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Abstract

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Characterization of tantalum and tantalum nitride films on Ti6Al4V substrate prepared by filtered cathodic vacuum arc deposition for biomedical applications

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Abstract

This paper explores tantalum and tantalum nitride thin films produced by filtered cathodic vacuum arc deposition method as bio-stable surface treatment for Ti6Al4V titanium alloy. Effect of nitrogen to argon gas ratio on microstructure of the deposited film has been investigated. Corrosion behaviour of the films in simulated biological fluid solution was evaluated by electrochemical impedance spectroscopy. It was found that both the Ta and Ta-N films enhanced corrosion resistance of the Ti6Al4V substrate with the best protective characteristics achieved by the Ta-N film deposited at 0.25 N₂/Ar gas ratio. The protective characteristic was attributed to the formation of tantalum oxide and oxynitride compound at the surface, as verified by X-ray photoelectron spectroscopy. Increasing N₂/Ar gas ratio increased susceptibility to localized corrosion. The corrosion resistance decreased as the thickness of the film increased to 1 µm.

Keywords:

Tantalum films; tantalum nitrides; X-ray photoelectron spectroscopy; electrochemical impedance spectroscopy

1. Introduction

Prosthetic implants made of titanium alloys are widely used because of their high strength-to-weight ratio and superior biocompatibility [1]. The thin oxide surface layer (1.5–10 nm) that forms naturally on the titanium metal exposed to air at room temperature improves the stability of the surface [2]. Unfortunately, titanium implants have poor wear resistance and the generated wear debris accelerates the rate of electrochemical reactions between the surface of the implant and the surrounding physiological environment [3]. Therefore, further development is needed to extend the longevity of these implants and decrease the risk of an agonizing repeat implantation procedure. Over the past few decades transition metal nitride and oxide films have been utilized to improve the corrosion and mechanical properties of biomedical components due to their excellent resistance to wear resistance and corrosion [4–6]. The metal nitride and oxide coatings have a strong covalent bonding which leads to high hardness and chemical stability, attributes which make them suitable candidates for bioactive coatings. Several studies have shown that Ta-N films can promote cell proliferation [7,8], enhance corrosion resistance [9], has good blood compatibility and they are suitable for fabricating artificial heart valves [10]. Tantalum oxide and oxynitride (TaON) have a better corrosion resistance than the native oxide film on uncoated Ti6Al4V [11,12]. The stability of the TaON coating is attributed to the release of ammonia (NH_4^+) or nitrate ions which consumes protons and increases the overall pH [13]. Possible acidification caused by dissolution of metal is then neutralized by the ammonia. The NH_4^+ ions

undergo further reaction to form stable compounds of nitrate and nitrites, which act as either corrosion inhibitors or stabilize the passive films and improve resistance to pitting corrosion [12,14].

Despite the promising chemical and mechanical properties of metal nitrides, investigations into the tantalum nitrides prepared by filtered cathodic vacuum arc deposition (FCVAD) are limited. Tantalum films enhanced the mechanical and corrosion properties of the titanium alloys substrate, have been reported in our previous studies [15–17]. The corrosion resistance decreased with increasing nitrogen level in the sputtered Ta-N films, as reported by Mendizabal et al. [18]. The cathodic-arc Ta-N thin films improved the corrosion resistance of AISI 317L stainless steel substrate [19]. In this study, we evaluate the performance of the Ta and Ta-N films deposited on Ti6Al4V substrate for biomedical applications. The N_2/Ar gas ratio during film deposition was varied to study its effect on the structure and electrochemical properties of the films. In order to simulate and examine the performance of the coating in the human body and understand the reaction mechanisms, Ta and Ta-N films was tested in simulated biological fluid using electrochemical impedance spectroscopy (EIS). The surface chemistry of the samples before and after corrosion tests was then examined using X-ray photoelectron spectroscopy for a better understanding of the corrosion protection mechanism.

