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Effect of pulse current on structure and adhesion of apatite electrochemically deposited onto titanium substrates

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Apatite films were deposited onto titanium (Ti) metal substrates by an electrodeposition method under a pulse current. Metastable calcium phosphate solution was used as the electrolyte. The ion concentration of the solution was 1.5 times that of human body fluid, but the solution did not contain magnesium ions at 36.5 °C. We used an average current density of 0.01 A/cm² and current-on time (T_{ON}) equal to current-off time (T_{OFF}) of 10 ms, 100 ms, 1 s, and 15 s. The adhesive strength between apatite and Ti substrates were relatively high at $T_{\text{ON}} = T_{\text{OFF}} = 10$ ms. It is considered that small calcium phosphate (C-P) crystals with low crystallinity were deposited on the Ti surface without reacting with other C-P crystals, H₂O, and HCO₃⁻ in the surrounding environment. This resulted in relaxation of the lattice mismatch and enhancement of the adhesive strength between the apatite crystals and Ti substrates.

I. INTRODUCTION

Titanium (Ti) metal and its alloys are widely used for orthopedic and dental applications because they exhibit high mechanical properties and biocompatibility.¹⁻³ Ti metal has a passive layer of TiO₂ on its surface, which is responsible for its chemical stability and therefore its biocompatible characteristics. This oxide layer naturally has a thickness of a few nanometers, but the thickness can be increased to a few micrometers by chemical and thermal treatments.⁴ Although Ti metal is biocompatible, it is not bioactive and hence it cannot directly bond to living bones. To improve its bone-bonding property, apatite coating is applied on Ti implants.⁵ Several coating methods, such as plasma spraying,⁶ biomimetic precipitation,^{7,8} and electrodeposition,⁹⁻¹¹ have been successfully used to deposit an apatite layer on Ti or titanium alloy implants. Each of these methods has some advantages as well as drawbacks. Among these coating tech-

niques, only the plasma spraying of apatite is clinically used. However, the high temperature of the plasma flame may adversely affect the implant structure and the homogeneity of the coating, resulting in delamination of the coating.¹²⁻¹⁴

The electrodeposition method is an alternative process that uses aqueous solutions at low temperatures; this method hardly affects the implant structure and can be applied to complex shapes.¹⁵ In addition, the apatite layer can be rapidly formed on Ti substrates by electrodeposition in a metastable calcium phosphate solution whose ion concentration is 1.5 times that of a normal simulated body fluid (SBF), and does not contain MgCl₂·6H₂O.¹⁶ However, this method increases the pH at the interface between Ti and the electrolyte due to electron incorporation to form OH⁻ ions and H₂ through water reduction.¹⁷ The H₂ gas evolution at the interface leads to a heterogeneous coating.¹⁵ Furthermore, the adhesive strength between apatite and the substrate is often low and the total amount of apatite crystals deposited on the Ti substrate is much smaller than that theoretically calculated from the total electric charge.¹⁸ To solve this problem, new approaches are needed in the electrodeposition

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method. Our previous studies revealed that electrodeposition under a long pulse (>5 s) was more effective for crystal growth and apatite formation than that under direct current.¹⁹ The studies also revealed that when the Ti metals were previously etched in 50% or 75% sulfuric acid (H₂SO₄) followed by electrodeposition and then heat treatment, a dense and uniform apatite layer with good adhesive properties was formed onto the Ti substrates.²⁰ However, pyrophosphate (P₂O₇⁴⁻) compound, an inhibitor of mineralization,²¹ was formed in apatite crystals during heat treatment.

The purpose of the present work is to coat an apatite layer on Ti substrates with good adhesion in metastable calcium phosphate solution by using various pulse widths without subsequent heat treatment and to investigate the effect of pulse width on the apatite coatings and adhesive strength.

II. EXPERIMENTAL

Rectangular specimens of Ti substrates (purity: 99.5%, The Nilaco Corporation, Tokyo, Japan) with a volume of 10 × 10 × 1 mm³ were abraded with a #400 diamond plate and then washed with acetone, ethanol, and ultrapure water in an ultrasonic cleaner.

Before the electrodeposition under pulse current (PE), the Ti substrates were etched in 30 cm³ of H₂SO₄ with a concentration of 75%, which was prepared by diluting 97% H₂SO₄ (Nacalai Tesque Inc., Kyoto, Japan) with ultrapure water at 60 °C for 30 min. After etching, the specimens were washed with ultrapure water.

