



Electrochemical characterization of Metal injection moulded Ti6Al7Nb used for dental applications under varying saliva conditions

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Abstract

In avoidance of metal ions into the human bloodstream, Ti₆Al₇Nb became a subsequent alloy for various implant's purpose. In this study, different corrosion techniques have been utilized to characterize corrosion behavior of Ti₆Al₇Nb alloy in pure saliva simulating electrolyte (SSE), varying fluoride conditions, and different peroxide concentrations as well as under severe plaque conditions when Ti₆Al₇Nb alloy is used for dental applications. With an exhibition of passivity range within most engaged conditions, Potentiodynamic polarization (PDP) curves generally revealed Ti₆Al₇Nb as good material for dental applications. Nonetheless, the increment of fluoride, peroxide (H₂O₂) concentration and severe plaque conditions showed a negative impact on Ti₆Al₇Nb alloy. This observation was clearly observed from electrochemical impedance spectroscopy (EIS) curves as the most compressed semicircles were found under those conditions. A critical fluoride concentration on the surface of the Ti₆Al₇Nb alloy was found to be 5g/l. In attesting to the later, surface morphologies under pure SSE was noble, while it was rapidly degraded under severe plaque condition.

Keywords: Corrosion, Ti6Al7Nb, PDP, SSE, EIS

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Citation: Onoyivwe Monday Ama et al. (2018), Electrochemical characterization of Metal injection moulded Ti₆Al₇Nb used for dental applications under varying saliva conditions. *Int J Nano Med & Eng.* 3:7, 71 - 77

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Received: October 05, 2018

Accepted: October 15, 2018

Published: November 29, 2018

Introduction

Due to low specific weight, good mechanical properties, low young modulus, very good strain-controlled and notch fatigues resistance, high strength-to-weight-ratio, superior corrosion resistance, etc., titanium (Ti) and its alloys lead on the utilization of biomedical and dental implants [1-4]. The Versatility of Ti and its alloys is from three categories it's normally found from: α, (α+β) and β phases [5, 6]. Within these categories, different problems exist.

Ti₆Al₄V (α+β phases) has a complexity of toxicity elements which release into the body fluid during their applications as implants [7-9].

In particular, aluminium (Al) and vanadium (V) release lead to inflammation and cytotoxicity in the surrounding tissue of implants. In fact, an accumulation of metallic ions in body fluid causes hypersensitivity and thrombogenesis, and these consequent to implant loosening. The worst consequence of ions release into the body entails difficulties such as revision and/or removal of implants [9-12]. In avoidance of toxicity elements release into the blood fluid, Ti₆Al₇Nb has subsequently been developed [13],[14],[15],[16].

Corrosion measurements for pertaining to wimplant usage has previously being done by numerous authors, and different findings have been documented [17],[18],[19],[20]. Furthermore, in consideration of localized corrosion occurrence which goes with non-uniformity pH around implants [21], varying electrolyte conditions have been found to play a critical role on the lifespan of the implant. Fojt et al. [11] utilized corrosion measurements on the characterization of porous Ti39Nb using phosphate-buffered saline solution as an electrolyte. It was found that porosity above certain limit could cause localized corrosion. With an utilization of Ringer solutions at different pH corresponding to severe conditions human body experiences due to surgery, infections and inflammations, Calderon Moreno et al. [22] provided in roads through utilization of electrochemical measurements. Most of all, fluoride which is normally used in toothpaste were found to be problematic at certain concentration [23-25].

Metal injection moulding (MIM) as the state-of-art powder metallurgy manufacturing technique which produces alloys on a mass production of complex small-medium parts has got quite a number of advantages over conventional manufacturing techniques which can be found elsewhere [26] [19, 27]. On the research of the components fabricated by MIM, most authors studied and reported on mechanical properties,

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chemical and biocompatibility properties of newly improved materials [26, 28-36]. However, there are no corrosion measurements, particularly, on deeper characterization of passive layer which forms on Ti_6Al_7Nb manufactured by MIM when it's exposed dynamic conditions for dental usage. The present study engaged for corrosion measurements to focus on the empirical occurrence of localised corrosion on Ti_6Al_7Nb manufactured by MIM processes when it is used for dental applications.