2. Experimental procedures

2.1. Deposition conditions

Ti6Al4V sheet substrates of $20 \times 20 \times 2 \text{ mm}^3$ were ground and polished using $15 \text{ }\mu\text{m}$ diamond polycrystalline suspension and then polished with an oxide polishing suspension consisting of 2 ml ammonia and 1 ml hydrogen peroxide. The deposition was carried out using an in-house built filtered cathodic vacuum arc deposition and the deposition parameters are summarized in Table 1. The arc cathode (the target) was made of high purity tantalum (purity, 99.99 %) with a diameter of 58 mm. The chamber in the deposition system reached a base pressure of $1.0 \times 10^{-2} \text{ Pa}$ prior to deposition. The DC arc current and substrate voltage bias were set to 200 A and -100 V , respectively for all the samples. The N_2/Ar gas ratio was varied from 0–0.75, and is defined as the ratio of the mass flow rate of nitrogen to the total mass flow rate of argon and nitrogen. Gas pressure during deposition was between 0.35–0.40 Pa, as measured with a MKS capacitor gauge. To investigate the influence of film thickness on the electrochemical performance, two samples with N_2/Ar gas ratio of 0 and 0.50 were prepared using a longer deposition time of 30 min. The thickness of the 5 minutes deposited films was found to vary in the range of 233–365 nm. The thickness of the films with a deposition time of 30 minutes at 0 and 0.50 N_2/Ar gas ratio were 0.7 μm and 1 μm , respectively.

2.2. Surface analysis

Grazing-incidence X-ray diffraction (XRD) tests of the deposited films were carried out with a GBC MMA diffractometer equipped with Cu $K\alpha$ incident radiation. The XRD scan was in the 2theta mode at a glancing angle of 5° from 30° to 80° in 0.02° steps. Surface topography and roughness of the films were examined using a Veeco DimensionTM 3100 atomic force microscope (AFM) operating in contact mode over a scanning area of $2 \times 2 \text{ }\mu\text{m}^2$ with a scanning

rate of 1 Hz. The contact angle of the surface was measured by the sessile drop method in a contact angle goniometer (Rame-Hart Instrument). 2 μl of the testing liquid, i.e. deionized water and fetal bovine serum (FBS) respectively, was used to measure the contact angle on the surface, these measures were repeated five times and an average value is reported. The thickness of the films was measured from the cross-section transmission electron microscope (TEM) specimens prepared using focused ion beam (FIB) technique on an FEI xT Nova Nanolab 200 work station.

2.3. Electrochemical measurements

Measurements of electrochemical impedance spectroscopy (EIS) were taken with a Gamry Reference 600 potentiostat (Gamry Instruments, Warminster, PA, USA). A conventional three-electrode electrochemical cell was used with platinum and Ag/AgCl as auxiliary and reference electrodes, respectively. The working electrode surface area of 0.8 cm^2 was exposed to the test electrolyte via a glass cell affixed to the surface by an O-ring and clamp. A simulated biological fluid (SBF) solution was used as the test solution because its ionic composition is similar to human blood plasma. The SBF was prepared by dissolving analytical grade mixtures of NaCl, NaHCO_3 , Na_2CO_3 , KCl, $\text{K}_2\text{HPO}_4 \cdot 3\text{H}_2\text{O}$, $\text{MgCl}_2 \cdot 6\text{H}_2\text{O}$, CaCl_2 , and Na_2SO_4 in distilled water. The HEPES (4-(2-Hydroxyethyl) piperazine-1-ethanesulfonic acid) was dissolved in 0.2 M NaOH solution in a separate volumetric flask before adding it into the mixture. The pH of SBF solution was 7.4 ± 0.5 . The electrochemical cell was placed in a Faraday cage in order to minimize interference from other electromagnetic devices. The sample was exposed to the electrolyte for 60 min at 37 $^\circ\text{C}$ prior to the electrochemical testing in order to establish the steady-state potential. Impedance measurements were conducted in the frequency range of 0.01 Hz–10 kHz with a data