PE was performed in a two-electrode cell configuration. The Ti specimens were used as the cathode, and a platinum plate (HX-C2, Hokuto Denko Corp., Tokyo, Japan) was used as the counter electrode. At a current-on time (T_{ON}) of 15 s, the pulse current was applied by a function generator (Agilent 33120A, Agilent Technologies Inc., Tokyo, Japan). At a T_{ON} of 10 ms, 100 ms, and 1 s, the pulse current was applied by a controlled current pulse generator (HC-110, Hokuto Denko Corp., Tokyo, Japan). The average current density was 0.01 A/cm², and T_{ON} was equal to the current-off time (T_{OFF}) irrespective of the pulse width. The electrolyte was 60 cm³ of 1.5SBF (simulated body fluid) without magnesium Mg²⁺ ions at 36.5 °C. In this study, to enhance the growth of apatite, the ion concentration of the solution was increased up to 1.5 times that of normal SBF,²² but Mg²⁺ ions, an inhibitor of hydroxyapatite precipitation and growth,²³ were eliminated in the electrolyte. Table I shows the amounts of reagents for the preparation of 1.5SBF without Mg²⁺ ions. This electrolyte was prepared by dissolving reagent grade chemicals of NaCl, NaHCO₃, KCl, K₂HPO₄·3H₂O, CaCl₂, and Na₂SO₄ (Nacalai Tesque Inc., Kyoto, Japan) in ultrapure water. The pH of the electrolyte was adjusted to 7.40 with tris(hydroxy-

TABLE I. Amounts of reagents for preparation of 1.5SBF without Mg²⁺ ions.

	Reagent	Amount/g	Purity/%
1	NaCl	11.994	99.5
2	NaHCO ₃	0.525	99.5
3	KCl	0.336	99.5
4	K ₂ HPO ₄ ·3H ₂ O	0.342	99.0
5	1.0M-HCl	48 cm ³	...
6	CaCl ₂	0.417	95.0
7	Na ₂ SO ₄	0.107	99.0
8	(CH ₂ OH) ₃ CNH ₂	9.086	99.0
9	1.0M-HCl	~15 cm ³	...

methyl)aminomethane [(CH₂OH)₃CNH₂] and 1.0 mol/dm³-HCl aqueous solution (Nacalai Tesque Inc., Kyoto, Japan) at 36.5 °C. The deposition times were 90 min, because our previous study¹⁶ demonstrated that uniform apatite deposition was achieved by electrodeposition over 90 min. After the electrodeposition, the specimens were gently washed with ultrapure water and then dried at 40 °C for 24 h.

The surface structure of the specimens was investigated by field-emission scanning electron microscopy (FE-SEM; S-4500, Hitachi Ltd., Tokyo, Japan), thin-film x-ray diffraction (TF-XRD; Rigaku RINT-2500, Tokyo, Japan), and Fourier transform infrared reflection spectroscopy (FTIR; Nicolet Magna 860, Thermo Electron K.K., Kanagawa, Japan). The thickness of the products deposited on the Ti substrates by PE was measured by FE-SEM. The surface roughness (arithmetic average roughness: R_a) of the substrates was measured by atomic force microscopy (AFM; SPA300, Seiko Instruments Inc., Tokyo, Japan).

The adhesion between apatite and the substrates was evaluated by a peeling test using Scotch tape (810, Sumitomo 3M Ltd., Tokyo, Japan) and by a scratch test using a surface property tester (Type 14DR, Shinto Scientific Corp. Ltd., Tokyo, Japan) equipped with a sapphire needle (diameter of 50 μm). In the scratch test, constant loads of 50, 100, and 200 g were applied to the scratch needle.

III. RESULTS AND DISCUSSION

Figures 1 and 2 show the FE-SEM photographs and the TF-XRD patterns of the surface of both the unetched Ti substrates and those etched in 75% H₂SO₄, respectively. We confirmed that microstructural roughness was formed by the acid etching. The R_a of the unetched substrate was 0.116 μm, while that of the etched substrate was 0.186 μm. The reactions taking place during the etching of the Ti substrate in 75% H₂SO₄ are shown below.²⁴

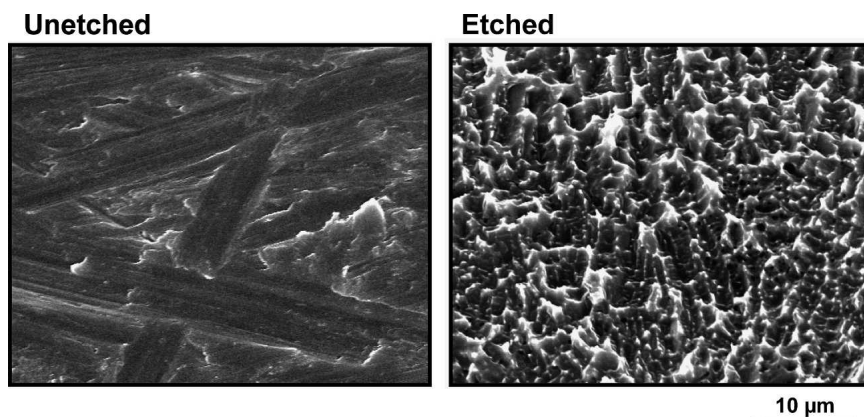


FIG. 1. FE-SEM photographs of surfaces of Ti substrates unetched and etched in 75% H₂SO₄.

