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Chams Eddine RAMOUL, BELIARDOUH NASSER EDDINE, R. BAHY, Abdelghani DJAHOUDI, Michael J. WALOCK, Corinne NOUVEAU - Surface performances of PVD ZrN coatings in biological environments - Tribology - Materials, Surfaces & Interfaces - Vol. 13, n°1, p.12-19 - 2018

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Surface performances of PVD ZrN coatings in biological environments

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ABSTRACT

Zirconium nitride (ZrN) thin films were deposited by reactive RF magnetron sputtering on Ti-6Al-4V and Si (100) substrates for potential use in biomedical applications. The tribological behaviour was evaluated against bovine bone in dry condition using a pin-on-disc apparatus. Abrasion is the primary wear mechanism observed in ZrN/bone contact. The corrosion properties were determined through two electrochemical techniques: potentiodynamic polarization and electrochemical impedance spectroscopy. The coatings with reduced oxygen content provided: (i) good resistance against corrosion when exposed to physiological solution and (ii) better anti-bioadhesion against *Staphylococcus aureus* bacteria.

KEYWORDS

PVD coating; ZrN; wear; corrosion; bioadhesion

1. Introduction

In the field of orthopaedic implants, the challenge is to develop alloys with high biocompatibility and the requisite material properties to effectively serve as substitute bones and joints for a long period without rejection. Here, bone-implant integration is closely related to the biocompatibility (immunology/bioadhesion), wear and corrosion behaviour of both the implant (artificial organ) and surrounding bone tissue.

Transition metal nitrides, oxides and intermetallics, especially Titanium-based alloys, have been widely studied and show great potential for the protection of orthopaedic applications. Diffusion hardened Zr-2.5Nb, known as ('Oxinium'), is another category of surface engineered material used for bio implants. As recently reviewed by Dearnley [1], it appears as the better non-coated material.

In order to eliminate the implant rejection risks due to bacterial infections, newer approaches are currently being evolved. The goal is to convert the surfaces of biomedical devices into antimicrobial surfaces while preserving the corrosion and wear resistance. Nanostructured surfaces and antimicrobial agents, such as silver, used to reduce infections are the best coating strategies and could represent an alternative to antibiotics.

Recently, it was demonstrated that bacterial adhesion on Zirconium alloy films was minimized by tailoring the surface with silver ions surrounded by amorphous carbon phases using a sputtering system [2]. Unfortunately, accumulation of silver ions (Ag^+) is well known to have negative effects on general health [3]. So, the challenge becomes the development of

multifunctional surfaces with simultaneous opposite responses; on the one hand towards human cells, i.e. providing adhesion molecules for attaching tissues and cells (osseointegration) and against bacterial adhesion/proliferation (antibacterial activity) on the other hand.

Over the past two decades, ZrN coating has attracted interest for its excellent erosion resistance, biocompatibility, good ductility and excellent chemical stability [4]. In addition, there is evidence that ZrN coatings have better corrosion resistance than TiN coatings [5].

In this work, a ZrN/Zr system was evaluated for its suitability as a protective coating in biological environments. The corrosion resistance in physiological solution (PS) and tribological performances against bovine bone were investigated systematically. This was followed by a study of the material's antibacterial activity against *Staphylococcus aureus* (*S. aureus*).

This study was conducted taken into account two fundamental aspects (i) an economic aspect, i.e. a low-cost biomaterial and (ii) an environmental aspect; because recovering (recycling) only one metallic element (Zr) from an oxide is much easier than recovering two metallic elements or more from a complex structure.

2. Experimental

2.1. Coatings deposition

Zirconium Nitride films were realized by the radio frequency magnetron sputtering technique (NORDIKO

type 3500). A pure Zirconium (99.98%) disc (101.6 mm diameter, 3 mm thick) was used as the target material. The substrates are Titanium alloys (Ti-6Al-4V) intended for tribological tests and Si (100) wafers ($10 \times 10 \times 0.380 \text{ mm}^3$) for corrosion and biological tests.

Metallic substrates (cylinder $\varnothing = 20 \text{ mm} \times 4 \text{ mm}$) were polished by SiC papers grade 800. All substrates were ultrasonically cleaned for 15 min in acetone and dried in air. Before deposition, the target disc and the substrates were *in situ* etched for 5 min at 10 μbar in an Ar discharge. A pure Zr buffer layer of approximately 100 nm was first deposited (10 min, 5 μbar) to improve adhesion between the ZrN topcoat and the substrates. The target to substrate distance was fixed at 80 mm and the target bias voltage was -900 V (Zr).