density of 10 points per decade at their respective open circuit potentials. The measurements were carried out by applying sinusoidal wave perturbations of 10 mV RMS in amplitude. The samples were then analyzed by the JEOL JSM-6490 SEM. X-ray photoelectron spectroscopy spectrometer (XPS) was performed using a Specs SAGE 150 XPS operated with Mg K α X-ray source at 10 keV and 10 mA. The data were calibrated to the adventitious C1s peak present at 284.8 eV to minimize sample charging.

3. Results and discussion

3.1. Film structure and surface topography

Fig. 1 shows the XRD patterns of uncoated Ti6Al4V substrate, Ta and Ta-N films prepared at various N₂/Ar gas ratios. The XRD pattern of uncoated Ti6Al4V substrate agrees with the characteristics peaks of Ti6Al4V substrate reported in the literature [20–22]. The Ta and Ta-N films exhibit peak broadening from 33° to 38°, indicating a significant amount of amorphous phase in the film deposited. Some peaks from the substrates are observed in the spectra of the film samples due to the thin film thickness. At pure Ar gas pressure the film has a tetragonal β -Ta structure (JCPDS-25-1280), but as the N₂/Ar gas ratio increased to 0.25, 0.50 and 0.75, minor peaks which correspond to hexagonal Ta-N (JCPDS-39-1485) appeared, as did tetragonal β -Ta. The result is consistent with the work reported by Kim and Cha [23], except that orthorhombic Ta₃N₅ was not found in this work. The structure of tantalum nitrides is formed by the closely packed Ta atoms with N atoms inserted in interstitial sites [24]. A wide range of nitrogen-rich compounds (Ta₅N₆, Ta₄N₅, Ta₃N₅) may form as the amount of nitrogen increases. The

complexity of the Ta-N compounds is well reported in Ref. [25]. The equilibrium and metastable phases formed depend on the deposition procedures and conditions.

Fig. 2 presents the AFM deflection images of the Ta-N films at different nitrogen to argon gas ratios, with a Z-range of 30 mV. Pure Ta film (0 N₂/Ar gas ratio) has a very smooth surface with an average roughness of 3 nm. The smooth surface of the pure Ta film is attributed to the growth of an amorphous film, which is in accordance with the XRD result. As the N₂/Ar gas ratio increases the surface topography shows island style growth of about 100 nm (Fig. 2 (b)). The surface of the film with a 0.50 N₂/Ar gas ratio, as shown in Fig. 2 (c) tends to disintegrate between the colonies of the islands and makes the surface rougher, a result attributable to the reduced surface mobility of the atoms added by the formation of Ta-N bonding during deposition. The average surface roughness of the TaN film is 25 nm and 17 nm for 0.25 and 0.50 N₂/Ar gas ratios, respectively. The surface becomes smoother ($R_a=10$ nm) in the Ta-N film of 0.75 N₂/Ar gas ratio, probably due to the prolonged time for the migration of surface atoms due to a very low growth rate.

3.2. Wettability test of the Ta and Ta-N film surfaces

Surface wettability and surface biocompatibility are important parameters which influence cell growth. The average contact angle of water and FBS is reported in Table 2. The contact angles of water and bovine serum of the Ta-N coated samples are higher than the contact angle of the uncoated sample. However, the sample deposited at pure Ar gas pressure (0 N₂/Ar gas ratio) has a high water contact angle and a low contact angle when the sample was exposed to the FBS.

This means that while the pure Ta surface presents hydrophobicity, it does not restrain the affinity between FBS and the film surface as much. The Ta-N films (N_2/Ar gas ratio of 0.25, 0.50 and 0.75) has similar contact angles for both water and FBS from 81° – 88° , indicating little change in the contact angle in relation to the presence of Ta-N phase in the films. Literature suggests that cell proliferation increases with increasing surface wettability [26,27]. Here, a contact angle less than 90° for Ta and Ta-N films indicates a less hydrophobic surface that is expected to promote cell adhesion and proliferation.

