



Laser surface deposition of niobium and titanium-niobium on Ti6Al4V substrates for biomedical applications

S.L. Pityana^{1,2}, N. Baloyi¹, M. Tlotleng^{2,3}, and P. Popoola¹

Affiliation:

¹Tshwane University of Technology, Pretoria, South Africa

²CSIR, Pretoria, South Africa

³University of Johannesburg, South Africa

Correspondence to:

S. Pityana

Email:

SPityana@csir.co.za

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ORCID ID:

<https://orcid.org/0000-0002-9273-2043>

Synopsis

The advantages of using Nb and Ti-Nb coatings for improving the hardness and corrosion resistance of Ti6Al4V were investigated. It has been reported that Ti6Al4V used in orthopaedic implants tends to release toxic Al and V ions into the surrounding tissue. Thin layers of Nb and Ti-Nb were deposited on Ti6Al4V substrates using a laser metal deposition technique. The deposited material was analysed using optical microscopy, scanning electron microscopy coupled with energy dispersive spectroscopy, and X-ray diffraction. The corrosion behaviour of the deposited layers was investigated using a Metrohm Autolab PG Stat101 compact potentiostat at 25°C in simulated body fluid. A Vickers hardness system was used to study the mechanical properties. Both Nb and Ti-Nb coatings exhibited good metallurgical bonding with the substrate. The microstructure and the XRD analyses for the Nb system showed that the α -Nb phase was most dominant, while the Ti-Nb system comprised a mixture of the α and β phases. The average hardness of the Ti6Al4V substrate was 350 HV_{0.3}, with a slight increase for the Nb coating (363 HV_{0.3}) and a much higher hardness from the Ti-Nb coating at 423 HV_{0.3}. The corrosion results show that the deposited Nb was more corrosion-resistant in the solution than either the Ti6Al4V substrate or the Ti-Nb coating.

Keywords

Corrosion, laser surface deposition, niobium (Nb), titanium (Ti), titanium-niobium (Ti-Nb), Ti6Al4V.

Introduction

Metals such as CoCrMo, 316L stainless steel, and NiTi alloy are used for manufacturing biomedical implants because of their distinct properties, including low Young's modulus and resistance to corrosion (Ryan, Pandit, and Apatsidis, 2006). These implants are required to remain in the body of the patient for many years. Due to a favourable combination of mechanical and corrosion properties, titanium (Ti) and its alloys have emerged as the material of choice for biomedical implants. Nevertheless, their use is not entirely without challenges. For example, in total hip replacement, surface degradation and release of metal debris from metal erosion caused by friction limit the service life of the implant (Oldani and Dominguez, 2012; Manivasagam, Dhinasekaran, and Rajamanickam, 2010). Ti6Al4V is generally regarded as the material of choice, but it has been found that over extended periods of time, the combination of surface corrosion and erosion releases toxic aluminium (Al) and vanadium (V) ions into the body (Khan, Williams, and Williams, 1996). This has led to the investigation of titanium alloys containing niobium (Nb), zirconium (Zr), tantalum (Ta), and molybdenum (Mo) as substitutes for the aluminium- and vanadium-containing Ti6Al4V (Okazaki *et al.*, 1997).

The objective of this work is not to alter the mechanical properties of Ti6Al4V, but rather to improve its surface properties by deposition of a biocompatible layer with improved wear resistance when subjected to friction. It is known that the titanium matrix can be stabilized as the α -Ti (hcp) or β -Ti (bcc) phase, depending on the element used; and it is acknowledged that alpha and beta alloys have different mechanical properties (Petrzhik, 2013). According to Rosenberg, Starosvetsky, and Gotman (2003), Zorn *et al.*, (2005), and Godly, Starosvetsky, and Gotman (2004), Nb alloys are noteworthy due to their remarkably low elastic modulus, excellent corrosion resistance, and the presence of a metastable β phase when dissolved in the Ti structure (Oldani and Dominguez, 2012; Kuphasuk *et al.*, 2001). As a result, the β -Nb phase has a higher hardness than Ti6Al4V. This β -Nb phase is also useful as it creates a passive layer when exposed to a corrosive environment such as body fluid (Kuphasuk *et al.*, 2001; Lee, Ju, and Chern Lin, 2002).

