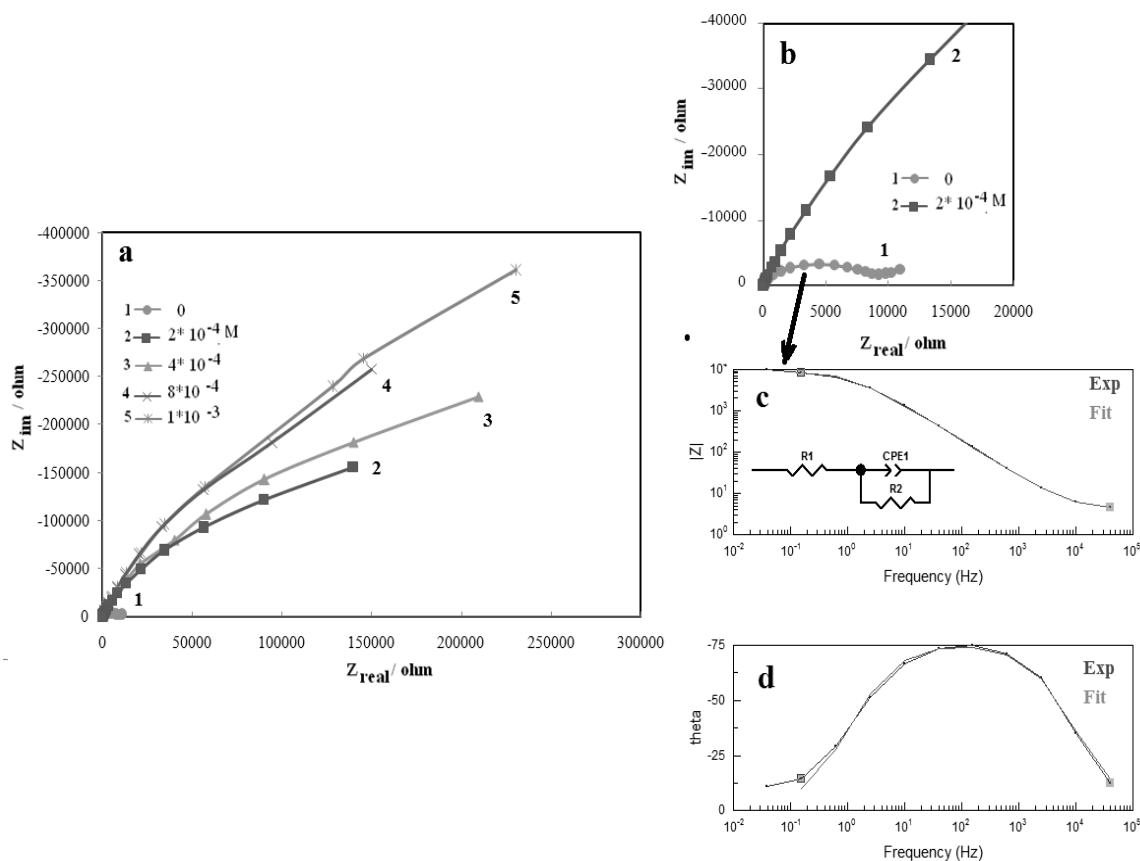


Fig. 2. (a, b) Nyquist plots of SS in 0.5 M H<sub>2</sub>SO<sub>4</sub> solution containing different concentrations of the NBTA in different magnification. (c) Electrical equivalent circuit used for fitting.

inhibition efficiency increased with inhibitor concentration.

### 3. 2. Electrochemical Impedance Spectroscopy

Electrochemical impedance spectroscopy is one of the best techniques for analyzing the properties of conducting polymer electrodes and charge transfer mechanism in the electrolyte/electrode interface. This has been broadly discussed in the literature using a variety of theoretical models [23-27]. Impedance measurements were performed under potentiostatic conditions after 1 h of immersion. Nyquist plots of uninhibited and inhibited solutions containing different concentrations of inhibitor molecules were performed over the frequency range from 100 kHz to 100 mHz and are shown in Fig. 2. The similarity in the shapes

of these graphs throughout the experiment indicates that the addition of inhibitor molecules does not cause any noticeable change in the corrosion mechanism [22]. The Nyquist diagrams show one capacitive loop at high frequencies. The capacitive loop at high frequencies represents the phenomenon associated with the electrical double layer. The above impedance diagrams (Nyquist) contain depressed semicircles with the centre under the real axis. Such behavior is characteristic of solid electrodes and often referred to as frequency dispersion, attributed to different physical phenomena such as roughness, inhomogeneities of the solid surfaces, impurities, grain boundaries, and distribution of surface active sites. The ideal capacitive behavior is not observed in this case and hence a constant phase element CPE is introduced in the circuit to give a more accurate

