Letter







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Consume or Conserve: Micro-Roughness of Titanium Implants towards Fabrication of Dual Micro-Nano Topography

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<u>Abstract</u>

Combining the micro-roughness of current titanium implants, which promotes accelerated bone integration, with nano-topography for enhanced bioactivity/drug-release may be an ideal solution to address therapeutic challenges inside the bone micro-environment. We hereby present a single-step electrochemical anodization using conditioned electrolyte to enable fabrication of aligned titania nanopores with preserved micro-scale features of the underlying titanium implant. Applicability towards the fabrication of mechanically robust and clinically translatable next-generation of orthopaedic/dental implants with dual-topography including 'gold-standard' micro-roughness and superimposed 'bioactive' nanotopography.

Electrochemical anodization (EA) of titanium resulting in self-ordering of titania nanostructures (TNSs), such as nanotubes and nanopores, has gained considerable interest with applications including bioactive and drug-eluting implants.¹⁻² It is accepted that electrolyte preparation via repeated anodization (or ageing) results in improved ordering and adherence of TNSs, especially in the case of organic electrolytes such as ethylene glycol.³⁻⁶ It is worth noting that ageing optimizes the fluoride and water composition (especially in a sealed electrochemical cell) of the electrolyte, which are crucial parameters for the formation of TNSs.⁷ These parameters determine oxide growth/dissolution rates, O₂ evolution, barrier layer thickness and the compressive stress at the titanium-oxide interface, which are inter-dependent with each other and greatly influence the yield of the pores/tubes.⁷

Thus far, very few studies have focussed on understanding the influence of ageing on EA of Ti, and hence there is a significant research gap and limited understanding of EA in this domain. Most studies utilize an EA setup whereby the moisture uptake by the hygroscopic electrolyte is a major consideration, as it is constantly increasing water content and hence conductivity.³⁻⁶ Alternatively, more recently we have presented an extensive investigation detailing the fabrication of TNS via electrolyte ageing in a sealed electrochemical setup.⁷ With this approach, water content continuously reduces and so does the conductivity.³ It was established that ageing the electrolyte enables improved ordering and the formation of stable nanostructures on the complex geometry of curved surfaces, representing a wider implant market and hence easy translation. However, to advance our knowledge with respect to electrolyte ageing and allow successful implant nano-engineering, it is crucial to understand what happens at the metal-oxide interface, how the substrate topography influences the EA and how the mechanical properties differ for TNS from fresh- and aged-electrolyte anodizations.

In the case of biomedical applications of nano-engineered Ti, such as bone/dental implants, the preservation of the underlying Ti implant features/roughness may be particularly important for enabling initial interlocking/stability from the micro-features, and then superimposing nanotubes/pores which may enable enhanced bioactivity or facilitate drug release.⁸ These hybrid micro-nano topographies have been prepared in the past using more complex lithography, laser-patterning or 3D printing, which have been combined with EA to create nanotubes/pores.⁹⁻¹¹ Bypassing the conventional surface smoothening steps and complex micro-patterning strategies, we hereby showcase a single-step EA on micro-machined Ti implants which yields optimized micro-nano topographies. Titania nanopores

are fabricated instead of conventional nanotubes to augment the stability of the anodic film, crucial for implant applications. The optimizations are performed using two electrolyte systems: pre-utilized (aged) and un-used (fresh), while other parameters were controlled, to also enable better understanding of the role of electrolyte conditioning in preserving the underlying implant micro-scale features. Additionally, the mechanical testing of these nanostructures obtained from varied electrolyte anodizations was also compared. The fabrication of stable nanotopography with preserved micro-scale features of commercial orthopaedic/dental implants could allow for multiple bioactive and therapeutic functionalities, while enabling easy translation into current implant market.