In this work, Nb and a mixture of Nb and Ti were investigated as surface coatings on a Ti6Al4V substrate. Thin layers of Nb and Ti-Nb were deposited on Ti6Al4V substrates using a laser deposition technique. The corrosion behaviour of the coatings was determined in a simulated body fluid (SBF) environment using a Metrohm Autolab PG Stat101 compact potentiostat at 25°C.

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Table 1

Powder specifications and process parameters for laser deposition

Type of alloy	PSD (μm)	Composition (wt% Nb)	Power (kW)	SS ($\text{mm}\cdot\text{s}^{-1}$)	BD* (mm)	PFR* ($\text{r}\cdot\text{min}^{-1}$)	Carrier gas flow ($\text{L}\cdot\text{min}^{-1}$)
Nb	210-10	100	0.5, 1.0, 1.2	9	4	3	5
Ti-Nb	350-10	87-13	1.0, 1.2, 1.5	9	4	3	5

* PSD: particle size distribution; SS: scanning speed; BD: beam diameter; PFR: power feed-rate

Experimental set-up

Laser deposition

A Rofin Sinar DY044, CW Nd:YAG laser system incorporating a three-way nozzle was used for the deposition. The laser head, mounted on a Kuka robot arm, was set at a standoff distance of 12 mm above the substrate. The laser beam spot and the scanning speed were kept constant at 4 mm and $9 \text{ mm}\cdot\text{s}^{-1}$, respectively. Nb was deposited on Ti6Al4V using laser beam powers of 500, 1000, and 1200 W. Similarly, the laser beam power used for depositing Ti-Nb was 1000, 1200, and 1500 W. In order to prevent oxidation, the process was conducted in a rectangular container which was covered and purged with argon gas at $5 \text{ L}\cdot\text{min}^{-1}$. The process parameters used in these experiments are summarized in Table 1.

Material preparation and characterization

Single and multiple tracks of Nb and Ti+Nb were laser-deposited onto the surface of Ti6Al4V. The Nb powder had a mean particle size (d_{50}) of $100 \mu\text{m}$. The Ti-Nb powder was obtained by mechanical mixing and pre-alloying of elemental Ti and Nb powders, using a Restch PM 400M ball mill. The particle size range of the pre-alloyed Ti-Nb powder was $10\text{--}50 \mu\text{m}$. Ti6Al4V substrate specimens, with dimensions of $30 \times 30 \times 8 \text{ mm}$, were sandblasted and cleaned with acetone prior to the laser metal deposition process.

After deposition, the samples were sectioned in the transverse direction through the deposited layer for phase and microstructure analyses. The sectioned samples were mechanically polished and etched with either Keller's (Nb coatings) or Kroll's (Ti-Nb coatings) reagents. The microstructures of the Nb and Ti-Nb layers were examined by optical microscopy and scanning electron microscopy (SEM) using a JOEL instrument, and the chemical composition was analysed by SEM using energy dispersive X-ray spectroscopy (EDX). The microhardness of the layers was measured using a Matsuzawa Vickers hardness tester with a load of 300 g at intervals of $100 \mu\text{m}$ from the top of the cladding to the cladding-substrate interface.

Phase analyses of the clad layers were performed by X-ray diffraction (XRD) at 40 kV and 30 mA. X-ray crystallography was performed using a Ni-filtered Cu K radiation source. Their matching phases were identified with the characteristic peaks of those in the files of the Joint Committee on Powder Diffraction Standards (JCPDS) references (JCPDS, 1995).

