A Comparative Study of 316L Stainless Steel and a Titanium Alloy in an Aggressive Biological Medium

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Abstract—The electrochemical behavior of stainless steel and titanium alloys is affected after prolonged contact with basic or acidic solutions, indicating a change in their surface properties. The human body often rejects invasive devices that aim to alter the biological or chemical composition of blood or other body fluids. Stents, fixation plates and screws, spinal implant devices, aneurysm clips, intramedullary nails and stems, temporary fixation devices and surgical instruments, etc. have been made from stainless steel AISI 316L for several years. Although the mechanical performance of implants and devices may be governed by their bulk properties, their interaction with the environment is managed by the characteristics of their superficial layer. In the case of biomedical devices, resistance to corrosion and biocompatibility has paramount importance. This study compares the corrosion behavior of 316L stainless steel and a titanium alloy in a Hank solution. The obtained results show that the titanium alloy has a higher potential than 316L stainless steel and lower corrosion current.

Keywords—corrosion; 316L stainless steel; titanium; passivation; biocompatibility

I. INTRODUCTION

Implants are organs and devices in order to replace a missing or a damaged organ of the human body, ensuring its vital functions optimally. Biomaterials, one of the great therapeutic advances in implantation, work under biological stress and play an increasingly crucial role in our health and quality of life [1]. In order to protect people against the possible dangers and inconveniences of implants and devices, it is necessary to test sufficiently their elements without putting in danger the patient’s life. This is why large-scale studies and clinical trials are required [2]. Experiments carried out over the years have shown that cells’ response to the surface of the biomaterial varies according to their type and state of maturation. In vitro and in vivo tests have led several authors to conclude that the attachment of osteoblasts is reinforced by the roughness of the surface [3, 4]. The first tests, regarding the resistance to corrosion in physiological environment, in vitro and in vivo, seem conclusive and should continue in the coming years [5]. Research was carried out for orthopedic applications, such as knee and hip prostheses [6, 7]. This type of prosthesis undergoes significant mechanical frictional stresses, causing the release of metal particles in large quantities.

Metallic biomaterials have been used in various biomedical applications for a century, due to their low cost, excellent mechanical properties and inertness [8, 9]. Stainless steel and titanium alloys are frequently employed in many biomedical applications, including cardiovascular stents/valves, orthopedic prosthesis and other devices and implants used in biomedicine, because of their biocompatibility and mechanical properties, such as high strength and toughness [10, 11]. Titanium alloys are extensively used due to their attractive combination of high strength, low density, and good corrosion resistance [12,13]. Moreover, the 316L stainless steel is known for its good ductility, fatigue resistance, and high strength toughness [14, 15]. However, there is always a concern about their corrosion resistance in physiological medium and their biocompatibility [16, 17]. The effects of surface treatment and metallic coating on the corrosion behavior and biocompatibility of surgical 316L stainless steel implants were evaluated in [18]. The experimental results indicated that Nb coating and surface treatment of the stainless steel improved its corrosion behavior. The structural and bio-corrosion barrier performance of Mg-substituted fluorapatite coating on a 316L stainless steel human body implant was investigated in [19], concluding in improved corrosion resistance and biocompatibility. In [20] the microstructural evolution, mechanical, sliding wear and corrosion behavior of 316L stainless steel multi axially forged at 600°C was studied, showing that severely deformed ultra-fine-grained metals have the potential to deliver improved implant performance. The electrochemical tests conducted in [21] showed that the surface enriched by molybdenum improved the corrosion resistance of the 316L stainless steel.

Recently, a comparative assessment of the corrosion intensity of orthodontic arch wires made of alloy steel, nickel-titanium and titanium-molybdenum alloys, in laboratory conditions, was presented in [22]. The highest resistance to corrosion was observed in nickel-titanium alloy arch wires, while steel wires had the lowest. Surface modification allows improvements on the materials’ characteristics without compromising their important bulk properties. A review on stainless steel surface modification methods provides guidance...
The purpose of this study is to compare the corrosion behavior of 316L stainless steel and titanium alloy in a Hank's solution, utilizing electrochemical measurements, such as potentiodynamic and cyclic polarization.