Micro-machined Ti ('Micro-Ti') discs (6.24mm diameter, 2mm height, commercially pure grade 4 Titanium) were supplied by AstraTech Dental (Sweden). Flat titanium foil (99.5% purity, 0.2mm thick, 1 cm²) was purchased from Nilaco (Japan). Ethylene glycol and ammonium fluoride (NH₄F) were purchased from Sigma-Aldrich (Sydney, Australia). Surface topography of Ti implant substrates and the fabricated TNSs were investigated using a scanning electron microscope (SEM, Zeiss Sigma FESEM). Titania nanopores (TNPs) were fabricated via a single-step EA in a two-electrode electrochemical cell at 60-100V for various times at room temperature using a DC power source (Keithley 2460) with the current precisely monitored. The Micro-Ti discs with micro-groove features were used as an anode and flat Ti foil served as the counter electrode. In this study, we performed anodizations in a moisture-controlled system with an ethylene glycol electrolyte (with 0.3 wt% NH₄F and 1 vol% deionized water). Two different electrolytes were used for this study: fresh and aged. "Fresh electrolyte" represents the newly prepared electrolyte without any previous use, whereas "aged electrolyte" represents a previously used electrolyte, which was repetitively anodized for 10h using dummy Ti, as previously described.⁷ To quantify the mechanical properties of the fabricated nanoporous structures, elastic modulus and hardness measurements were performed by nanoindentation testing, using a Berkovich 3-sided pyramidal tip with a maximum force of $10,000 \mu N$.

Figure 1 shows top-view SEM images of bare Micro-Ti, as well as anodized Micro-Ti with either fresh or aged electrolyte. As received Micro-Ti (Figure 1a,b) shows machined micro-scale lines and features, characteristic of the micro-roughness of commercial implants. The average width of the 'sinusoidal' micro-features as determined from SEM images is approximately $36 \mu m$. Upon anodization, the micro-features of the implant are better preserved with aged EA, with the 'crest-and-trough'

like features of Micro-Ti being markedly diminished with the fresh EA. Both systems resulted in the formation of nanopores along the machined lines of the Micro-Ti. The average diameter of nanopores on titanium for fresh and aged EA is 28 nm and 30 nm, respectively. The formation of pores instead of tubes can be attributed to: reduced water content, less voltage/time, lack of surface smoothening and the EA performed in a controlled environment.¹² Interestingly for aged EA, the nanopores were superimposed on the underlying micro-groove like architecture, thereby resulting in a dual micro- and nano-scale surface (Figure 1g-i). The micro-scale groves of the 'asreceived' machined Ti (Figure 1a,b) served as the template, allowing self-ordering of the nanopores along these lines. Clearly, for the aged system the nanopores can be seen aligned (Figure 1i) in the direction of the machining imprints (micro-scale lines, as seen in Figure 1b). On the contrary, for the fresh EA system, both of these features were diminished, with reduced alignment of the nanopores (Figure 1f) along the micro-machining lines of the Micro-Ti.

Interestingly, 1st anodized nanoporous structures have been reported previously, however have not been applied towards specific applications (especially orthopaedic/dental implants).¹³ Conventionally, these 1st anodized anodic films are removed to perform a 2nd EA on a template surface. A few studies have aimed at utilizing the dual micro-nano architecture to display augmented cellular functions, but only using multiple technologies combined with EA.⁹⁻¹¹ The ability to achieve such features via a single-step EA can enable easy and cost-effective clinical translation. Furthermore, more recently we have demonstrated that aligned nanopores can enable selective bioactivity, whereby proliferation and adhesion of osteoblasts and fibroblasts is enhanced, and macrophage functionality is reduced.¹⁴ Additionally, osteoblasts and fibroblasts aligned in the direction of the nanopore arrangement, clearly suggesting strong mechanical stimulation from these nanostructures.¹⁴ It is also noteworthy that a variety of cells are able to respond to mechanical and topographical cues, and hence changing the micro- and nano-dimensions (simply by varying anodization conditions) can further aid in achieving tailored cellular and therapeutic responses.^{2,15}