Experimental

Materials and Solution: Ti_6Al_7Nb specimens manufactured by MIM were provided in dimensions of $2cm^2$. The Standard electrolyte used in this work was Fusayama-Meyer artificial saliva, and the composition was synthesized from the following chemical powders: NaCl (0.4 g/L), KCl (0.4 g/L) $CaCl_2 \cdot 2H_2O$ (0.796 g/L), $NaH_2PO_4 \cdot 2H_2O$ (0.690 g/L), $Na_2S \cdot 9H_2O$ (0.005 g/L) and Urea (1 g/L). The saliva simulating electrolyte (SSE) was prepared using deionised water with a resistivity of 18.2 MV cm. Certified reagents supplied by Alfa Aesar, USA. NaF (Sigma Aldrich, USA) was added to the artificial saliva solution to simulate different fluoride media, and it was varied up until a critical concentration was obtained. As based from the literature, for the simulation of the plaque conditions, sulphuric acid (H_2SO_4) was used to lower the pH of the pure SSE to 3.0, while the peroxide solutions in SSE (0.1% and 10%) were obtained following steps that can be found elsewhere [37, 38].

Electrochemical Measurements: An electrochemical cell with three imperative openings was used in this study to simulate localized corrosion occurrence on bare Ti_6Al_7Nb in the dynamic SSE. Three openings were for engaged electrodes: a working electrode (Ti_6Al_7Nb), a platinum wire (counter electrode) and 3M Ag/AgCl as a reference electrode. The whole setup was computerized from advance autolab PGSTAT302 N potentiostat.

Prior to any electrochemical measurement, the OCP was run at room temperature for at least 2 hrs. After OCP measurements,

Potentiodynamic polarization (PDP) and Electrochemical impedance spectroscopy (EIS) measurements were carried out. Firstly, the potential of 250 mV was scanned in the reverse direction (cathodically polarized) to remove oxide which formed on the surface of the LCS electrode during an attainment of steady-state conditions under OCP. Secondly, the anodic and cathodic curves were measured at the scan rate of 0.16mV/s; starting from -1 V to 1.5 V.

For EIS measurements, NOVA 2.0.1 software was utilized. The Measurements were carried out at OCP over a frequency scan ranging from 100 kHz to 100 mHz. The wave type was single sine at an integration time of 0.125s per cycle and the amplitude used was 10 mV. Ultimately, after an acquisition of EIS spectra, an electrical equivalent circuit (EEC), which was attained through Boukamp model, was utilized to interpret the EIS data. For reproducibility of the results, all the electrochemical measurements were triplicated.

Surface Characterization: Scanning electron microscopy (SEM) coupled to energy-dispersive spectroscopy (EDS) was used to inspect the surface phase morphologies and elemental compositions of the engaged samples. The used SEM was JEOL JSM electron microscope, and it was operated at 20 kV.

Results and Discussion

Corrosion Measurements results: Fig. 1 below shows the OCP curves for all engaged Ti_6Al_7Nb electrodes in the dynamic SSE used in this study. As can be seen from the Figure, the OCP for 0.1% H_2O_2 is more positive than the other OCPs. This could be due to a momentary oxidation which occurs on the surface of the electrode. In comparison with the higher content (10% H_2O_2), an exhibition of more negative CP potential can be seen in Fig. 1. Furthermore, it can also be seen that increasing of the fluoride content and existence of plaque condition led to OCP values shifting from noble values. This could be due to an aggressive effect on the passivation layer of Ti_6Al_7Nb within those electrolyte media [39, 40].

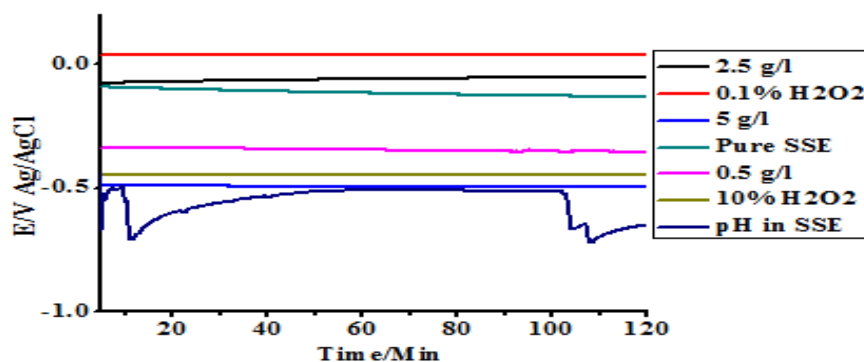


Fig.1: OCP curves for Ti_6Al_7Nb in different SSE conditions

On the other hand, Fig. 2 shows PDP curves, and it was intended to characterize the corrosion behavior of Ti_6Al_7Nb in engaged SSE conditions. As can be seen in Fig. 2, Ti_6Al_7Nb shows active-passive behavior as it starts reacting with electrolyte under all engaged conditions. As time goes on, it forms a compact passive layer which prohibits further active transition, particularly under pure SSE. However, with inclusion and increment of fluoride concentration, anodic curves deviate a bit from noble region, and the critical concentration was found to be 2.5 g/l NaF. An initial introduce of peroxide (0.1% H_2O_2)

