Effect of processing time on microstructure of surface and corrosion resistance of coatings resulting from plasma electrolytic oxidation on titanium alloy in hydroxyapatite nano-particles electrolyte

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Abstract. This study investigated the effect of coating processing time on the microstructure of surface and corrosion resistance of coatings resulting from Plasma Electrolytic Oxidation (PEO) on the substrate of TiAl6V4 alloy. The coating processes in hydroxyapatite nano-powder electrolytic were carried out under the condition of constant voltage of 600 V and three different times of 125, 250, and 350 seconds. Studying the microstructure of coatings identified that the coating formed in 125 (s) had a more compact and steady structure with fine surface cavities and less porosity. The X-ray diffraction pattern of coating demonstrates that this coating consists of oxide phases of titanium (rutile and anatase) and hydroxyapatite. Moreover, the study of corrosion resistance of coatings by potentiodynamic polarization and electrochemical impedance spectroscopy in corrosive solution of chloride sodium 3.5% showed that the coating formed in 125 s had the most significant corrosion resistance potential and the least \(I_{corr}\) and, thus, the highest resistance to corrosion.

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1. Introduction

Titanium and its alloys are highly applicable to the aerospace, chemical, biomedicine, energy, and marine industries [1,2] due to the perfect properties such as high rigidity relative to weight, almost low density, good corrosion resistance, high melting point, good bio-compatibility, and mechanical behavior [3,4]. In addition, the formation of an oxide and stable layer on titanium and its alloys while exposing air or moisture makes them able to be protected against oxidation and corrosion resistance. Furthermore, the thin thickness of this layer (5 to 70 angstrom) may lead to the rapid destruction of some environments such as sulfuric, hydrochloric, and phosphoric acid [5,6]. On the other hand, tribological applications of titanium and its alloys are limited due to the low resistance to erosion, low rigidity, and high friction coefficient [7]. There are several approaches to coating titanium and alloys including physical and chemical deposition of vapor, penetration processes, plasma spray, and laser and nitrating operation. The above-mentioned approaches may improve the properties through the formation of rigid coatings [8]. Most of the mentioned methods are expensive and time-consuming; there is a weak possibility for desirable adhesion [9]. Therefore, the plasma electrolyte oxidation process is derived from the
anodizing method and may cause ceramic oxide coatings that have been created in recent years. In PEO, mechanical properties, corrosion resistance, and erosion on light metals, such as titanium and aluminum, have improved with 10-100 micron coatings [10,11]. This coating approach is known as an electrical discharge process (unstable, fine, and short-life sparks) on the anode surface due to breakdown voltage with gaseous evolution [12,13]. PEO process is often carried out in weak and environment-compatible basic aqueous solution consisting of silicate, phosphate, and aluminate elements [14]. The PEO process consists of substrate oxides and complex oxides [15]. There are different factors such as process factors (voltage, frequency, work cycle, and oxidation time) and characteristics of substrate and coatings (chemical composition, concentra-
tion, and temperature), affecting the properties of PEO coatings [16]. In this paper, the effect of coating process time on microstructure of surface and corrosion resistance of coatings formed on Ti6Al4V layer in hydroxyapatite nano-particles was investigated under constant voltage and in three different time periods.

2. Methodology

Ti6Al4V samples are selected with 20 millimeters in diameter and 5 millimeters in thickness as substrate in the coating process.

Before any coating process, the surface and edge of the sample are polished by number 60 to 1500 grinders, washed with distilled water, and dried with hot air flow. A coating bath of hydrate acetate calcium and di-hydrogen phosphate hydrate with concentrations of 0.04 and 0.01 M is used for the process, respectively. The processes are carried out using a 15 KW self-made coating device with a direct feeding resource and in constant conditions of 600 V and in three periods of 125, 250, and 350 seconds with a constant current density of 0.1 A/cm². PH balance and electrolyte conductivity were carried out by adding 10 g/l hydroxyapatite nano-powder to electrolyte. In this process, the titanium samples are used as anode, and a coat of stainless steel was used as cathode. The temperature was fixed at 30-35°C. After coating, the coated sample was taken out of electrolyte bath, washed with water, and dried by air. Table 1 demonstrates the properties of samples and process conditions.