3.3. Electrochemical measurements

Fig. 3 shows the impedance spectra of the Ta-N films produced at different N_2/Ar gas ratios after exposure to SBF for 1 hour. The Nyquist plot, Fig. 3 (a), shows a relatively larger semi-circle for the coated samples at 0 and 0.25 N_2/Ar gas ratios, which indicates an increased resistance of the surface layer with the most capacitive behavior for the sample prepared at 0.25 N_2/Ar gas ratio. Fig. 3 (b) shows a magnified section of the Nyquist plot in the lower range of values in Fig. 3 (a). In Fig. 3 (b) and Fig. 3 (d), it is evident that the Ta and Ta-N coated samples have much higher impedances than the uncoated Ti6Al4V substrate, which implies that the passive film formed on the coated surfaces is more protective than for the uncoated Ti6Al4V in SBF. Fig. 3 (b) and Fig. 3 (d) also suggest that there is a threshold for N_2/Ar gas ratio after which the impedance begins to decrease. Low-frequency (<1 Hz) impedance is lower in the thick 30 min Ta film with 0 N_2/Ar gas ratio, whereas the thick 30 min Ta film with 0.50 N_2/Ar gas ratio has a higher impedance than the thinner 5 minute film. The mid-frequency region of the Bode plots can be used to characterize local surface defects, while the reactions on the surface, e.g. charge transfer, are

indicated by the low-frequency region (<1 Hz) [28]. As Fig. 3 (c) shows, a significant change in the phase angle in the low-frequency region for the samples of 0.50 N_2/Ar gas ratio+5min and 0.75 N_2/Ar gas ratio+5min, showing that surface was acting like a non-ideal or leaking capacitor. This is possibly due to the passive film of the sample becoming unstable and defective after immersion in SBF, causing a large variation in the phase angle. While the phase shift reached 85° for the sample of 0.25 N_2/Ar gas ratio+5min, implying a passive film present on the surface with highly capacitive response. Ti6Al4V substrate has the lowest phase shift in the low-frequency range of the Bode plot, as shown in Fig. 3 (d). However, the passivating layer on top of the Ta-N films acts like an insulating layer that offers corrosion protection in a buffered biological environment. Similar protective properties of Ta-N films on 316L stainless steel attributable to the formation of a protective surface layer, was reported by Alishahi et al [29].

Table 3 presents a quantitative analysis of EIS data by fitting the appropriate electrical equivalent circuit (EEC). The EEC model with two loop elements, as shown in Fig. 4 (a) produced a good fit on the impedance data for Ti6Al4V substrate and Ta-N coated samples except for the film with 0.25 N_2/Ar gas ratio+5min. The impedance data for the sample of 0.25 N_2/Ar gas ratio+5min fit best with the EEC model with one time constant (loop element), as shown in Fig. 4 (b). In the circuit, R_s represents the resistance of the solution between the working and reference electrodes, $Q_{oxide\ film}$ and $R_{oxide\ film}$ are constant phase element (CPE) and resistance associated with the oxide film on the surface. The constant phase element (CPE) is indicative of the capacitive contribution and its deviation from ideal dielectric behaviour due to surface heterogeneities such as roughness, inhomogeneous surface, absorption of ions or possible variations in the physical properties of the covering film [30]. The exponent n indicates a

variation from ideal capacitor/resistor where $n = 1$ for the pure capacitor, and $n = 0$ for the pure resistor. Q_{dl} and R_{ct} are the CPE and resistance associated with the electrical double layer and charge transfer resistance, respectively. The simulation model with one time constant for the 0.25 N₂/Ar gas ratio+5min indicates the formation of a uniform oxide layer without defects. The 0.25 N₂/Ar gas ratio+5min has the highest charge transfer resistance of 3607 kΩ cm² of the tested samples. This result indicates that the surface itself affords the best corrosion protection due to the stability of passive film.