In addition to improve the adhesion, a functional buffer layer is an efficient strategy for overcoming corrosion substrate problems (contact between the electrolyte and the substrates surface because of pores and other coating defects) and improving the tribological properties. Four (4) samples (ZrN-01; ZrN-02; ZrN-03 and ZrN-04) were deposited by changing Ar/(N₂+O₂) ratio, as summarized in Table 1. The total working pressure (5 μbar), deposition time ($\sim 242 \text{ min}$) and substrate temperature (150°C) were kept constant. As-deposited and post-test films were characterized by the scanning electron microscopy (SEM; JEOL JSM-5900LV-10 kV) with energy dispersive X-ray spectroscopy (EDS; Oxford INCA x-act) and optical profilometry (Veeco-Wyko NT1100).

2.1.1. Bone pins preparation

The bone pins were prepared from the femur of a bovine. The complete femur was immersed in distilled water at a constant temperature (100°C) for 2 h to remove residual soft tissue attached to the bone. From this clean bone, the pins were machined as cylinders (cylinder $\varnothing = 6 \text{ mm} \times 20 \text{ mm}$), ultrasonically cleaned (15 min) in acetone, dried in air and finally kept at $3\text{--}4^\circ\text{C}$ in a fridge (until ready for use in the pin-on-disc tribometer).

2.2. Coating characterizations

2.2.1. Tribology

Dry friction tests were conducted in ambient air at room temperature and relative humidity of $40\% \pm 5$

Table 1. Chemical composition and roughness of ZrN/Zr coatings.

Sample	Ratio Ar/(N+O ₂) (%)	Total thickness (μm)	Chemical composition (Weight %)			Roughness (μm)	
			N	O	Zr	R _a	R _q
ZrN-01	80/20	1.82	16.5	1.5	81.9	0.024	0.031
ZrN-02	70/30	1.67	16.4	1.8	81.8	0.027	0.053
ZrN-03	60/40	1.75	12.2	4.9	82.9	0.018	0.024
ZrN-04	50/50	1.72	12.9	6.1	81	0.014	0.021

using a pin-on-disc tribometer (CSM Instruments, Switzerland) in accordance to the ISO 7148, ASTM G99-95a and ASTM G133-95 standards to determine the coefficient of friction (COF). Bone pins were used as counterparts against rotating ZrN-coated Ti-6Al-4V discs.

When it is impossible to make a ball geometry, e.g. from organic matter of animal origin (bone, cartilage, etc.), the pin-on-disk configuration provided by [®]CSM Instrument tribometer is perfectly adapted for this case (Figure 1). The pin holder is positioned at 45° ; this allows to make several tests with one pin. The tribological test parameters are listed in Table 2.

2.2.2. Electrochemical analyses

The electrochemical corrosion tests were carried out using a conventional three-electrode cell of 300 ml capacity and Gamry[®] Potentiostat/Galvanostat Reference 1000. The cell was fitted with a coated Si sample as working electrode (WE), saturated calomel electrode (SCE) as the reference electrode and platinum as a counter electrode (CE). The corrosion behaviour was evaluated by measuring the open circuit potential (OCP) after 3600 s in 150 ml of 0.9% NaCl PS at room temperature with a scan rate $s = 1 \text{ mV/s}$. The corrosion potential (E_{corr}) and the corrosion current density (I_{corr}) were deduced from the Tafel plot (i.e. $\log i$ vs. E plot). Electrochemical impedance spectroscopy (EIS) tests were conducted in a frequency range from 10^5 to 10^{-2} Hz at a rate of 5 points/decade, in E_{ocp} vs. SCE. The potentiodynamic polarization test was raised from -1 to $+1 \text{ V}$, at a scan rate of 1 mV/s . The surface contact of the sample with the electrolyte was 0.5 cm^2 . EIS parameters were calculated using the ZView software (Scribner Associates, Inc).