In the primary stages of the PEO process, the current increased rapidly as gradually applied voltage (to 600 volts) increased, and severe oxidation and gaseous evolution occurred in coating baths. The surface microstructure of coatings was investigated by Scanning Electron Microscope (SEM). The size of cavities and the thickness of coatings were determined by software. Further, MIP software was used in order to determine the porosity percentage of coatings. The phase composition of coating was studied via the analysis of X-ray pattern by KO Cu ray in a vector of 20 to 100 degrees. The corrosion resistance of titanium samples without coats and with different coats was investigated using electrochemical impedance spectroscopy and potentiodynamic polarization in 3.5% sodium chloride corrosive solution at room temperature. These tests were carried out on a three-electrode flat cell. In this cell, the tested sample with a floating level of 0.4-centimeter square is the electrode, platinum sticks as a supplementary electrode, and silver wire as a referenced electrode in the silver chloride-saturated solution (saturation in 3-molar potassium chloride solution). NOVA 11.1 software was used for extracting data resulting from curves and modeling the electrochemical impedance spectroscopy curves. Prior to conducting corrosion resistance tests, the samples floated in corrosive solution for 1800 seconds in order to achieve stability and measure the open-circuit potential. The electrochemical impedance spectroscopy test was carried out at frequency rates of 0.01 to 100000 Hertz and ±10 millivolts. By applying −250 millivolts potential toward open-circuit potential to 500 millivolts potential, the potentiodynamic polarization test was carried out at a rate of 1 millivolt per second on the samples. The tests tripled.

3. Discussion and Results

3.1. Coating Development

When applied voltage reached breakdown voltage, dielectric breakdown occurred in weak points such as cracks or impure ions on the surface [17]. After reaching the breakdown voltage of oxide layer, the electric discharge phenomena occurred on the surface of titanium sample and the current decreased smoothly. In the step coating, a non-finite number of white, fine, and short-life sparks are created on the anode surface. In fact, when oxygen is absorbed in the form of oxygen compositions, the electric discharge is known as a combustion phenomenon and an exothermic reaction including oxide factor [18]. The spark can alter temperature and local pressure of evacuation channels from 1000 to 10000 degrees (Kelvin) and from 100 to 1000 (mega Pascal), respectively, and may lead to the eruption of the lava from evacuation channels and

| Sample code | Time (s) | Frequency (HZ) | Electrolyte composition |
|-------------|---------|----------------|-------------------------|
| A           | 125     | 1000           | 10 g/l nano-HA          |
| B           | 250     | 1000           | 10 g/l nano-HA          |
| C           | 350     | 1000           | 10 g/l nano-HA          |
the formation of volcanic-crater-like cavities on the surface [19,20]. Over time, intensity and size of sparks increase as the number decreases. Their color gradually changes to yellow and, then, to orange. Since the coating bath and other conditions except time were the same in all of the three processes, the characteristics of sparks such as light and sound, number, size, intensity, and uniformity were similar until 125 s. By increasing the time of the process to 250 and 350 s, their size and intensity increase. The characteristics of sparks influenced the microstructure of surface, size, and uniformity of micro cavities and corrosion resistance properties of the formed coatings in different periods.

### 3.2. Microstructure of surface

In order to characterize the coating created on substrate, the pictures of the scanning electron microscope were used. Figure 1 shows the picture of the scanning electron microscope taken from Ti6Al4V substrate before the coating process. This picture shows a uniform surface with lines and cavities resulting from the surface polishing process. Figure 2 reveals the surface microstructures of the formed coatings in different time periods. As can be observed, the surface of all coated samples at different times is non-uniform and porous in comparison with uncoated ones due to the formation of micro sparks in the substrate during the process. The average size of surface micro cavities and computed porosity percent are given in Table 2 by different coating analysis software products. An increase in the time of processing may lead to greater intensive and larger evacuation, resulting in a larger structure of the coating surface due to the formation of relatively large cavities [21]. Therefore, the data presented in Table 2 showed that time increment from 125 to 350 s might lead to the growth of surface structure and the size of micro cavities. Nevertheless, according to the size development and the decreasing number of micro-cavities because of a pressure drop in evacuation channels during processing [22], the surface porosity percentage increased and, then, decreased over time. It is obvious that the surface microstructure of coatings in 125 s has the least surface micro cavities, the least porosity percent, and the highest density. In fact, the size of cavities based on processing time increment increased due to the application of less electrical current and dielectric stability [22]. Moreover, by increasing the time of processing from 125 to 350 s, the thickness of coatings increased, too (Table 2).