shifted corrosion potential to the more electropositive value, and shift anodic curve to the noble side as can be seen in Fig. 2. As H_2O_2 is a strong oxidizing agent, in its initial impact it thickens the oxide layer and increases cathodic current [41]. In contrast, when 10% H_2O_2 was applied, the potential and current density increase rapidly. This could be due to defects which are created on the oxide layer by the presence of peroxide in the SSE [41]. Even for the 0.1% H_2O_2 , it is known that the observations are attributed only to a short period of time which H_2O_2 inclusion was exposed to Ti_6Al_7Nb . On the previous study

where H_2O_2 was exposed to Ti alloy, it was realised that it becomes worst as time goes on. On the other SSE modification, it was observed

that maximum rapid corrosion occurs under plaque conditions (SSE modified to pH 3.0 by H_2SO_4). That is evidenced by quazi-broken oxide layer and a higher current density as can be seen in Fig. 2.

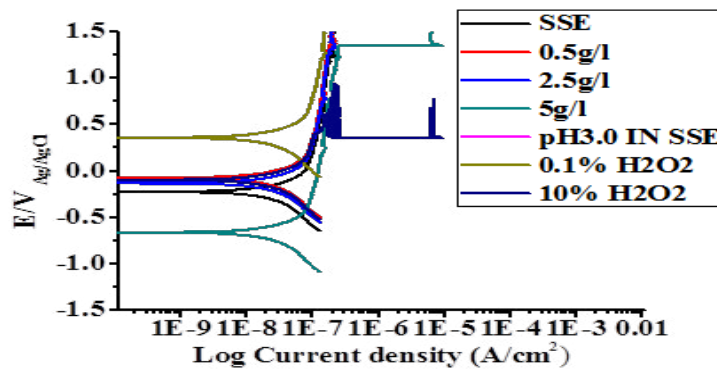


Fig. 2: PDP curves for Ti_6Al_7Nb in different SSE conditions: Pure SSE, 0.1% H_2O_2 , 0.5g/l NaF, 2.5g/l NaF, 5g/l NaF and pH 3.0 in pure SSE (plaque condition).

PDP observations were furthermore corroborated by EIS Nyquist diagrams shown in Fig. 3. On a deeper analysis of the diagrams, it can be seen that the semicircles became more compressed when fluoride concentration increases and under severe plaque condition in SSE. This is the exact view that was observed in previous studies^[17], and it is attributed to the acidified conditions which became involved therein. In Fig. 4, a higher impedance values were noted at low frequency for pure SSE and 0.1% H_2O_2 . This once again proves the negative impact of higher fluoride concentration on Ti_6Al_7Nb . Higher fluoride concentration has been reported to lower corrosion resistance of

implants alloys^[42], and the findings herein are consistent with the previous studies. Most importantly, for engaged Ti_6Al_7Nb from novel fabrication method (MIM), the fluoride content of about 5 g/l noticed to be critical. In Fig. 5, a constant phase angle curve can be seen at a low-frequency range in Fig. 5 (i), and that shows an intact passive layer which is anticipated to alloys which are used for implants^[43, 44]. For this study in particular, the intact oxide layer underneath obviously porous layer (TiO_2) is Nb_2O_5 , and it is shown in Fig. 6. However, as other conditions are introduced, the phase angles started to decline within the intermediate frequency range, and that is attributed to the dissolution of oxide layers^[39].