3. Results and discussion

3.1. ZrN coatings microstructural characterization

Figure 2 shows SEM images of an as-deposited ZrN coating on a Si substrate. There are some sputtering defects, e.g. pits and droplets, as shown in Figure 2

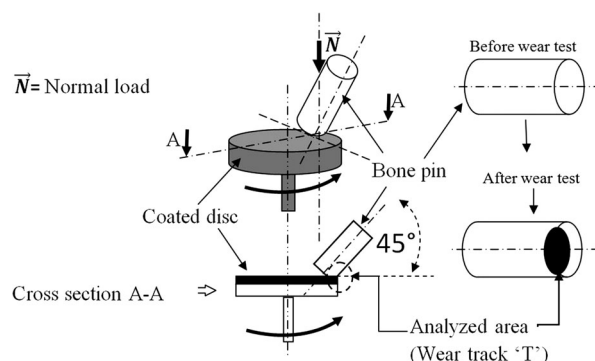


Figure 1. Schematic of the pin-on-disc tests.

Table 2. Pin-on-disc test conditions.

Parameters	Value
Velocity (cm/s)	2
Applied load (N)	2
Total sliding distance (m)	5
Wear track diameter (mm)	4
Environment	Air
Temperature (°C)	20 ± 3
Humidity (%)	40 ± 5

(a–b). All films have an overall thickness of ~ 1.6 to $1.9 \mu\text{m}$ and exhibit a columnar structure, as shown in Figure 2(c). The surface roughness of the samples is shown in Table 2. No remarkable differences in roughness parameters (R_a , R_q) are observed. The ZrN-01 and ZrN-02 samples have a relatively low oxygen content in comparison to ZrN-03 and ZrN-04.

3.2. Wear behaviour

3.2.1. COF of the ZrN coatings

Figure 3 shows the friction coefficient curves of the ZrN films sliding against bovine bone in dry conditions. It shows that the friction coefficients of the four coatings possess similar features, i.e. the friction curves of ZrN films are characterized by two different stages. The first stage represents run-in and ends with a rapid increase of the COF; the second stage was the steady state wear regime.

During dry sliding tests, the following scenario most probably occurs; at the beginning of the wear test, there is a severe material transfer between the coating and the pins, which leads to an increase of the COF. When the steady state is reached, the material transfer is slowly reduced and stabilized after a large percentage of the wear track area is covered with the transferred material. At the steady state, the highest COF value (~ 0.58) was observed for the ZrN-02 coating surface, whereas lower COF values (~ 0.38) were obtained for the other three ZrN coatings (Figure 3). Thus, the COF is fairly high. That is the disadvantage of hard ceramic coatings (ZrN), e.g. their brittleness and relatively high COF. The COF of the uncoated substrate at the



Figure 2. SEM images of the as-deposited ZrN-01 coating on Si substrate: (a) global view of the surface, (b) magnified image of the region enclosed by the white square (area A) and (c) cross-section image of the same sample.

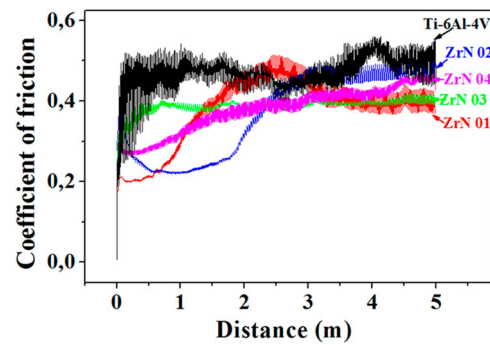


Figure 3. COF of the ZrN coatings sliding against bone counterparts.

steady state was ~ 0.58 . From Figure 3, there is clear evidence that, after 3 m of sliding distance the coatings are destroyed.

The wear volume loss of the ZrN coatings/bone couple was not measurable as it was of the order of the surface roughness of the film. In addition, the wear scars on the bone pin could not be estimated because of the low interaction between bone and light in profilometry. So, quantitative data of wear rate were not possible in this study. Only a qualitative analysis of the wear tracks was conducted using SEM/EDS.