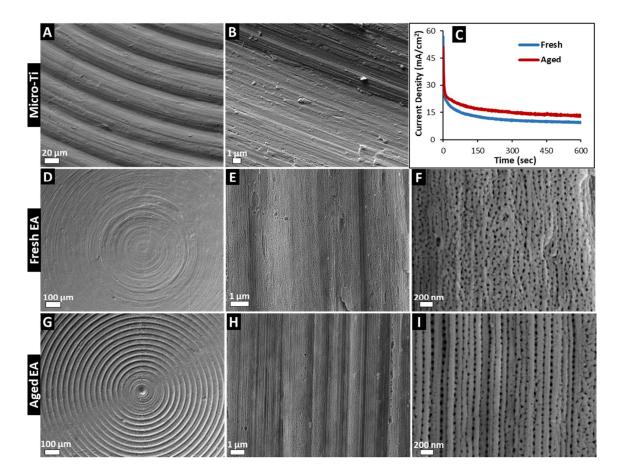


Figure 1. Electrochemical anodization (EA) of micro-machined Ti implants (Micro-Ti). Top-view SEM images: (a-b) bare Micro-Ti, (d-f) Micro-Ti anodized with fresh electrolyte, and (g-i) anodized in aged electrolyte. (c) Current density vs time plots comparing EA using fresh and aged electrolytes. EA performed at 60V 10m.

It is well known that nanotubes/pores grow perpendicular to the surface of the substrate, in the direction of the electric field. If the surface contains inconsistencies in the form of artefacts or impurities, the high concentration electric field at these points may result in very high local current densities, yielding defects in the anodic film.^{7,16} Besides, the electric field intensity is expected to be higher at the topographical minima of the substrate, i.e. minima of the 'crest and trough'-like architecture. The implant substrate used in this study not only has micro-scale features (crests and troughs), but also has a surface which is entirely decorated with a unidirectional machined lining (Figure 1a-b). The high intensity of the electric field at these points ('weak spots') further enhances the oxidation-dissolution rates, which is more prominent for the fresh EA system ('consumption').¹⁷ The growth phase is also

characterized by the continuous dissolution and rebuilding of the compact barrier layer (BL) at the pore bottom-electrolyte interface; and for a high electric field concentration (fresh electrolyte); high growth rates can effectively lead to consumption of substrate micro-features.¹⁸⁻¹⁹ This can explain the 'blunting' of the micro-features of the machined titanium substrate, yielding an almost flat surface, for the fresh EA system (Figure 1d-f).

In contrast to other electrolyte ageing approaches where the conductivity increases (due to higher TiF_6^{2-} and water content in the electrolyte); in a moisture-controlled system the conductivity continuously reduces with each EA, which is mainly attributed to the consumption of water.⁷ We have also observed an increase in O₂ bubbles at the oxide-electrolyte interface for fresh EA, as compared to a sparse bubble production for the aged EA. We have also shown previously that the increased time required to reach an EA equilibrium stage, as well as the reduced rates of formation and dissolution of the oxide, contribute towards the generation of mechanically stable nanostructures, especially on a complex substrate with many 'weak spots'.⁷

As presented in Figure 1g-i, the aged EA enables generation of a hybrid micro-nano architecture whereby an anodic film with nanopores is superimposed on the micro-patterned Ti implant. Furthermore, as observed in the SEM images (Figure 1f,i), it is clear that the underlying micro-machining enables pore alignment, whereby the pores/tubes aligned in the direction of the grooves on the mechanically polished Ti. However, it is interesting to note that the alignment of pores along the underlining machined surface is better for aged-EA, as compared to fresh-EA. This again can be attributed to increased preservation of these micro-machining lines (serving like a template) in the aged EA (water and fluoride constraints impeding both oxidation and dissolution reaction rates). It is noteworthy that the dimples/template features of the Ti substrate acts as initiation sites for tube/pore growth, and hence can result in specific arrangements.

Presence of active fluoride ions in the electrolyte can significantly influence the EA process.^{1,7} Fluoride accumulation at the nanotube/nanopore bottoms in the form of soluble species (Ti-O-F, Ti-F complexes) also forms an additional F-rich layer between the BL and the substrate.²⁰ In addition, for a fluoride-rich fresh electrolyte, the fluoride attack can be more prominent at the defect sites (in our study: presence of micro-machining lines, and micro-architecture: crest and troughs). In fact, it has been

reported that water content in fluoride-containing electrolytes can caused increased expansion [Pilling–Bedworth ratio (PBR)] that can lead to stress and volume difference between anodic layer and substrate.²¹ For an aged system, with reduced water (and fluoride content) reduced expansion rates can be assumed, which in our investigation led to improved preservation of both the micro-machining lines (better nanopore alignment on the micro-templated Ti) and the 'crest and trough' like micro-architecture (dual micro-nanoporous).