### 3.3. Phase composition of coating

Two groups of coating baths were used in PEO processes. The first group may cause metal oxides with oxygen in coating, and the second group includes anion elements that insert other elements (except oxygen) in coatings [9]. Figure 3 shows the Energy Dispersive X-ray Spectroscopy (EDS) analysis of the coated sample in different times. The results demonstrated that the coating consisted of Al as anion of coating and Ti as substrate of titanium. The X-ray pattern of the coated sample is given in Figure 4 for studying the phase composition and EDS analysis. Titanium oxide (TiO$_2$) used in many industries, such as cell phones,
vehicles, and medical implements, can be formed on
titanium and alloys through the PEO process, rutile,
anatase, and brookite are the three forms of titanium
oxide. Rutile has high sustainability and enjoys good
mechanical properties due to high rigidity and better
protection properties in comparison with anatase. In
XRD pattern of Figure 4, rutile and anatase titanium
oxide phases and a few hydroxyapatite are observed.
In addition, X-ray can easily penetrate titanium sub-
strate, and titanium-related peaks appear in XRD
pattern because of porosity and thinness of coatings.
Hydroxyapatite phase is visible in all three coatings,
and the amount of this phase in coating A is the
most. Rutile phase available in coating A having more
amount than others and the simulation of this phase
with hydroxyapatite phase can be a good combination
for applying this kind of coating in medical care such
as implements.

3.4. Corrosion resistance
Electrochemical impedance spectroscopy and poten-
tiodynamic polarization tests were carried out in order
to evaluate the resistance to corrosion in coated samples
in different periods and their comparison.

3.4.1. Electrochemical impedance spectroscopy test
Nyquist curves obtained from electrochemical
impedance spectroscopy tests of coated and uncoated
titanium samples are given in Figure 5. In these
curves, the imaginary part of impedance is drawn
based on real part in each frequency after 1800 s
floating in corrosive solution of 3.5% sodium chloride.
As can be observed, the Nyquist curve of uncoated
titanium substrate has a half-ring capacitor in every
frequency. This behavior is because of forming an
electrical layer on the sample surface and a time
constant in the circuit, as presented in Figure 6(a).
The corrosion resistance was calculated by measuring
the diameter of the half ring in substrate Nyquist
curve. In contrast, the Nyquist curves of coated

Figure 3. Energy Dispersive X-ray Spectroscopy (EDS) analysis of the coated samples in (a) 125, (b) 250, and (c) 300 s.

Figure 4. XRD pattern for the coated samples.

Figure 5. Electrochemical impedance spectroscopy tests of coated and uncoated titanium samples in corrosive solution of 3.5% sodium chloride.
samples have 2 half-ring capacitors. The larger ring appearing at low frequencies is related to the internal dense layer, and the smaller ring at high frequencies is related to the external porous layer [23]. The internal dense layer forms due to the voltage applied to metal, and the porous layer forms due to the plasma reaction with metal and coating bath during processing [1]. The Nyquist curve of coating formed in 350 s had a smaller diameter in comparison with two other coatings and even uncoated titanium substrate. This shows weaker corrosion resistance of this coating, compared with other samples. As explained in Figure 4, the coating C has less rutile and hydroxyapatite than others, and it can be due to the relative destruction of the upper layer on titanium during the PEO process [24]. In contrast, the diameter of rings in Nyquist curves was related to the coatings formed in 125 and 250 s from the diameter of rings related to the bigger titanium substrate because of rutile and hydroxyapatite phases available. This shows better corrosion resistance of these two coatings in comparison with that of uncoated substrate. The extracted corrosion resistance of internal layers from equal circuit is reported in Table 3. Since the resistance of internal layer is more than that of the external layer, the corrosion resistance of coatings depending on internal and external layers plays no significant role. Nyquist curves of the coatings formed in 180 s have the largest ring diameter and, therefore, the biggest corrosion resistance among all samples. In this sample, the corrosion resistance is about 37 times more than the uncoated substrate resistance.