Fig. 5 shows the SEM images of the specimens after one hour immersion in SBF and then 20 minutes of EIS testing. A layer of corrosion product was clearly visible on the pure Ta film after EIS testing, as shown in Fig. 5 (a). An EDS analysis of the corroded surface shown in Fig. 6 confirmed the presence of oxygen, tantalum, titanium, aluminium, vanadium, chlorine, sodium, calcium and potassium; the calcium, sodium, chlorine and potassium came from the SBF. Fig 5 (b), Fig 5 (c) and Fig. 5 (d) show the SEM images of Ta-N films at 0.25 N₂/Ar gas ratio+5min, 0.50 N₂/Ar gas ratio+5min and 0.75 N₂/Ar gas ratio+5min, respectively. Under the same magnification, the formation of small pits was more obvious as the N₂/Ar gas ratio increased. Fig. 7 (a) demonstrates the corroded surface at other regions of the 0.50 N₂/Ar gas ratio+5min Ta-N. It is apparent that the Ta-N film is susceptible to pitting corrosion, where the corrosive liquid diffuses into the coating and leads to corrosion pits, as shown in the Fig. 7 (b) at a magnification of $\times 1000$. Ta-N films with more nitrogen and high volume fraction of TaN is expected to result in sharper change in Young's modulus and hardness across the film/substrate interface, and hence poor film adhesion and coating delamination. The multiple phase combination of Ta, TaN and some sub stoichiometric compounds in the 0.25 N₂/Ar gas ratio sample may also benefit the

toughness and resistance to crack propagation inside the film [4, 18]. Fig. 5 shows coating delamination and more pits with large exposed areas for TaN films with 0.5 and 0.75 N₂/Ar gas ratios. The TaN film with 0.25 N₂/Ar gas ratio exhibits less defects and therefore its corrosion resistance is better than other samples with higher nitrogen level. With the Ta-N films at a deposition time of 30 minutes, i.e. Fig. 5 (e) and Fig. 5 (f), film detachment was clearly evident. The corroded surface of the thicker film is prone to formation of micro cracks. There is the possibility of a more distinct difference in reactivity between the grain boundaries and the rest of surface in the thicker film compared to that of the thinner film. As reported by Wei et al. [31], the larger stresses in a thicker coating create more local defects such as micro-fractures. Those defects allow more eroding ions to enter into the coating and worsen the corrosion resistance of the oxide coating. Similarly, the thicker Ta-N films in this study exhibit coating defects which expedited the formation of micro cracks after immersion in SBF.

The corrosion seen in Fig. 5 (a) and Fig. 5 (e) and the EDS analysis in Fig. 6 confirms the formation of a surface layer of calcium phosphate. The pure Ta film exhibits a somewhat protective effect because the pores are sealed by the precipitation of calcium phosphate in the electrolyte. Hydrated oxide film (Ta-OH group) was formed by a slow hydration of tantalum oxide layer in the SBF solution. This Ta hydroxide reacts with calcium ion and phosphate ion on the metal surface and build up a layer of calcium phosphate [32,33]. Similar results were reported by Valero Vidal and Igual Munoz [34], on the protective effect that phosphate had on the stainless steel in phosphate buffered saline solution that improves the passive behaviour by increasing the charge transfer resistance. In Ta-N coated Ti6Al4V substrate, passive film forms and protects the substrate from degradation and generates a capacitive response.