Polarization and qualitative electrochemical impedance spectroscopy measurements on the coated surfaces were carried out in 50 ml of Hanks solution at room temperature (approx. 25°C) using a Metrohm Autolab PG Stat101 compact potentiostat, consisting of a three-electrode cell: a saturated calomel electrode ($\text{Hg}/\text{Hg}_2\text{Cl}_2$) in saturated KCl as the reference electrode and platinum as the counter-electrode. The samples were placed in a glass cell and a sample area of approximately 1.1 cm^2 was exposed to the electrolyte. The system was interfaced with a computer to determine the open circuit potential (OCP). The anodic and cathodic Tafel plots were recorded using NOVA software.

Results and discussion

Optical and scanning microscopy

The microstructures of typical single-layer Nb and Ti-Nb coatings produced by the laser deposition process on Ti6Al4V substrates are shown in the optical images in Figure 1. The images show that the deposited layers had bonded well with the substrate metal and formed coatings. The metallurgical interaction between the coatings and the substrate is more visible in the case of the Ti-Nb coating, which displays a pronounced dilution region as shown by the arrow in Figure 1b. The large, visible pores in the coatings are thought to be due to vaporization of the coating material. Of note is the good metallurgical bond between the cladding and the substrate.

Figure 2 shows optical micrographs of the cross-sections of the thin Nb layers deposited on the Ti6Al4V substrate at laser powers of 500 W, 1000 W, and 1200 W. The images indicate the presence of honeycomb structures at 500 W, a mixture of honeycomb and needle-like structures at 1000 W, and needle-like structures at 1200 W.

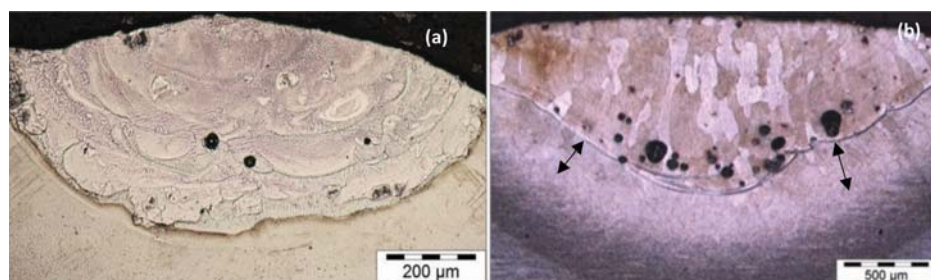


Figure 1—Typical single laser-deposited tracks of (a) Nb and (b) Ti-Nb coatings

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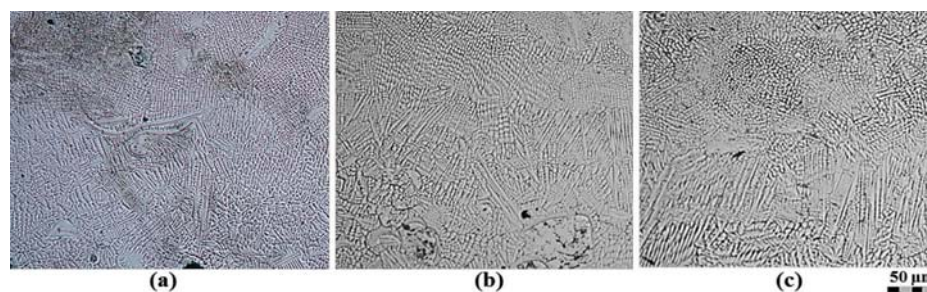


Figure 2—Cross-sections of Nb coatings deposited at (a) 500 W, (b) 1000 W, and (c) 1200 W

Table II

EDX analysis (wt%) of Nb coatings deposited at different laser powers

Element	500 W	1000 W	1200 W
C	4.77	5.26	3.84
O	2.31	1.83	1.71
Al	1.86	1.88	2.14
Ti	27.87	33.09	41.65
V	1.13	1.23	1.89
Nb	62.05	56.71	48.76

Table II lists the compositions of the Nb coatings. The data indicates the effect of laser power on the dilution effect of Nb deposited coating in terms of Nb, Ti, Al, and V content. At 500 W laser power, the Nb content is 62%, decreasing markedly as the laser power increases. This decrease is attributed to increased dilution of Nb by the Ti matrix. The contents of Al and V remain within the range 1–2%, which is lower than that of Ti6Al4V (substrate).