To further investigate why the micro-features were poorly or well preserved in specific electrolyte systems, we observed the early time points in the EA, using 60V and 100V for 60sec. The top view SEM images and corresponding current densitytime plots are presented in Figure 2. For both 60 and 100V, in the first 15 sec, there is no significant difference between the current density values for fresh and aged systems, however after 15 sec, clearly J-aged > J-fresh (Figure 2e-h). This co-relates to the delayed time to reach equilibrium (established through the barrier oxide layer at the nanopore bases) and thicker barrier layers (for aged EA).⁷ It has also been recognized that the fluoride-rich layer at the metal-oxide interface is attributed to faster inward fluoride ion migration under high electric fields.²²⁻²³ Clearly for a fresh system with an abundance of fluoride ions, rapid fluoride migration to the metal-oxide interface occurs, which is followed by chemical dissolution of these fluoride-rich layers in water, resulting in a tubular morphology (consumption of surface microfeatures over time).²⁴ On the contrary, in a fluoride and water deficient aged system. the formation/solubility of this layer is impeded, thereby preserving the underlying substrate features and continuing with the EA-assisted growth of nanotubes/pores. The perpendicular growth of tubes/pores thereby results in surface defects (due to field concentration effects), as shown in Figure 2j,l, for the aged-EA. This finding is similar to the observed cracks/voids on the anodic film on curved surfaces of Ti wire, as reported previously.⁷ However, it is worth noting that over time, these surface imperfections are reduced with continued EA, as shown for 60V-10m EA in the agedsystem (Figure 1h-i).

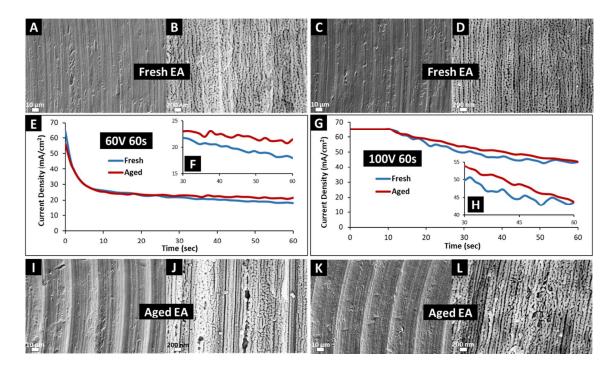


Figure 2. Comparison of electrochemical anodization (EA) of micro-machined Ti in fresh and aged electrolyte systems at 60/100V for 60 sec. Top-view SEM images: (a-b) Fresh EA 60V, (c-d) Fresh EA 100V, (i-j) Aged EA 60V, and (k-l) Aged EA 100V. Current density vs time plots: (e-f) 60V 60sec, and (g-h) 100V 60sec.

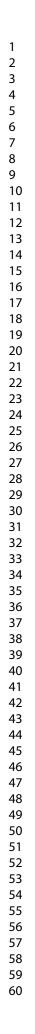
To simplify, we have schematically represented the EA procedure (Figure 3) for fresh and aged electrolyte systems to correlate with the SEM images shown in Figure 1-2. It is well established that the electrochemical equilibrium established as per the following chemical reactions tunes the balance between TiO_2 formation and its dissolution, leading to the self-ordering of nanotubes/pores.^{1,10}

Oxide Formation:	$Ti + 2H_2O \rightarrow TiO_2 + 4H^+ + 4e^-$	(1)
Oxide Dissolution:	$TiO_2 + 4H^+ + 6F^- \rightarrow TiF_6^{-2-} + 2H_2O$	(2)