II. STUDIED MATERIALS

AISI 316L stainless steel and titanium alloys are commonly used in the manufacturing of prostheses. The chemical composition of these materials was determined by spectroscopic analysis (Tables I and II). Samples were cut in the form of a 14mm in diameter disc from cylindrical bars delivered in a raw state.

III. SURFACE PROPERTIES AND POLISHING TECHNIQUES

Optimizing the properties of surface corrosion resistance is a wide and important research topic, given its potential application in different biomedical fields. Corrosion resistance is one of surfaces' major characteristics that must be optimized in order to improve the integration of implants. In this section, focus is set on roughness and electrochemical properties. Anodic dissolution of the sample can produce excellent polishing results without deforming its surface. The sample is polarized at the anode and connected to the cathode through an electrolyte bath concentrated in acid. Applying voltage creates a current that travels through the bath and thus preferentially electrolyte bath concentrated in acid. Applying voltage creates a current that travels through the bath and thus preferentially

distributes the concentration of the electrochemical system around the working electrode. In this section, focus is set on roughness and electrochemical properties. Anodic dissolution of the sample can produce excellent polishing results without deforming its surface. The sample is polarized at the anode and connected to the cathode through an electrolyte bath concentrated in acid. Applying voltage creates a current that travels through the bath and thus preferentially dissolves the salts without attacking the metal [25].

B. Electrochemical Analysis

1) Used Solution

The Hank solution was used as an aggressive environment, assimilated to the physiological medium prepared from the compounds given in Table V.

TABLE V. CHEMICAL COMPOSITION OF THE HANK SOLUTION

<table>
<thead>
<tr>
<th>Compositions</th>
<th>Quantity (g/l)</th>
</tr>
</thead>
<tbody>
<tr>
<td>NaCl</td>
<td>8</td>
</tr>
<tr>
<td>CaCl₂</td>
<td>0.15</td>
</tr>
<tr>
<td>KCl</td>
<td>0.40</td>
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<tr>
<td>NaHCO₃</td>
<td>0.35</td>
</tr>
<tr>
<td>Glucose</td>
<td>0.1</td>
</tr>
<tr>
<td>MgCl₂·6H₂O</td>
<td>0.1</td>
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<tr>
<td>Na₂HPO₄·2H₂O</td>
<td>0.6</td>
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2) Experimental Conditions

The samples were characterized electrochemically under the following conditions:

- A simulated physiological environment (Hank's solution of PH = 7.1)
- Environment temperature: 37°C (human body temperature)
- Sweep rate of the potential: 0.5 mV/s

3) Evolution of the Free Potential

This technique makes possible to follow the evolution over time of the sample potential (working electrode) in an open circuit (OCP) relative to a reference electrode. The sample was kept immersed in the electrolyte and the free electrode potential was measured as a function of time. The open-circuit potential (Ea) is determined from the potential follow-up curves as a function of time. This potential is a characteristic of the modification of the interface between a metal and its surrounding environment.

IV. RESULTS AND DISCUSSION

A. Microhardness

The microhardness tests were carried out under a load of 20g, load from which the impression is visible under the microscope, using a Vickers microdurometer equipped with a microscope. Vickers HV hardness is related to the length of the diagonal of the impression (d) and the applied load (m) by:

\[ H_V = 1.854m / d^2 \]

with \( m \) in [kgf] and \( d \) in [mm]. The microhardness results for both materials are shown in Table IV and represent the average of 10 measurements. It can be noted that the titanium alloy is mechanically stronger than 316L stainless steel.

TABLE IV. MICROHARDNESS OF THE TESTED SURFACES OF THE STUDIED MATERIALS

<table>
<thead>
<tr>
<th>Materials</th>
<th>Microhardness</th>
</tr>
</thead>
<tbody>
<tr>
<td>316L stainless steel</td>
<td>166.8 ±0.2Hv</td>
</tr>
<tr>
<td>Titanium alloy</td>
<td>354.0 ±0.2Hv</td>
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for selecting the appropriate modification routes, tailored for specific biomedical applications [23].
The variation of the potential value with immersion time can be explained by the electrochemical reaction which begins to take place between the medium (Hank’s solution) and the exposed surface of the metal substrate, more or less adherent, causing a relative stability of the potential for corrosion abandonment.