Figure 3 shows optical micrographs of the cross-sections of the Ti-Nb layers deposited at different laser powers. Due to the effect of pre-alloying Nb and Ti, the microstructures obtained from this combination are fine and granular. The grain size varies with increasing laser power due to slow cooling.

Table III shows the EDX analyses of the Ti-Nb coatings deposited at different laser powers. The composition of the coating does not follow any trend with relation to change in laser power. The concentration of Al and V remained in the range 1.5–4%, whereas the Nb content varied from 6.5% to 10%.

X-ray diffraction

Figure 4 shows the XRD patterns of the Nb and Ti-Nb coatings

Table III

EDX analysis (wt%) of Ti-Nb coatings deposited at different laser powers

Element	1000 W	1200 W	1500 W
C	6.14	5.56	5.83
O	4.35	4.37	5.13
Al	3.34	3.91	3.09
Ti	75.23	77.74	74.28
V	1.98	1.81	1.41
Nb	8.95	6.59	10.26

deposited at various laser powers. There are no significant phase changes in either system when the laser power is increased. The pattern for Nb shows an orthorhombic (α') phase and a metastable β phase. The α' phase is attributed to the Ti6Al4V substrate; and the β phase is due to the Nb in the melt pool. The XRD results for the Ti-Nb system also show the occurrence of hexagonal (α' -Ti) phase and the presence of β phase that seem to occur at the same angle for all the samples.

Microhardness

Microhardness profiles of the Nb and Ti-Nb deposits are shown in Figure 5. Figure 5a represents Nb deposition at various powers. All the curves indicate that hardness increases from the top of the layer to the heat-affected zone (HAZ) and then decreases again towards the substrate. Figure 5b depicts the profile for Ti-Nb deposition at various powers. All the curves indicate a steady decrease in hardness value from the top of the cladding to the HAZ and the substrate. The average hardnesses of the deposited materials relative to the substrate are plotted in Figure 6. The Nb deposit has an average hardness of 363 HV_{0.3} on the points taken on the deposited layer and 350 HV_{0.3} closer to the substrate. The average hardness of the Ti-Nb deposit was 423

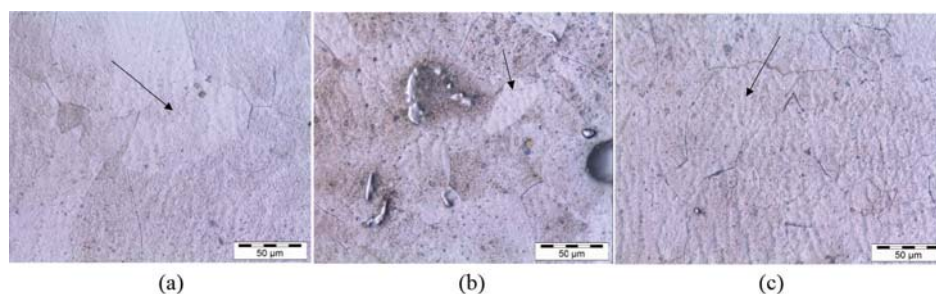


Figure 3—Cross-sections of Ti-Nb coatings deposited at (a) 1000 W, (b) 1200 W, and (c) 1500 W

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HV_{0.3}, the hardness of the layers gradually decreasing to that of the substrate. This trend was apparent in all the layers. In this system (Ti-Nb), a hardness improvement was observed and the hardness of the deposit was approximately 70 HV_{0.3} higher than that of the substrate.