3.2.2. Wear mechanisms of the ZrN-coated discs

Figure 4 shows the wear tracks of ZrN coatings on Ti-6Al-4V substrates tested against bone pins as counterparts in dry conditions. The primary wear mechanism appears to be abrasion, as shown in Figure 4(a–d). In addition to grooves and scratches parallel to sliding direction, wear debris accumulated in the wear tracks are also observed as deep furrows from trapped particles (Figure 4(c)). The elemental composition of some isolated particles, analysed by EDS, indicates the presence of both bone elements (Na, Mg, Ca, K, etc.) and coating elements (Zr and oxygen) in the adhered film (points 1, 2, 3, 4 in Figure 4(a–e)). So, the wear debris is a mixture of the antagonist pair components. From this, one can conclude that bone adhesion to the disc is the second wear mechanism observed in the ZrN/bone contact pair. SEM analysis also revealed some areas within the wear track where the coating did not appear to be damaged after the wear tests. No plastic deformation was detected in the wear tracks on any of the coatings.

3.2.3. Wear mechanisms of the bovine bone pins

Bone is a heterogeneous and anisotropic material. It has a hierarchically organized structure and is often classified as a ‘composite material’ [6]. At the highest hierarchical level (with feature sizes ~ 1 to 5 mm), there are two types of bone: cortical bone (which is the type used in this study) and trabecular bone. Cortical bone consists of repeating units called Haversian

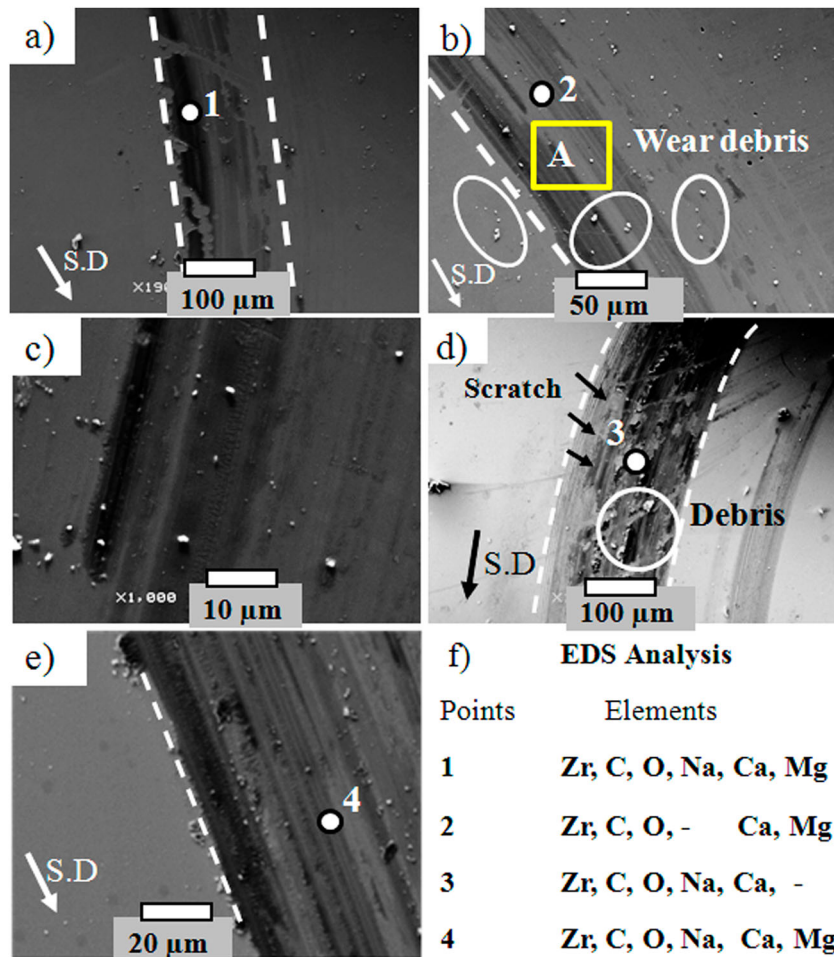


Figure 4. Wear mechanism of ZrN coatings: abrasion (scratches-grooves) and bone adhesion wear; (a) wear track of ZrN-01, (b) wear track of ZrN-02, (c) zoom of the square A, (d) wear track of ZrN-03, (e) wear track of ZrN-04 and (f) EDS analysis results at points 1, 2, 3, 4; (S.D – sliding direction).

systems or osteons. [6]. The macrostructure of the bone pin specimen (cortical bone) is shown in Figure 5.