fit [28-30]. Obviously, the corrosion behavior of SS in acidic solution is influenced to some extent by mass transport since the Warburg impedance is observed in Nyquist plots. Anodic dissolution of SS is mass-transport limited and leads to the formation of sulfate compound. However, the diffusion step was ascribed either to sulfate ions transport to the surface or to the transport of formed compound diffusion to the bulk solution. The mechanism of corrosion remains unaffected during the addition of inhibitor molecules. The simplest equivalent circuit is represented in Fig. 2c, which is a parallel combination of the charge transfer resistance ( $R_{ct}$ ) and the constant phase element (CPE), both in series with the solution resistance ( $R_s$ ). The impedance function of a CPE can be represented as:

$$Z_{CPE} = Y_0^{-1} (j\omega)^{-n} \quad (4)$$

where  $Y_0$  is the CPE constant,  $\omega$  is the angular frequency, and  $n$  is the CPE exponent, which can be used as a gauge of the heterogeneity or roughness of the surface [25-31]. Warburg element in equivalent circuit presents Warburg impedance. In the present work, the value of  $n$  has a tendency to decrease with increasing inhibitor

concentration, which may be attributed to the increase of inhibitor concentration resulted in the increasing surface roughness. For a circuit including a CPE, the  $C_{dl}$  could be calculated from CPE parameter values  $Y_0$  and  $n$  using the expression [22]:

$$C_{dl} = Y_0 (\omega_m'')^{-n} \quad (5)$$

where  $C_{dl}$  is the double layer capacitance and  $\omega_m''$  is the frequency at which the imaginary part of the impedance has a maximum. As seen in table 2, the double layer capacitance ( $C_{dl}$ ) decreases with increase in concentration. This can be attributed to the gradual replacement of water molecules by the adsorption of the organic molecules at metal/solution interface, which is leading to a protective film on metal surface. In addition, the more the inhibitor is adsorbed, the more the thickness of the barrier layer increases according to the expression of the Helmholtz model [32]:

$$C_{dl} = \frac{\epsilon\epsilon_0 A}{d} \quad (6)$$

where  $d$  is the thickness of the protective layer,  $\epsilon$  is the dielectric constant of the medium,  $\epsilon_0$  is the

**Table 2.** Impedance parameters for the corrosion of SS in 0.5M H<sub>2</sub>SO<sub>4</sub> containing different concentration of NBTA at 298 K.

Concentration x (M)	$R_s$ ( $\Omega$ )	n	$R_{ct}$ ( $k\Omega$ )	IE%
0	4.17	0.88	7.81	
$2.0 \times 10^{-4}$	7.65	0.85	347.70	97.75
$4.0 \times 10^{-4}$	14.12	0.84	413.50	98.11
$6.0 \times 10^{-4}$	16.65	0.84	587.87	98.67
$8.0 \times 10^{-4}$	17.24	0.82	614.39	98.72
$1.0 \times 10^{-3}$	17.19	0.80	687.14	98.86

vacuum permittivity and A is the surface area of the electrode. The equation used for calculating the percentage inhibition efficiency is:

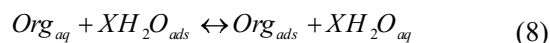
$$\eta(\%) = 100 \left( \frac{R_{ct}^* - R_{ct}}{R_{ct}^*} \right) \quad (7)$$

where  $R_{ct}^*$  and  $R_{ct}$  are values of the charge transfer resistance observed in the presence and absence of inhibitor molecules. Impedance parameters are summarized in table 2. The results obtained from the EIS technique in acidic solution were in good agreement with those obtained from the polarization method. As observed in table 2, the adsorption of NBTA molecules on SS surface modifies the interface between the corrosive medium and metal surface and decreases its electrical capacity. The increase in  $R_{ct}$  values with increase in NBTA concentration can be interpreted as the formation of an insulated adsorption layer. At the highest inhibitor concentration of  $10^{-3}$  mol/L, the inhibition efficiency markedly increases and reaches 98.86%. Thus, it can be deduced that NBTA has a clear role in metal protection at the concentration of  $10^{-3}$  M.