Figure 6 represents the microhardnesses of the materials. Nb coatings show variable hardness behaviour, with an average hardness of 364 HV_{0.3}. Ti-Nb coatings display a steady decrease in hardness, with depth Figure 5b), with an average hardness value of 423 HV_{0.3}. This indicates that Ti had a positive effect on the coatings, due to the presence of harder phases.

The increase in hardness value for Nb coatings was approximately 5%, which is attributed to dilution of Nb by the substrate. The 8.4% increase in hardness for the Ti-Nb coatings is due to the presence of α -Ti, which results in a martensitic microstructure.

Corrosion behaviour

The potentiodynamic polarization curves of all the Nb coatings in simulated body fluid (Hanks solution) electrolyte, under a standard working environment are plotted in Figure 7. The results presented were selected because of the nearest mean values of the current densities, and they indicate constant behaviour, especially during the first five minutes of immersion. The Tafel calculations have been deduced from the available data and are reported in Table IV.

The joint polarization curves in Figure 7 and the results in Table IV illustrate that the Nb coatings have better corrosion properties than the Ti-Nb coatings and the substrates. It is also observed that the corrosion potential (E_{corr}) of the Ti-Nb coating was slightly lower than that of the substrate (Table IV), while the Nb coating had the highest E_{corr} , 0.709 V higher than that of Ti-Nb.

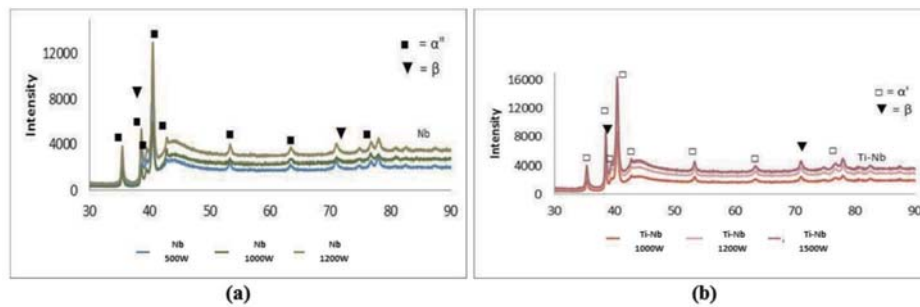


Figure 4—X-ray diffraction (XRD) patterns of laser-deposited (a) Nb and (b) Ti-Nb coatings

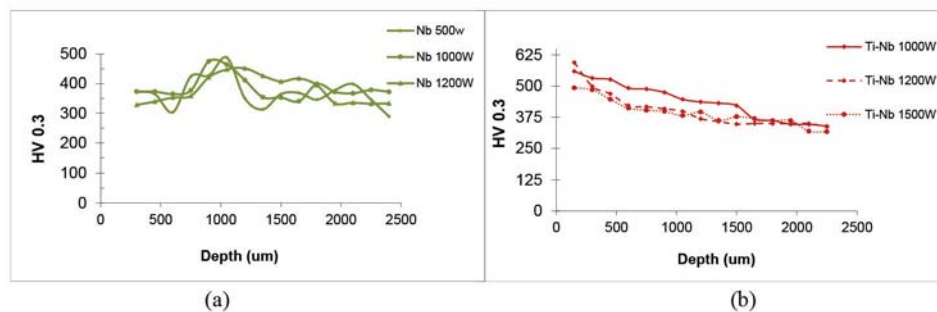


Figure 5—Hardness profiles measured from the edge of the coating. (a) Nb and (b) Ti-Nb