3.4. X-ray photoelectron spectroscopy analysis

Fig. 8 compares the Ta 4f core-level XPS spectra of pure Ta film before and after corrosion tests. The deconvolution of all four Ta 4f spectra was conducted with a constraining condition where all the $4f_{7/2} - 4f_{5/2}$ doublets are of a fixed area ratio of 4:3 and an energy separation of 1.91 eV [35,36]. As shown in Fig. 8 (a), Ta 4f spectrum is de-convoluted to four doublets consisting of Ta_2O_5 , Ta sub-oxides with lower oxidation states (TaO_2 and TaO) and metallic Ta^0 . The Ta $4f_{7/2}$ and Ta $4f_{5/2}$ peaks located at 26.3 eV and 28.2 eV are attributed to the most stable form of pentavalent state, Ta_2O_5 , as reported in the literature [37]. The Ta $4f_{7/2}$ and Ta $4f_{5/2}$ peaks located at 25.0 eV and 26.9 eV are corresponding to TaO_2 , peaks located at 23.5 eV and 25.4 eV, and are categorized to the TaO. The binding energies of the suboxides are close to the values reported by Moo et al [38]; these sub-oxides may exist in the surface layer or may occur due to a reduction of the oxide by the preferential removal of oxygen during Ar etching before XPS scanning [38]. The binding energies of the low energy-doublet (22.0 eV and 23.9 eV) are attributed to metallic tantalum, Ta^0 [36]. Note that the Ta 4f spectrum of pure Ta film has a relatively large shoulder at the low energy doublet, which implies the deposited film contains mainly metallic Ta. After corrosion test, the peak width (Fig. 8 (b)) at the low energy doublet decreased significantly and the spectrum shows two primary peaks of Ta $4f_{7/2}$ and Ta $4f_{5/2}$ peaks located at 26.3 eV and 28.2 eV; this result indicates that the formation of a passive layer of Ta_2O_5 .

For Ta-N film at 0.25 N_2/Ar gas ratio+5min, the deconvolution of Ta 4f spectrum is shown in Fig. 9 (a), where the Ta $4f_{7/2}$ and Ta $4f_{5/2}$ peaks positioned at 26.3 eV and 28.2 eV correspond to

Ta₂O₅; there are small peaks corresponding to the TaO₂ are also observed (25.0 eV and 26.9 eV). The doublet with the Ta 4f_{7/2} and Ta 4f_{5/2} peaks is in close agreement with the values of 25.6 eV and 23.7 eV reported for hexagonal TaN [39]. The TaN peaks are consistent with the result obtained for XRD, where hexagonal TaN phase is observed as the N₂/Ar gas ratio increased. Representative peaks of metallic Ta⁰ are also observed in the surface. After corrosion test, the Ta 4f peaks of Ta₂O₅ were shifted to lower binding energies by 0.5 eV to 25.8 eV and 27.7 eV for the 4f_{7/2} and Ta 4f_{5/2} peaks, respectively (Fig. 9 (b)). The peak position of O 1s shifted to a lower energy by 0.5 eV after corrosion test, as shown in Fig. 9 (c). The high resolution Ta 4p_{3/2}/N 1s spectra in Fig. 9 (d) show similar peak position before and after corrosion tests. The XPS result indicates the formation of TaON layer on the surface. This result is consistent with the work reported by Xu et al [12], where the binding energies of the Ta 4f peaks for pure Ta₂O₅ coating shift to a lower energy by 0.6 eV for pure TaON coating. The shift in binding energies is caused by covalent bond between Ta and N which is greater than between Ta and O where the oxygen is more electronegative than nitrogen. The peak intensity and peak width of the 4f_{5/2} and Ta 4f_{7/2} peaks (21.4 eV and 23.3 eV) at low-energy doublet corresponding to metallic Ta⁰ decreased after the corrosion test. The TaOx phase on the surface layer acts like a charge barrier layer and protects the substrate from corrosion. As reported by Xu et al. [12], the TaON coating with a lower charge carrier density, exhibited a higher corrosion resistance than the β -Ta₂O₅. The charge donor density (N_d) is highest for the passive film formed on bare Ti6Al4V alloy ($3.76 \times 10^{20} \text{ cm}^{-3}$), followed by the β -Ta₂O₅ ($9.89 \times 10^{18} \text{ cm}^{-3}$) and TaON ($7.19 \times 10^{18} \text{ cm}^{-3}$) coatings as calculated from Mott-Schottky plots. This means that the protective oxide film provided by the TaON coating is the most stable and insulating, therefore offers more protection to Ti6Al4V alloy than the β -Ta₂O₅ coating [12]. The Ta-N film at 0.25 N₂/Ar gas ratio+5min displayed a higher