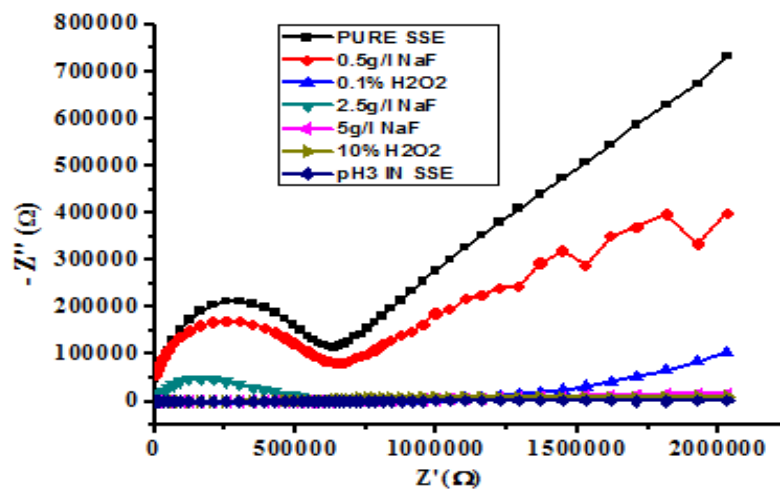


Fig. 3: EIS Nyquist diagrams for Ti_6Al_7Nb in different SSE conditions : Pure SSE, 0.1% H_2O_2 , 0.5g/l NaF, 2.5g/l NaF, 5g/l NaF and pH 3.0 in pure SSE (plaque condition).

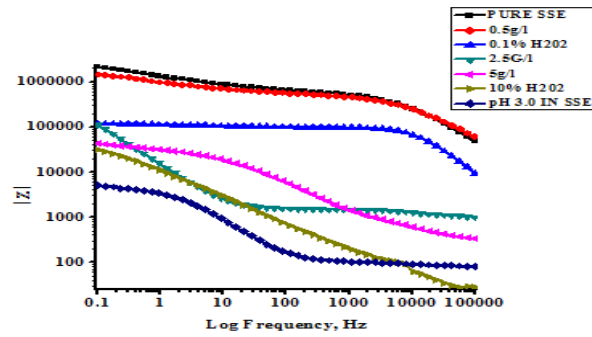


Fig. 4: EIS bode impedance plots ($|Z|$) for Ti_6Al_7Nb in different SSE conditions : Pure SSE, 0.1% H_2O_2 , 0.5g/l NaF, 2.5g/l NaF, 5g/l NaF and pH 3.0 in pure SSE (plaque condition).

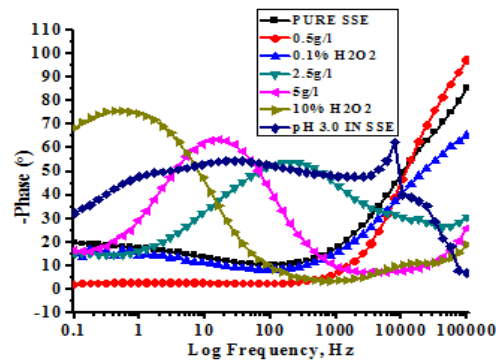


Fig. 5: EIS bode phase plots ($^{\circ}$) for Ti_6Al_7Nb in different SSE conditions: Pure SSE, 0.1% H_2O_2 , 0.5g/l NaF, 2.5g/l NaF, 5g/l NaF, and pH 3.0 in pure SSE (plaque condition).

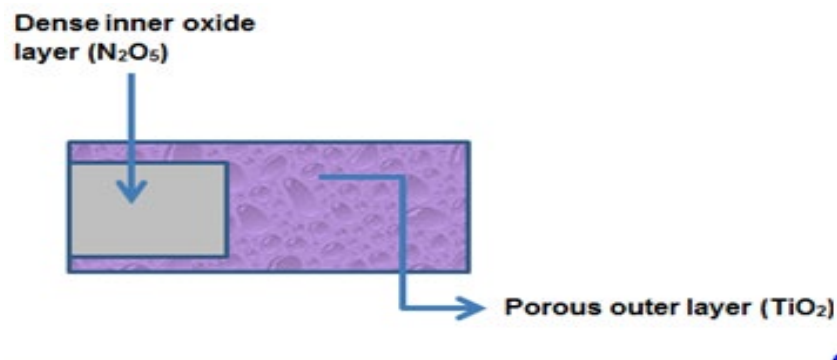


Fig. 6: Two layers forming on Ti_6Al_7Nb under exposure to electrolytes

Surface morphologies: Fig. 7(a) to (g) show surface morphologies of Ti_6Al_7Nb alloy when exposed to aforementioned electrolytes engaged in this study. Vividly, intact oxide layer can be seen when Ti_6Al_7Nb alloy exposed to pure SSE. The morphology is composed of black and white parts as can be seen in Fig. 7a. The observation revealed alloying elements (Al and Nb) to the titanium, and this shows negligible occurrence of the degradation^[45]. On 0.5 g/l NaF and 0.1% H_2O_2 usage as electrolytes, a uniform corrosion oxide products can be seen all over the surfaces (Fig 7(b) and (c)), and that shows mild corrosion reactions on the surface of Ti_6Al_7Nb from the engaged electrolytes^[46].