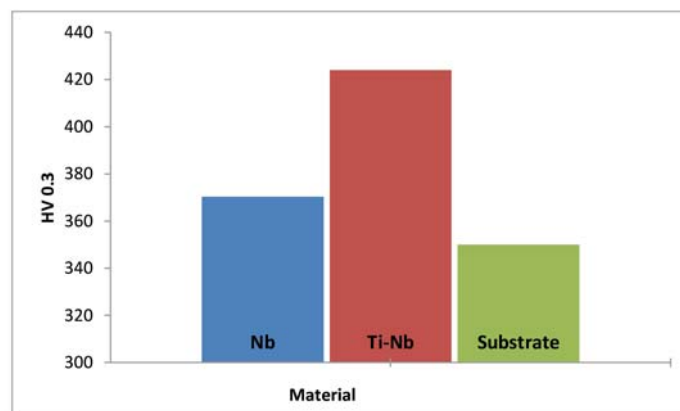


Figure 6—Average microhardness property plots for Nb and Ti-Nb coatings

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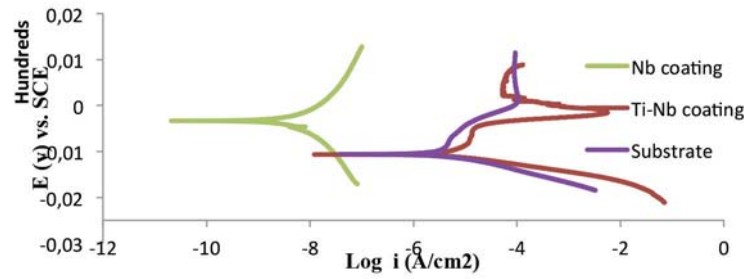


Figure 7—Potentiodynamic polarization curves of Ti-Nb and Nb deposited onto Ti6Al4V after immersion in a Hanks solution electrolyte at the standard condition

The Nb coating also exhibits the lowest corrosion current density while the Ti-Nb coating and the substrate displayed similar i_{corr} values in the same decade. This also indicates that the Nb coating is more corrosion-resistant.

According to Osorio *et al.*, (2011), the corrosion resistance of alloys depends on the microstructure and the presence of defects. This was also observed in this study. The coatings with microstructures formed in the presence of oxide layers showed passivation behaviour and consequently mitigate against further corrosion. This is seen in the microstructures of the Nb coatings; the highest resistance to corrosion was exhibited where less dilution and fewer pores were present. Furthermore, these coatings had the lowest hardness due to the low Ti content. All these factors are connected to the latent heat of materials. During heating, niobium reacted with oxygen to form an oxide layer that is resistant to corrosion. Furthermore, the refractory nature of Nb also contributes to its corrosion resistance. Similar to Nb, molten Ti (Ti-Nb) would easily react with oxygen and form a passive TiO_2 layer, which is also corrosion-resistant. This corrosion reaction of Ti-Nb seems to resemble that which occurs on Ti6Al4V surfaces when exposed to simulated body fluids, as explained by Mythili, *et al.*, (2007). The overall performance of Nb surpassed that of Ti-Nb and Ti6Al4V; irrespective of the passive layer and improved hardness due to the presence of TiO_2 , the Nb coating performed better in Hanks solution.

Conclusions

Nb and Nb-Ti were successfully coated onto Ti-6Al-4V alloy substrates using laser deposition. The hardness and corrosion behaviour of the coatings were investigated, resulting in the following observations.

- Ti-Nb deposited on the Ti6Al4V substrate resulted in a fine microstructure consisting of α' and β phases that have a higher hardness than that of the substrate. This alloy displayed a similar corrosion resistance to the Ti6Al4V substrate.
- Nb deposited on the Ti6Al4V substrate resulted in a dendritic microstructure with an uneven hardness profile. The XRD pattern indicated the presence of the α' orthorhombic phase, which is significantly more resistant to

corrosion than the substrate and Ti-Nb layers.

- The high latent heat of Nb influences the dilution, microstructure, and hardness of the coating.
- Based on corrosion results alone, Nb-cladded Ti6Al4V will be a better option than Ti-Nb as a coating to improve the surface properties of surgical implants.

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Sample	i_{corr} (A/cm ²)	E_{corr} (V)	Corrosion rate (mm/a)
Substrate	4.37E-07	-1.06	1.79E-02
Ti-Nb	9.70E-07	-1.15	4.24E-03
Nb	1.30E-10	-0.33	2.16E-08