Figure 6 (a–d) shows micrographs of the bone pin wear scars. It appears that the edges of the bone pin (see Figure 1; *analyzed area (Wear track 'T')*) become larger after acting as counterparts against ZrN-coated discs. Wear scars also reveal deformation of the Haversian system, lacunae (crevasses indicated by white narrows), isolated wear particles (black narrows) and entrapped particles (ellipse). EDS analyses, carried out at different locations (isolated as well as entrapped

particles) within the wear scars, indicate that a mixture of both the antagonist elements in contact was found. Abrasive wear is the only wear mechanism observed on the bone pins.

When sliding against ZrN-coated discs, segments of the bone pins were initially scraped off by surface asperities along the wear track. This is due to the large difference in mechanical properties between the antagonists in contact. A literature study reveals the average hardness (H) and elastic modulus (E) of cortical bovine bone obtained from nanoindentation in

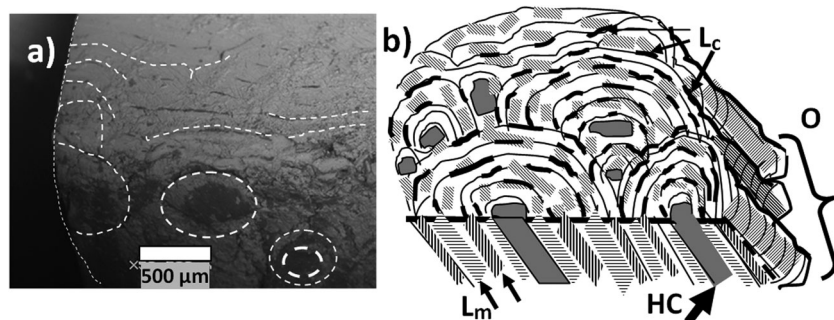


Figure 5. (a) SEM micrograph of a bone pin specimen. (b) A schematic diagram illustrating the macrostructure of cortical bone; O – Osteons (Haversian systems), L_m – concentric lamellae, HC – Haversian canal and L_c – lacunae.

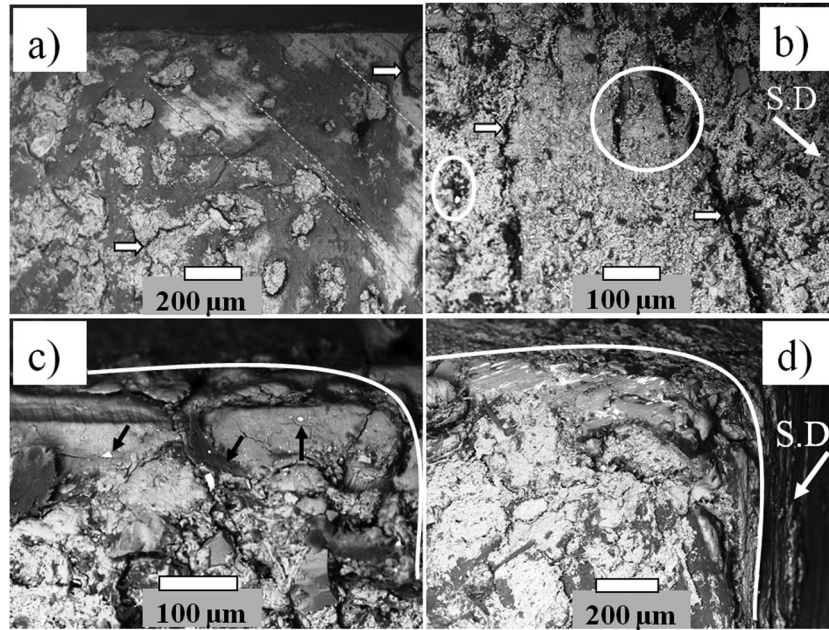


Figure 6. SEM images of wear scars of bone pins after pin-on-disc tests for: (a) ZrN-01, (b) ZrN-02, (c) ZrN-03 and (d) ZrN-04 (dash lines indicate scratch marks, SD – sliding direction).

transversal direction between 0.647 and 0.892 GPa and 19.8 ± 1.6 GPa, respectively [7]. ZrN coatings hardness and modulus values range from 15 to 34 GPa and 240 to 320 GPa, respectively [8]. As the number of sliding cycles increases, the transferred layer onto ZrN surface becomes brittle and is destroyed through fatigue; the resulting wear debris acts as a third body.