TABLE VI. FREE POTENTIAL RESULTS FOR STUDIED MATERIALS

<table>
<thead>
<tr>
<th>Material</th>
<th>E\text{corr} (mV/ECS)</th>
</tr>
</thead>
<tbody>
<tr>
<td>316L stainless</td>
<td>-63</td>
</tr>
<tr>
<td>Titanium alloy</td>
<td>-10</td>
</tr>
</tbody>
</table>

4) Potentiodynamic Curves log(I)=f(E)

A metal immersed in any electrolytic medium tends to dissolve and electrically charge creating a double electrochemical layer, comparable to an electric battery. After a long enough time, a steady state is established and the metal electrode takes, relative to the solution, a potential called the potential corrosion (E\text{corr}). The latter cannot be determined absolutely and it is marked with respect to a reference electrode. Using an external generator and a counter-electrode, a current passes through the metal electrode, its steady state is modified and its surface takes a new potential value (the curves E=f(I) or I=f(E) constitute the polarization curves). Figures 3 and 4 show the evolution of the polarization curve log(I) in terms of the potential E for both materials. The potential scanning range was set at a speed of 0.5mV/s. Calculation of the density \(i_{\text{corr}}\) and corrosion potential E\text{corr} is performed by extrapolating the anode and cathode tafel lines to the corrosion potential E\text{corr}. The point of intersection gives us directly \(i_{\text{corr}}\) and E\text{corr}. The corrosion rate V\text{corr}, expressed in (mm/year), is calculated by [27]:

\[
V_{\text{corr}} = \frac{10i_{\text{corr}}.d.M}{n.E.d} \tag{2}
\]

where \(i_{\text{corr}}\) is the corrosion current density (A/cm\(^2\)), \(t\) is the time corresponding to years, \(M\) is the average atomic mass of the studied material, \(n\) is the valence, and \(d\) is the density of materials.

In Figure 3, it is observed that the titanium alloy in the solution becomes passive and has a passivation plateau of the order of 500mV. The corrosion current is very low and is of the order of 0.01µA/cm\(^2\) (Table VII). So, the titanium alloy has a low rate of degradation (60×10\(^{-3}\)mm/year). In fact, the corrosion current can be likened to a residual current resulting from an ion diffusion phenomenon through the passive layer that has been formed. A break in the passivation layer was noticed beyond +500mV/ECS. It should be noted that this break was partial because the measured currents remained low. Beyond +1.0V/ECS, where the overvoltage is greater, the currents increase drastically, meaning that the layer is completely broken. This state gives a good behavior against corrosion. The 316L steel exhibits the same behavior as the titanium alloy. The polarization curve (Figure 4) shows a passivation level of +500mV/ECS and a film breaking potential of +480mV/ECS. The corrosion current remains low, which is of the order of 2µA/cm\(^2\) and the resistance to polarization is weak (10.7MΩ) (Table VIII).

According to the results obtained in the Hank solution for both alloys, significant differences were noted in their electrochemical characteristics values. The corrosion potential of the titanium alloy is close to -190.8mV/ECS and its corrosion current density was 9.8×10\(^{-3}\)µA/cm\(^2\). Compared to the 316L stainless steel values, titanium’s alloy values are...
clearly favorable. This alloy has greater potential than 316L stainless steel and lower corrosion current. The corrosion rate for the titanium alloy is $60 \times 10^{-3}$ mm/year. This value is very acceptable in the corrosion field, showing better corrosion behavior in the Hank solution (similar to physiological environment) compared to 316L stainless steel. Titanium-based alloys and stainless steels are passive materials sensitive to pitting corrosion in the presence of specific aggressive materials. These are most often chloride ions (Cl\(^-\)), which are the most aggressive and play an important role in the pitting mechanism.