corrosion resistance compared to the pure Ta film in this work most likely because of the good passivation effect by the TaON due to its lower charge carrier density. The XPS results confirmed the passive surface oxide and TaON on the surfaces of pure Ta film and Ta-N film at 0.25 N₂/Ar gas ratio+5min after corrosion process, respectively.

The formation of TaON could be attributed to electrochemical reactivity of TaN in aqueous media due to its low isoelectric point of 1.0 [40] and presence of negative charge on TaN surface at pH 7.4. This promotes absorption and interaction with the electronegative species thus formation of oxide or hydrated oxide TaN, i.e. TaON or TaON_xH₂O. Similar mechanism has been proposed elsewhere [41] for formation of TaO_xN_y upon exposure of TaN film to SBF. TaON layer would intrinsically exhibit higher inertness compared to TaN and therefore will afford more effective passivation. The TaON in the surface provided satisfactory resistance to corrosion and leads to a significant improvement of the corrosion resistance in the Ta-N film with a proper thickness of 300 nm and with mechanical integrity.

4. Conclusions

Ta and Ta-N films were deposited by filtered cathodic vacuum arc deposition technique. A hexagonal Ta-N phase in the resultant films was identified with an increase in the N₂/Ar gas ratio. Ta-N films enhanced the corrosion resistance due to the stability of the passive oxide and oxynitride layers. The low charge carrier density of TaON layer was verified by XPS results and the high electrochemical impedance. A combination of passive chemistry, uniformity and lack of defect in the film made resulted in optimized protective performance by Ta-N film at 0.25 N₂/Ar

gas ratio. Further increase in N_2/Ar gas ratio results in defective film and hence the unstable surface film is susceptible to corrosion attack in simulated biological fluid solution. These results show that Ta-N film prepared by filtered-cathodic-arc deposition technique in an optimized atmosphere and thickness showed improved corrosion resistance that makes Ta-N film a promising candidate for bio-implantable devices.

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Figure Captions

Fig. 1 X-ray diffraction patterns of the uncoated Ti6Al4V and Ta-N films at various N_2/Ar gas ratios.

Fig. 2 AFM deflection images of the Ta-N films at N_2/Ar gas ratio of (a) 0 (b) 0.25 (c) 0.50 (d) 0.75, all with a Z- full range of 30 mV.

Fig. 3 Nyquist and Bode impedance spectra of Ta-N films at different N_2/Ar gas ratios.

Fig. 4 Electronic equivalent circuits (EEC) used in the fitting procedure of the EIS experimental data.

Fig. 5 SEM images of Ta and Ta-N films after electrochemical measurement (a) 0 N_2/Ar gas ratio+5min; (b) 0.25 N_2/Ar gas ratio+5min; (c) 0.50 N_2/Ar gas ratio+5min; (d) 0.75 N_2/Ar gas ratio+5min; (e) 0 N_2/Ar gas ratio+30min; (f) 0.50 N_2/Ar gas ratio+30min.

Fig. 6 Energy dispersive X-ray spectrum of the sample 0 N_2/Ar gas ratio+5min. Data is associated with the area shown in Fig. 5 (a).

Fig. 7 SEM images of 0.50 N_2/Ar gas ratio+5min Ta-N film after electrochemical measurement at magnification of (a) $\times 50$ (b) $\times 1000$.