3.4.2. Potentiodynamic polarization test
Potentiodynamic polarization test was carried out in order to verify the corrosion resistance behavior of the coated and uncoated samples. Potentiodynamic polarization curves for coated and uncoated titanium substrates in corrosive solution of 3.5% sodium chloride can be seen in Figure 7. As can be observed, by applying a coating to the sample in 125 and 250 s, the curves are delivered to upside to the left and in 350 s to the right and up toward the curve of the uncoated sample. The sample coated in 125 s had the most transportation and the best behavior. This curve has the most positive corrosive potential and the least density of corrosive current. By increasing the coating time, the corrosion resistance potential becomes more negative, the current density increases, and the curves move on to the right side. While corrosion resistance potential shows the thermodynamic trend to corrosion resistance, the density of corrosive current shows the corrosion resistance rate of samples [25]. Since the density of corrosion resistance current decreases [25], corrosion resistance increases and the coated sample in 350 s has the highest density of corrosive current and the least corrosion resistance in comparison with the two other coated and uncoated substrates. However, corrosive potential related to this sample was more significant than the uncoated substrate. In fact, the more significant corrosive potential demonstrates higher chemical sustainability of the titanium substrate [1]. Since the only altered factor in this paper is the processing time, the surface microstructure is

![Figure 6.](image_url)  
**Figure 6.** Equivalent circuit for modeling of data from Electrochemical Impedance Spectroscopy (EIS): (a) Uncoated titanium and (b) coated titanium.

| Sample code | \( n \) | \( Y_0 \) | \( R_1 \) |
|-------------|--------|--------|--------|
| Uncoated (bare) | 0.838 | 51.5 | \( 5.87 \times 10^4 \) |
| A | 0.845 | 68 | \( 2.16 \times 10^6 \) |
| B | 0.829 | 98 | \( 1.1 \times 10^6 \) |
| C | 0.840 | 180 | \( 2.06 \times 10^4 \) |

![Figure 7.](image_url)  
**Figure 7.** Potentiodynamic polarization curves for coated and uncoated titanium substrates in corrosive solution of 3.5% sodium chloride.
the most important parameter effective in corrosion resistance of coatings. In fact, the micro cavities on the surface of coatings acted as paths for corrosive ions for reaching substrate and affecting corrosive resistance \cite{26}. Therefore, by increasing the time of the coating process from 125 to 350 s despite increasing the thickness of coating for increasing the size of cavities and porosities, the corrosive resistance of the coated sample decreased. Therefore, to effectively prevent coating corrosion, resistance should be free from structural deformations such as porosity and thickness. Corrosion potential and density of corrosion current were derived from potentiodynamic polarization curves, as given in Table 4. As can be observed, the coated sample in 125 s had the most significant corrosion potential, the least density of corrosive current, and the best corrosion resistance, compared to other samples. In fact, the density of corrosion resistance current in this sample is about 2.5 times less than that in the uncoated substrate.

4. Conclusion

This study investigated the effect of processing time on surface microstructure and corrosion resistance of coatings from the plasma electrolytic oxidation process on pure titanium substrate. The results are as follows:

1. Increasing the processing time of coating from 125 to 350 s may lead to an increase in the size of surface micro cavities, a decrease in the number of cavities, and magnification of the structure of coatings due to the formation of larger and more intensive sparks and also thickness. Therefore, the microstructure of coatings formed in 125 s had the least size of surface micro cavities, the least porosity percentage, and the highest density and uniformity;

2. X-ray pattern of coatings formed in 125 s demonstrated that this coating consisted of titanium oxide phases and hydroxyapatite;

3. Studying corrosion resistance behavior showed that more amounts of rutile and hydroxyapatite available in coating led to an increase in the corrosion resistance. However, the porosity percent, surface micro sparks, density, and coating uniformity have to be simultaneous with these phases to improve corrosion resistance considerably;

4. Increasing the processing time led to decreasing the corrosion resistance of coatings. Then, the coating formed in 125 s showed the best corrosion resistance. Corrosion resistance in this sample was about 37 times greater than the uncoated substrate resistance;

5. The corrosion resistance potential of coatings decreased and the density of corrosion resistance current increased as the processing time increased. Therefore, the coatings formed in 125 s had the most significant corrosion resistance potential and the least density of corrosion current and, as a result, the best corrosion resistance. The density of corrosion resistance current in this sample decreased about 2.5 times in comparison with the uncoated substrate.

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