3.3. Electrochemical behaviour

3.3.1. Potentiodynamic polarization measurements

Figure 7(a) shows the potentiodynamic polarization curves (plots of potential vs. log current density) of the ZrN coatings exposed to the PS for 2 h. All specimens form an anodic domain (labelled '1'), which corresponds to the formation of a self-passivating (Zr, N,

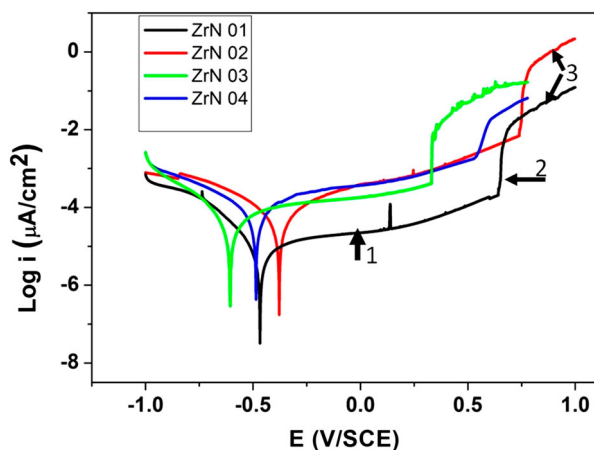


Figure 7. Potentiodynamic polarization curves of ZrN coatings in physiological liquid solution.

O) film after being soaked for a period of time. The size of this domain is indicative of the durability of the self-passivating film and the corrosion resistance of each coating. For example, the anodic plateau remained flat from -0.4 up to $+0.65$ V in the case of ZrN-01. The rupture of the passive film occurs in the trans-passivation domain (labelled '2'). In this domain, the current density abruptly increases as the materials start to corrode (labelled '3'). The polarization resistance (R_p) of Tafel curves, which takes into consideration both the corrosion current density and the slopes of polarization curves, can provide a quantitative ranking of coatings resistance to corrosion and can be estimated using the well-known Stern–Geary equation.

Table 3 summarizes the relevant values from the Tafel plots. From these results, it is clear that ZrN-01 is significantly more corrosion-resistant (high value of R_p and low value of I_{corr}) when compared to the other three coatings.

3.3.2. EIS analysis

Each sample's impedance was measured after 1 h of immersion in the PS. Nyquist plots of the real and imaginary parts of the impedance are shown in Figure 8(a). The plots show that all tested samples are characterized by two capacitive loops in the high and medium

Table 3. Potentiodynamic polarization test results.

Samples	E_{corr} (mV)	I_{corr} ($\mu A/cm^2$)	a (mV/dec)	c (mV/dec)	R_p (Ω)	$\chi^2 \cdot 10^{-4}$
ZrN-01	-466.76	0.007	174.8	544.3	$8.04 \cdot 10^6$	15.99
ZrN-02	-376.53	0.042	297.7	274.5	$1.47 \cdot 10^6$	13.42
ZrN-03	-612.98	0.037	512.7	220.2	$1.80 \cdot 10^6$	14.44
ZrN-04	-486.20	0.044	202.4	219.9	$1.04 \cdot 10^6$	9.10

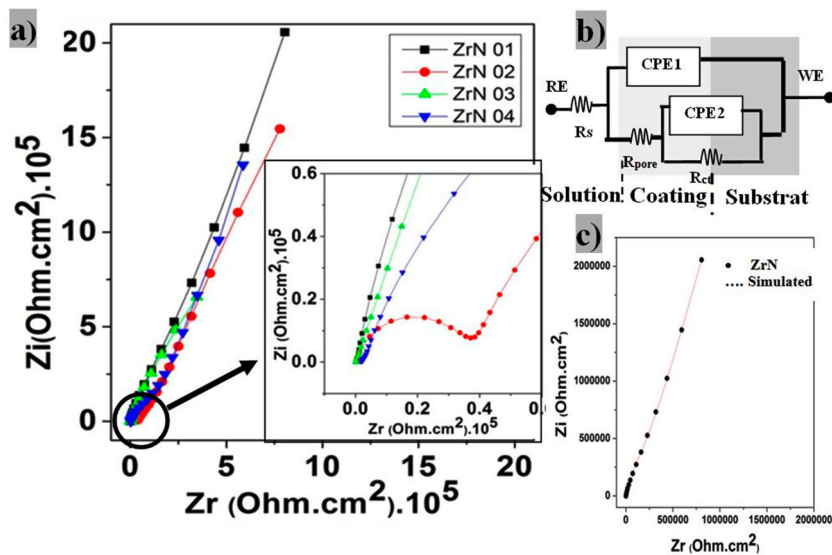


Figure 8. Electrochemical test results (EIS): (a) Nyquist plots of tested specimen in physiological liquid solution, (b) equivalent circuit model and (c) example of fitting curve (WE = working electrode and RE = reference electrode).