Fig. 8 X-ray spectroscopy spectra of the (a) pure Ta film before corrosion test; (b) pure Ta film after corrosion test.

Fig. 9 X-ray spectroscopy spectra of the Ta-N film at 0.25 N_2/Ar gas ratio+5min (a) Ta 4f spectrum before corrosion test; (b) Ta 4f spectrum after corrosion test; (c) O 1s spectra; (d) Ta 4P/N 1s spectra.

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Table 1.

Deposition parameters of the Ta and Ta-N films.

N ₂ /Ar gas ratio	Mass flow rate		Average ion beam current (A)	Deposition time (min)
	N ₂ (sccm*)	Ar (sccm*)		
0	0	20	0.04	5
0.25	5	15	0.06	5
0.50	10	10	0.08	5
0.75	15	5	0.10	5
0	0	20	0.04	30
0.50	10	10	0.08	30

*sccm (standard cubic centimetre per minute)

Table 2.

Contact angle values of each sample. Note that N₂/Ar gas ratio of 0 indicates that the Ta coating was deposited in pure argon atmosphere.

Sample	Ti6Al4V		N ₂ /Ar gas ratio			
	substrate	0	0.25	0.50	0.75	
Water contact angle (°)	70.3 ± 2	89.1 ± 2	87.1 ± 1	87.4 ± 2	86.5 ± 1	
Fetal bovine serum contact angle (°)	77.2 ± 1	73.7 ± 1	88.1 ± 3	85.4 ± 2	81.7 ± 1	

Data are shown as mean ± standard deviation.

Table 3.

Electrical components calculated by fitting an equivalent electrical circuit on the EIS data.

	$Q_{oxide\ film}$	n_1	$R_{oxide\ film}$	Q_{dl}	n_2	R_{ct}
	($\mu\Omega^{-1}\ s^n\ cm^{-2}$)		($k\Omega\ cm^2$)	($\mu\Omega^{-1}\ s^n\ cm^{-2}$)		($k\Omega\ cm^2$)
Ti6Al4V	31.3	0.90	18.3	855	0.91	9.58
substrate						
0 N ₂ /Ar gas	14.1	0.94	133	2.28	0.84	336
ratio+5min						
0.25 N ₂ /Ar gas				12.2	0.95	3607
ratio+5min						
0.50 N ₂ /Ar gas	12.4	0.80	36.7	58.3	0.80	604
ratio+5min						
0.75 N ₂ /Ar gas	13.2	0.95	13.6	18.9	0.73	282
ratio+5min						
0 N ₂ /Ar gas	29.0	0.87	733	47.5	0.87	903
ratio+30min						
0.50 N ₂ /Ar gas	21.9	0.89	43.1	10.7	0.89	494
ratio+30min						

Table 4.

XPS result for

Sample	Ta +TaOx (at. %)	N (at. %)	O (at. %)	C (at. %)	Ca (at. %)
Ti-6Al-4V substrate	10.5	-	49.6	37.7	2.3
0 N ₂ /Ar gas ratio	26.1	3.9	53.5	16.5	
0.25 N ₂ /Ar gas ratio	18.7	7.6	42.9	30.8	
0.50 N ₂ /Ar gas ratio	17.8	8.4	40.3	33.4	
0.75 N ₂ /Ar gas ratio	21.0	6.5	44.8	27.8	

Highlights

- Incorporation of nitrogen improves the corrosion resistance of Ta coated Ti6Al4V.
- XPS results confirm the formation of a TaON after corrosion of the TaN film surface.
- The improved corrosion resistance was attributed mainly to the formation of TaON.

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Figure 1

Figure 2

Figure 3

(a)

$R_s(Q(R(QR)))$

(b)

$R_s(QR)$

Figure 4

Figure 5

Figure 6

Figure 7

Figure 8

Figure 9