Following the Le Chatelier's equilibrium principle, for an aged system, clearly the scarcity of water in the electrolyte reduces the oxide formation rate (1), while the presence of TiF_6^{2-} reduces the oxide dissolution rate (2), both leading to a reduced rate of nanotube/nanopore growth. For fresh EA, representing a rapid formation and dissolution system, an increased build-up of compressive stress at the early EA phase occurs when metal converts to oxide (volume expansion: Pilling-Bedworth ratio or PBR).²⁴⁻²⁵ Over time, this results in the consumption of the micro-features of the Ti, as shown in Figure 3a. On the

contrary, in an aged system, whereby the dissolution is severely impeded, the compressive stress may be 'frozen' in the compact oxide layer, as reported elsewhere.²⁴⁻²⁵ Although with numerous parameters influencing the metal-oxide interface, it is difficult to elucidate the exact mechanism behind the varied EA behavior observed for fresh and aged systems, clearly reduced water and fluoride content for aged-EA induces surface defects, increased time to reach equilibrium current value, and reduced etching of grain boundaries, all leading to well preserved underlying substrate features with superimposed anodic nanopore film.²⁶⁻²⁷ Furthermore, it is established that EA on a nano-templated substrate (obtained after removal of the anodic film) results in improved ordering, however this approach may completely remove the underlying implant features. Our strategy (single step EA with conditioned electrolyte) may be beneficial for applications such as the ability to preserve the 'gold standard' micro-features of dental implant surfaces, which have been shown to improve early bone formation and osseointegration.⁸

In the current study, for EA in a shielded electrochemical setup (avoiding moisture uptake by the hygroscopic electrolyte: throughout ageing using dummy titanium and anodization of target substrate), the pH of the electrolyte increased from 6.65 for fresh electrolyte to almost 7.90 (averaged values) for a 10 hr aged electrolyte. It is interesting that other electrolyte ageing studies report a reduced pH, however, that may be attributed to a continued water absorption over time (which also leads to an increased conductivity).⁶⁻⁷ For our study, this effect can easily be explained by continued consumption of water which leads to reduced titanium hydrolysis (equation 1), lesser H^+ production and hence a relatively higher pH value.²⁸ Various investigations have also stressed the importance of pH on anodization yield, however, pH at the nanotube/pore tips and the bottoms is controlled by the bulk electrolyte and self-acidification process, respectively.²⁹ Here, the self-acidification at the tube-bottoms is in turn effected by the chemistry of the electrolyte: notably in aged electrolyte this is related to the water and fluoride constraints.³⁰ Briefly, it is established that for a basic electrolyte pH, both the oxide formation (equation 1) and the chemical dissolution (equation 2) rates are reduced.²⁹ Therefore, a decreased rate of electrochemical etching at the tube bottom for aged electrolyte (pH 7.9) as compared to fresh electrolyte (pH 6.65) may also explain the 'conserving' of the substrate micro-features.



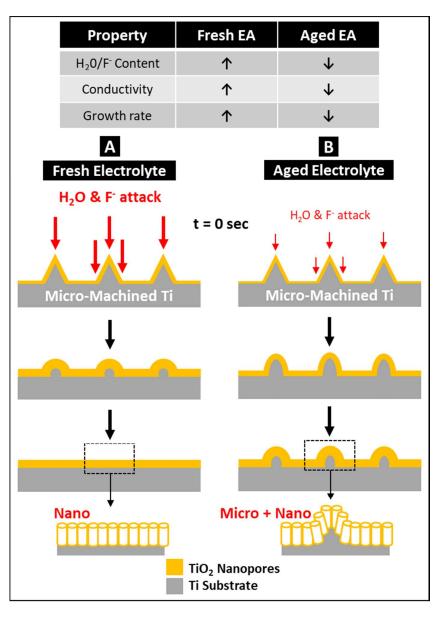


Figure 3. Schematic illustration of the varied electrochemical anodization (EA) reactions based on electrolyte age (fresh/unused and aged) on (a) 'consuming' [higher fluoride and water attack] and (b) 'conserving' [reduced fluoride and water attack] the underlying microroughness of a titanium substrate, towards generation of nanoporous and dual micronanoporous topographies, respectively. Table summarizes the difference in water/fluoride content between the fresh and aged electrolytes, as well as the corresponding EA properties.