frequency range. By comparing the diameters of the loops in the Nyquist plot, the sample's corrosion properties can be easily evaluated. Larger diameter loops equate to higher corrosion resistance. This analysis supports the previous conclusion: ZrN-01 has the best corrosion resistance of the four tested samples. An equivalent circuit (EC) model, which is shown in Figure 8(b), was selected for quantitative studies. This EC is extensively used in similar situations [9]. Figure 8(c) shows good agreement between the experimental and modelled values. The fitted line was plotted as a dashed line. The fitting errors for each configuration in this study were less than 9%.

In the selected equivalent circuit, R_s represents the electrolyte resistance. The Constant Phase Elements, CPE1 and CPE2, are the double layer capacitance and the coating capacitance respectively. R_{ct} is the charge transfer resistance and R_{pore} represents the pore resistance (and other fails of coating).

The overall corrosion resistance of the coating can be estimated by the sum of all resistances of the circuit model in accordance with the method described by Subramanian et al. [9]. Therefore, using the quantitative data given in Table 4, the ZrN-01 coating presents the highest resistance against corrosion in a physiological liquid solution of the four tested samples, as expected.

3.4. Biocompatibility

The bactericidal properties of ZrN coatings were assessed using *S. aureus* bacteria. This bacterium is the main pathogenic microbe responsible for implant associated infections [10]. Two kinds of samples are used: ZrN coating on silicon wafer and an uncoated Ti-6Al-4V substrate. Ti-6Al-4V is one of the most

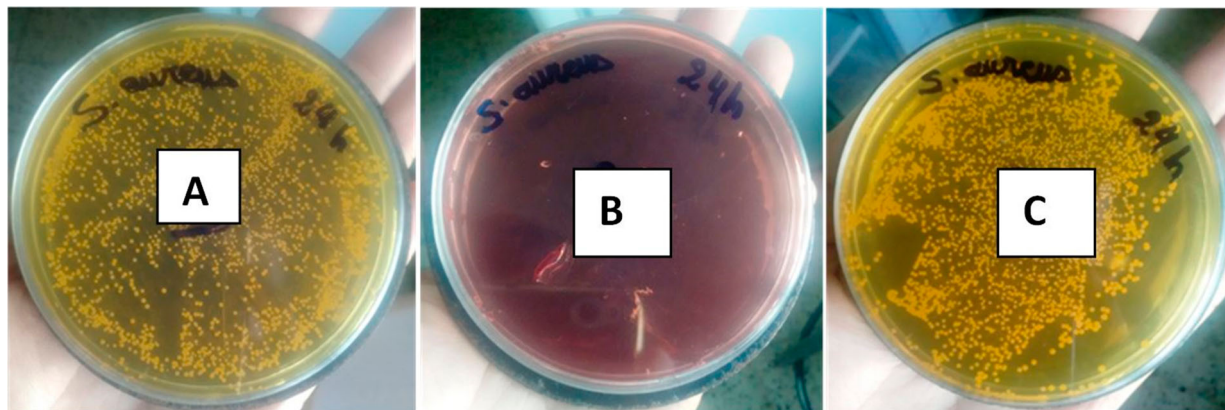
studied biomaterials, and its biocompatibility in various environments has been well documented. This sample acts as a 'reference' material for the comparison of the biocidal properties of ZrN coatings.

Following sterilization in an autoclave, three samples (bare Ti-6Al-4V, ZrN-01/Si and ZrN-03/Si) were exposed to bacteria suspended in a buffered glucose broth and incubated at 37°C for 24 h. To quantitatively assess the antibacterial effect of the three samples, the plate count method, in association with the dilution method, is used [11]. Specimen A is the liquid solution collected from the suspension which contained the metallic Ti-6Al-4V substrate. Specimen B and specimen C are the liquid solutions which previously contained the ZrN-01/Si and ZrN-03/Si samples, respectively. Twenty microlitres of each specimen are spread over a sterilized Petri dish coated with mannitol salt agar, also known as Chapman medium. The coated Petri dishes were incubated at 37°C for 24 h. Quantitative test results (counts) are given in term of colony forming units per millilitre (CFU/ml).