In opposite, when the alloy was subjected to further fluoride concentrations (2.5 and 5 g/l), aggressive surface degradation occurred on the surfaces as can be seen in Fig. 7 (d) and (e). Most notably, when the peroxide concentration was increased to 10% and under plaque conditions, aggressive surface degradation became apparent as can be seen in Fig. 7(f) and (g). All these are in agreement with previous analysis and some reported work that higher fluoride concentrations, the existence of peroxide and plaque conditions are toxic to the alloys surfaces used for implants^[47], and Ti_6Al_7Nb fabricated by MIM is no exception as well.

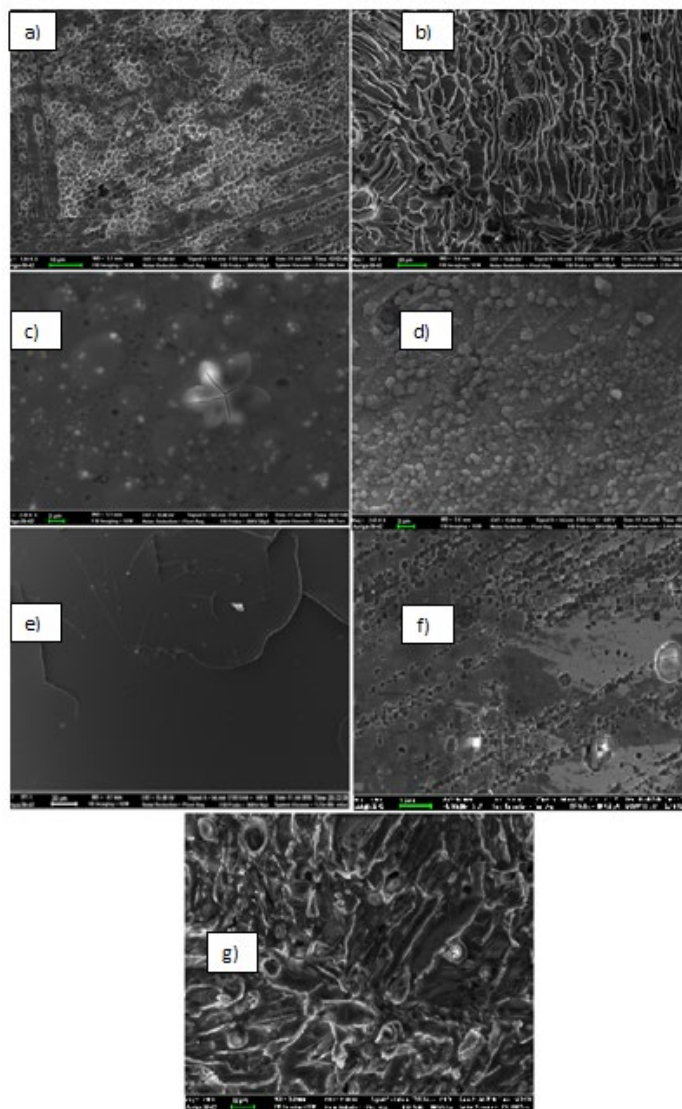


Fig. 7: SEM surface morphologies for Ti_6Al_7Nb under dynamic SSE Two layers:
a) Pure SSE, b) 0.5 g/l NaF, c) 0.1% H_2O_2 , d) 2.5 g/l, e) 5 g/l, f) 10% H_2O_2 , g) SSE at pH 3.0.

Conclusions

Corrosion behavior of a Ti₆Al₇Nb alloy, when is used as a dental implant, has been characterized in this study. From various dental application exposure, the following conclusions were drawn:

Excellent passive layer forms when Ti₆Al₇Nb is exposed to pure simulating saliva electrolyte (SSE).

The increase of the fluoride concentration increases corrosion potential and corrosion kinetics, and the critical fluoride concentration was 5g/l.

Peroxide at smaller concentration causes corrosion potential to shift to the electropositive value, while the higher concentration lowers stability of the anodic passive layer forms on Ti₆Al₇Nb.

A simulated severe plaque formation (SSE lowered to pH 3.0) causes the most accelerated corrosion rate on Ti₆Al₇Nb alloy.

Electrochemical impedance spectroscopy (EIS) measurements revealed a capacitive behavior with formation of two oxides layers (porous oxide and dense-compact oxide layer). Furthermore, high phase angle and impedance were noted at low and intermediate frequencies for pure SSE, but opposite occurrences were observed under dynamic modified SSE.

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