It is worth noting that for conventional nanotubes, the gap between the individual tubes can make them prone to fracture and hence can be less mechanically robust.²⁷⁻³¹ On the contrary, nanopores are compactly interconnected with the surrounding pores enabling

sharing/distribution of external compression forces.^{27,32} To evaluate the mechanical stability of the as-fabricated dual micro-nano Ti implants, we investigated and compared the elastic modulus and hardness of the nanopores anodized with fresh and aged systems (Table 1). Nanoindentation testing revealed a dependence of the modulus on the displacement depth of the indenter, and clearly for both systems, the modulus increased with depth, which can be attributed to the presence of underlying substrate and densification.^{27,33} However, aged EA yielded nanopores with increased modulus and hardness values as compared to the corresponding depth for fresh EA. These observations are also in agreement with our previous study, whereby we obtained improved stability of anodic films when more time was taken for the current to reach a steady-state value (equilibrium).⁷ Interestingly, the dual micro-rough and nanoporous structures fabricated using aged EA demonstrated increased modulus and hardness values. This is also in accordance with a study by Wang et al., which showed that a laser pre-treated micro-rough surface significantly enhanced the adhesive strength between nanostructure layers and the underlying substrate, mainly due to the expanded interfacial area.¹⁰

Displacement	Elastic Modulus (GPa)		Hardness (GPa)	
(nm)	Aged	Fresh	Aged	Fresh
50	44.34±15.1	44.90±9.5	2.32±1.5	1.73±0.5
100	55.45±10.3	46.09±2.4	2.18±0.4	1.36±0.1
150	59.84±11.1	50.33±3.3	2.45±0.5	1.33±0.1
200	69.21±16.7*	57.38±3.8* [#]	2.63±0.7	1.36±0.1
250	73.05±12.1*	58.85±7.1* [#]	2.83±0.6	1.43±0.2
300	73.91±10.6*	61.32±7.4* [#]	2.93±0.5	1.48±0.3

Table 1 Comparing the mechanical properties of the nanoporous surface on the micromachined titanium implants, anodized (60V 10m) using aged and fresh electrolytes, using Nanoindentation testing. Measurements represent a mean value of atleast 3 indentation measurements performed on each sample (total 3 samples) [*: p<0.05 compared with values from 50 nm within the same group, [#]: p<0.05 compared with values from 100 nm within the same group].

While the results of this study demonstrated the ability to decide between 'consumption or conservation' of substrate micro-features, towards fabrication of dual micro-and nanoporous titanium for implantable applications, there are a few challenges with respect to determining the precise mechanism involved (with respect to role of electrolyte ageing, based on current established models).³⁴⁻³⁵ These briefly include: (a) surface micro-features (crest and trough) and micro-machining lines cause extreme irregularity with respect to the electric field distribution; (b) the monotonic current density decrease with time causes issues with estimating J_{max} or J_{min} (and corresponding BL), especially for a short duration (and less voltage) of anodization; and (c) the complex chemical balance of fluoride and water constraints with excess Ti ions (aged electrolyte). Apolinário *et al.* have described these factors in detail for a more conventional Ti EA with fresh electrolytes, and longer EA times (with higher voltages).³⁴⁻³⁵ We believe the abovementioned challenges must be met to correctly estimate and match the established EA models used to describe the growth kinetics dependent on various EA parameters (more importantly on Ti substrate roughness and conditioned electrolyte: both of which were at play in this study).

The results of this study provide evidence that for electrochemical anodization of microrough titanium implant, electrolyte ageing can enable preservation of the underlying implant micro-features (especially relevant to ensure initial implant interlocking/stability), thereby yielding a dual micro-nano surface. The findings compared the short-time anodization between fresh (unused) and aged electrolytes, to elucidate how electrolyte conditioning contributes towards the 'consumption or conservation' of the underlying micro-roughness. Bypassing the complicated multi-step procedures, this cost-effective single-step anodization technique with aged electrolyte yielded mechanically-stable aligned nanopores superimposed on the micro-scale features of the implant, enabling multi-functionality (micro: initial stability, and nano: bioactivity and drug release) and thereby ensuring easy translation into current implant market.

Conflict of interest

There are no conflicts to declare.

Acknowledgements

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Consume or Conserve: Micro-Roughness of Titanium Implants towards Fabrication of Dual Micro-Nano Topography

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