### 3. 4. Adsorption Isotherms

The adsorption of an organic adsorbate at metal/solution interface can be presented as a

substitution adsorption process between the organic molecules in aqueous solution, ( $Org_{aq}$ ), and the water molecules on metallic surface, ( $H_2O_{ads}$ )  $Org_{aq}$



where X, the size ratio, is the number of water molecules displaced by one molecule of organic inhibitor. X is assumed to be independent of coverage or charge on the electrode [33]. Basic information on the interaction between the inhibitors and the steel surface is provided by the adsorption isotherm. The degree of surface coverage,  $\theta$ , at different inhibitor concentrations in 0.5 M  $H_2SO_4$  ( $\theta = IE(\%)/100$ ) at 25 °C. The plot of  $C/\theta$  against inhibitor concentration, C, displayed a straight line for the tested inhibitor (Fig. 3). The linear plot clearly revealed that the surface adsorption process of NBTA on the SS surface obeys the Langmuir isotherm. Likewise, it suggests that an adsorption process occurs, which can be expressed as follows [34]:

$$\frac{C}{\theta} = \frac{1}{K_{ads}} + C \quad (9)$$

where  $K_{ads}$  is the equilibrium constant of the adsorption process. Free energy of adsorption

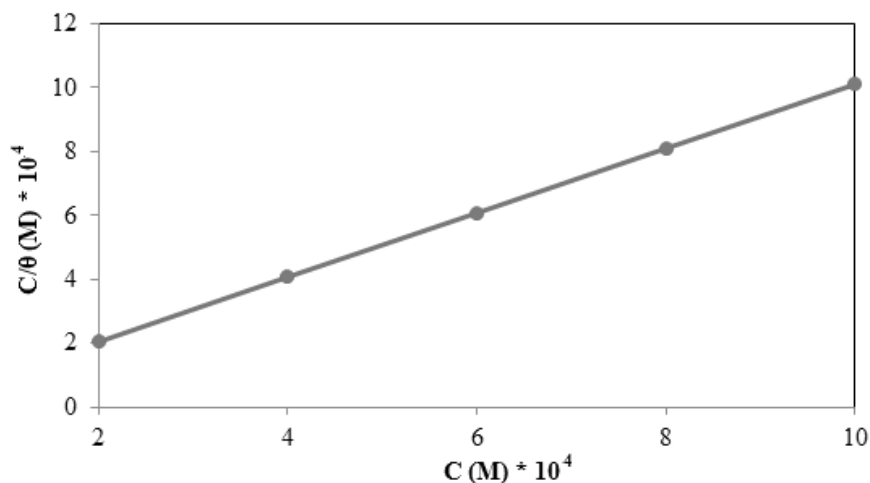


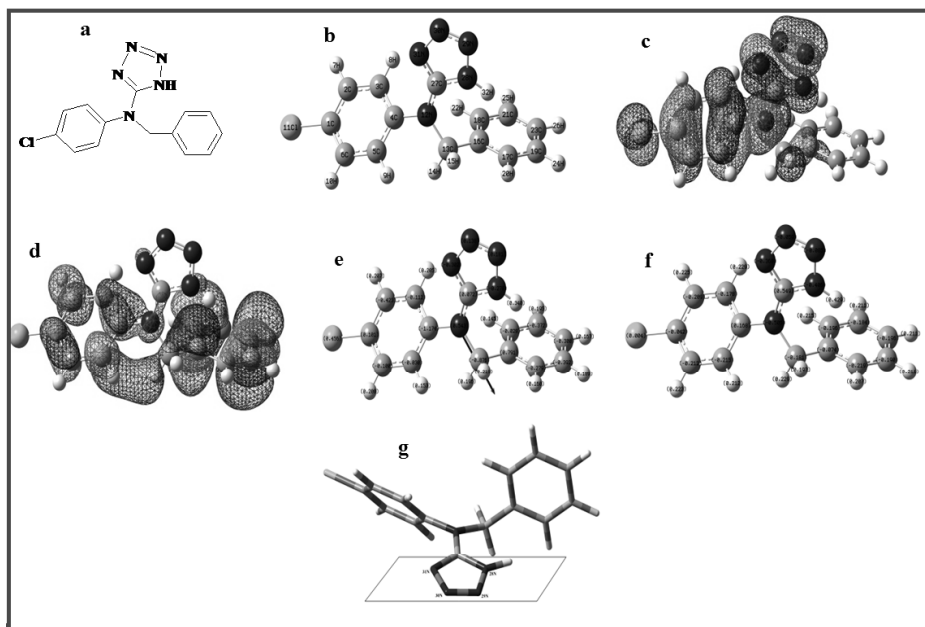
Fig. 3. Langmuir adsorption plot for SS in 0.5 M  $H_2SO_4$  containing different concentrations of NBTA.