### Table VII. Potentiodynamic Results for Titanium Alloy

<table>
<thead>
<tr>
<th>Corrosion parameter</th>
<th>$E_{corr}$ (mV/ECS)</th>
<th>$I_{corr}$ ($\mu$A/cm(^2))</th>
<th>$V_{corr}$ (mm/yr)</th>
<th>$R_p$ (M(\Omega))</th>
</tr>
</thead>
<tbody>
<tr>
<td>Titanium alloy</td>
<td>-190.8</td>
<td>0.0098</td>
<td>$60 \times 10^{-3}$</td>
<td>2.21</td>
</tr>
</tbody>
</table>

### Table VIII. Potentiodynamic Results for 316L Steel

<table>
<thead>
<tr>
<th>Corrosion parameter</th>
<th>$E_{corr}$ (mV/ECS)</th>
<th>$I_{corr}$ ($\mu$A/cm(^2))</th>
<th>$V_{corr}$ (mm/yr)</th>
<th>$R_p$ (M(\Omega))</th>
</tr>
</thead>
<tbody>
<tr>
<td>316L stainless steel</td>
<td>-267.5</td>
<td>2.155</td>
<td>$16.8 \times 10^{-3}$</td>
<td>10.7</td>
</tr>
</tbody>
</table>

Pitting corrosion is related to localized attacks of heterogeneities that contribute to the deterioration of the passive layer protecting the material. This process of material deterioration generally leads to a generalized corrosion phenomenon. Observation by an optical microscope of the samples for both studied alloys shows the attacking shape they experienced after a corrosion test in Hank's solution (Figure 5).

### 5) Cyclic Potentiodynamic Curves

The behavior of the oxide layer formed on the samples’ surface was studied by drawing the cyclic potentiodynamic curves. Figure 6 shows the titanium’s alloy cyclic polarization curves over a potential range for a half-hour immersion time in Hank's solution with a sweep rate of 0.5mV/s. There is a passivation stage with a slight increase in the corrosion potential compared to the previous value of EC for the “go” scan. On the other hand, during the “return” scan the corrosion potential is less great with higher current densities. This means that the formed oxide layer has instabilities. This degradation may be at the source of localized corrosion.

![Cyclic potentiodynamic polarization curve of titanium alloy](image)

Cyclic curves in the case of 316L (Figure 7) indicate a different behavior. Beyond the formation of the passivation layer on this alloy, an ennoblement of the potential corrosion in...
the anodic direction of the order of 300mV and very low corrosion currents were noted. Stable protective character can be established here. Tables IX and X summarize the cyclic potentiodynamic results for titanium alloy and 316L stainless steel. It is noted that the polarization resistance of the titanium alloy is higher than that of 316L.

| TABLE IX. CYCLIC POTENTIODYNAMIC RESULTS FOR THE TITANIUM ALLOY |
|---------------------|------------------|-----------------|-----------------|
| Corrosion parameter for the Titanium alloy | $E_{cor}$ (mV/ECS) | $I_{cor}$ ($\mu A/cm^2$) | $V_{ave}$ (mm/year) | $R_p$ (KΩ) |
|---------------------|------------------|-----------------|-----------------|
| 316L stainless steel | -419.0 | 0.347 | 53.82×10^{-3} | 62.51 |

- In the Hank’s solution significant differences were found in the electrochemical characteristics. The titanium alloy had higher potential than 316L stainless steel and lower corrosion current
- The use of cyclic intensity-potential curves, can help determining the quality of the passive film
- Finally, the study of the polarization resistance evolution as a function of the immersion time shows that the two alloys retain their resistance properties as a function of the immersion time in the Hank’s solution.

ACKNOWLEDGMENT

The authors thank the laboratory of the ALFON complex (Algerian Foundry Society of Oran) for their support and assistance in the determination of chemical compositions by spectroscopic analysis.

REFERENCES

Aroussi et al. : Comparative Study of Two Passive Metals in Aggressive Biological Medium