After incubation, the *S. aureus* colonies are clearly visible (Figure 9) and were easily counted. The results indicate that *S. aureus* bacteria were completely killed in specimen B, while the surviving bacterial colonies are 3.2×10^5 (CFU/ml) for specimen A and 5.2×10^5 (CFU/ml) for specimen C. Specimen B and Specimen C have approximately similar surface parameters (i.e. roughness, morphology, etc.) but there is a noticeable difference in the oxygen content of the samples. So, surface morphology effects cannot explain the difference in the biocidal effect of these two specimens (B and C). In addition, it has been reported that a surface roughness $<0.2 \mu\text{m}$ may not affect bacterial adhesion because most bacteria are $>0.2 \mu\text{m}$ [12].

Table 4. EIS data obtained by equivalent circuit simulation.

Samples	R_s ($\Omega \text{ cm}^2$)	CPE1 ($\mu\text{F}/\text{cm}^2$)	a_1	R_{pore} ($\Omega \text{ cm}^2$)	CPE2 ($\mu\text{F}/\text{cm}^2$)	a_2	R_{ct} ($\Omega \text{ cm}^2$)
ZrN-01	163.1	$2.68 \cdot 10^{-6}$	0.91	$675 \cdot 10^3$	$2.08 \cdot 10^{-6}$	0.73	$205 \cdot 10^6$
ZrN-02	155.3	$16.23 \cdot 10^{-9}$	0.86	$35.7 \cdot 10^3$	$3.92 \cdot 10^{-6}$	0.74	$31.08 \cdot 10^6$
ZrN-03	304.3	$5.11 \cdot 10^{-6}$	0.96	$4.6 \cdot 10^3$	$5.85 \cdot 10^{-6}$	0.72	$13.39 \cdot 10^6$
ZrN-04	365.2	$2.46 \cdot 10^{-6}$	0.78	$362.9 \cdot 10^3$	$3.92 \cdot 10^{-6}$	0.81	$48.85 \cdot 10^6$

**Figure 9.** *Staphylococcus aureus* colonies grown in sterilized Petri dishes of: (a) specimen A, (b) specimen B and (c) specimen C.

However, chemical interactions between microorganisms and biomaterial surfaces can have a significant effect on bioadhesion and the proliferation process [13,14]. Within liquids, oxide surfaces can be positively charged and bacteria typically are negatively charged [15]. Consequently, bioadhesion of bacteria is significantly enhanced in the presence of surface oxides. This result is in accordance with conclusions reported by Vadillo-Rodriguez et al. [14]. The presence of oxygen, and the development of surface oxides, can also affect the release of Zr ions. Zirconium ions interact with the bacterial cell wall, which may disturb its permeability and/or may inhibit bacterial metabolism [13].

Therefore, oxygen content appears to be a key limiting factor in the biocompatibility of ZrN surfaces. Oxygen content leads to higher bioadhesion aptitude and more colonies (proliferation). As a result, there is a higher probability of *S. aureus* survival and increased chances for infection.

4. Conclusion

From the above-mentioned results, it can be concluded that:

- (i) All coatings exhibited good wear properties against bone counterparts. The primary wear mechanisms were abrasive wear and bone adhesion for the ZrN-coated discs and third-body abrasive wear for the bone counterparts.
- (ii) All coatings exhibited good corrosion resistance. However, the oxygen content seems to play a more significant role here than in the wear tests. The sample with the lowest oxygen content,

ZrN-01 ($\text{O}_2 = 1.5 \text{ wt. } \%$), showed the highest corrosion resistance in both potentiodynamic testing and EIS.

- (iii) Bioactivity of an uncoated Ti-6Al-4V disc substrate and the ZrN-03 ($\text{O}_2 = 4.9 \text{ Wt. } \%$) coating shows a poor antimicrobial effect against the *S. aureus* bacteria. A strong biocide effect was observed with the surface of ZrN-01 ($\text{O}_2 = 1.5 \text{ Wt. } \%$) coating; no microbes were observed after 24 h of contact time.

Acknowledgements

The authors are thankful to Mr. Denis LAGADRILLERE for SEM/EDS analysis.

Disclosure statement

No potential conflict of interest was reported by the authors.

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