( $\Delta G_{\text{ads}}$ ) can be calculated by Eq. (10). The numeral of 55.5 is the molar concentration of water in the solution:

$$K_{\text{ads}} = \frac{1}{55.5} \exp\left(\frac{-\Delta G_{\text{ads}}^0}{RT}\right) \quad (10)$$

The value of  $\Delta G^0_{\text{ads}}$  for adsorption of NBTA was found to be  $-7.88 \text{ kJ mol}^{-1}$ . The negative value of  $\Delta G^0_{\text{ads}}$  suggests that NBTA is spontaneously adsorbed on the SS surface. Literature survey reveals that the values of  $\Delta G^0_{\text{ads}}$  around  $-20 \text{ kJ mol}^{-1}$  or lower are consistent

with the electrostatic interaction between the charged molecules and the charged metal (physical adsorption) [35]. The adsorption of an inhibitor on the metal surface can occur on the basis of donor-acceptor interactions between the p-electrons of the heterocyclic compound and the vacant d-orbitals of the metal surface atoms. Therefore, the energies of the frontier orbitals should be considered. Energy of LUMO shows the ability of the molecule to receive charge when attacked by electron pair donors, even as the energy of HOMO to donate the charge when attached by electron seeking reagents. As the energy gap between the frontier orbitals gets



**Fig. 4.** (a) Structure of NBTA; (b) Optimized molecular structure of NBTA, H atoms have been omitted for clarity; (c) The highest occupied molecular orbital (HOMO) of NBTA; (d) The lowest unoccupied molecular orbital (LUMO) of NBTA; (e) Mulliken charge population analysis and vector of dipole moment of NBTA; (f) Natural charge population analysis of NBTA and (g) The schematic representation of the adsorption behavior of NBTA on the surface of the SS.

**Table 3.** Orbital energies for HOMO, LUMO, HOMO-LUMO gap energy ( $\Delta E$ ) and dipole moment ( $\mu$ ) of Compound in the gaseous phase. All quantum chemical parameters calculated at DFT level using the 6-31G(d,p) basis set.

Compound	phase	$E_{\text{HOMO}}(\text{eV})$	$E_{\text{LUMO}}(\text{eV})$	$\Delta E(\text{eV})$	$\mu(\text{D})$
NBTA	G	-6.395	-1.396	4.999	7.3769
	A	-6.496	-1.153	5.343	9.6801

**Table 4.** Electronegativity ( $\chi$ ), global hardness ( $\eta$ ) and proportion of electrons transferred ( $\Delta N$ ) of NBTA.

Compound	phase	$\chi$	$\eta$	$\Delta N$
NBTA	G	3.895	2.499	0.621
	A	3.824	2.671	0.594

**Table 5.** Mulliken and Natural Charges (e) for atoms in NBTA.

NBTA	Mulliken	Natural
12N	0.543	-0.505
28N	-0.278	-0.406
29N	0.181	-0.085
30N	0.138	-0.059
31N	-0.302	-0.368

smaller, the interactions between the reacting species strengthen [36]. In this regard, the electronic properties such as highest occupied molecular orbital (HOMO) energy, lowest unoccupied molecular orbital (LUMO) energy and frontier molecular orbital coefficients have been calculated for prepared inhibitor. The natural bond orbital (NBO) analysis was applied to determine the atomic charges. Results are presented in figure 4 and tables 3, 4, 5. According to the results, HOMO location in the NBTA molecule is mostly distributed in the vicinity of the nitrogen, oxygen atoms. This indicates the reactive sites of the interaction between NBTA and the SS surface. Mulliken population analysis, presented in figure 4e, is further evidence for the interaction between SS surface and inhibitor active sites. It is clear from figure 4 that the nitrogen atoms of NBTA have considerable excess of negative charge than other atoms. Thus, the adsorption of NBTA as a neutral molecule on the metal surface can occur directly involving the displacement of water molecules from the metal surface and sharing of electrons between the nitrogen atoms and the metal surface. It should be noted that NBTA adsorbs mainly through electrostatic interactions between the positively charged nitrogen atom (since acidic solution can protonate the nitrogen atoms of NBTA and the

negatively charged metal surface (physisorption) as evident in the value of  $\Delta G^0_{ads}$  obtained.

#### 4. CONCLUSION

In the case of NBTA as a corrosion inhibitor for stainless steel in acidic solution, the results of potentiodynamic polarization, electrochemical impedance spectroscopy and quantum chemical analysis was studied in this work. The following points can be emphasized:

1. The potentiodynamic polarization revealed NBTA as a mixed-type corrosion inhibitor with a predominantly anodic action.
2. The results of quantum chemical analysis supported those from the electrochemical measurements relating to the effectiveness of the inhibitor.
3. The nitrogen atoms of NBTA have considerable excess of negative charge than other atoms. Thus, the adsorption of NBTA as a neutral molecule on the metal surface can occur directly involving the displacement of water molecules from the metal surface and sharing of electrons between the nitrogen atoms and the metal surface.
4. The adsorption of DAT on the surface was found to be in accordance with the Langmuir isotherm. By taking advantage of this isotherm, the adsorption process was suggested to take place through electrostatic interaction (physisorption) between inhibitor and iron.

#### 5. ACKNOWLEDGEMENTS

We gratefully acknowledge the support of this work by Qom University Research Council.

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