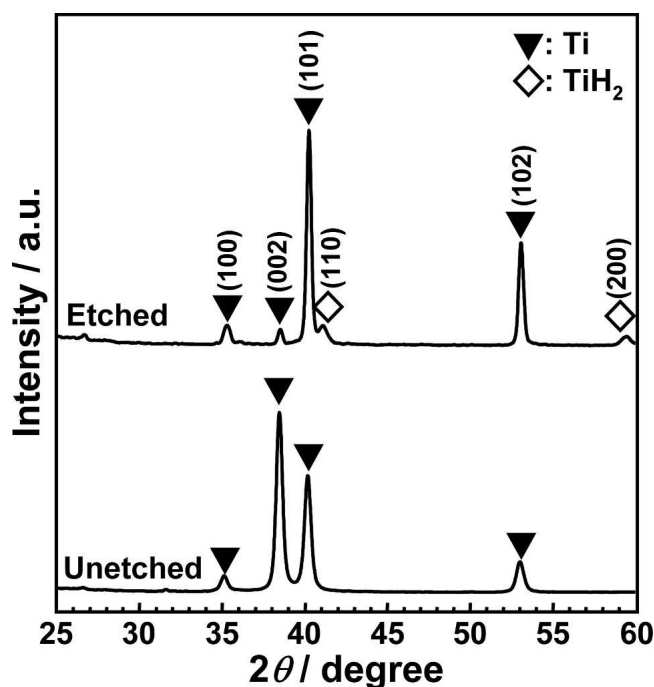
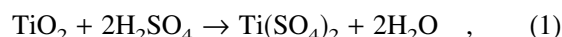


FIG. 2. TF-XRD patterns of surfaces of Ti substrates unetched and etched in 75% H₂SO₄.



According to the above chemical reaction (3), TiH₂ will be formed on the Ti substrates by the acid treatment. Indeed, we can confirm XRD peaks of TiH₂ around 40.9° and 59.6° in 2θ, as shown in Fig. 2 and our previous study.²⁰

Figure 3 shows the TF-XRD patterns of the unetched and etched Ti substrates subjected to PE at various T_{ON} . For all specimens, the peaks assigned to apatite were observed at around 26°, 28°, and 32° in 2θ.²⁵ Furthermore, the diffraction peak due to (202) of apatite was observed at around 36° in 2θ for the specimens at $T_{\text{ON}} = 15$ s. The existence of the diffraction peaks of apatite means that apatite was successfully deposited on the Ti metal surface by PE. The diffraction peaks of apatite for the specimens at T_{ON} above 1 s were stronger than those at T_{ON} below 100 ms. Figure 4 shows the total intensities of the diffraction peaks of apatite at various T_{ON} . The intensity of apatite tended to decrease with decreasing

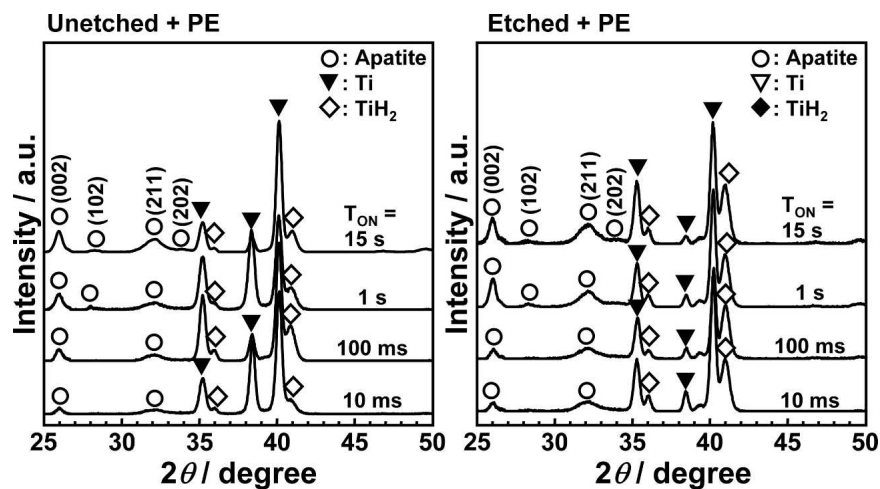


FIG. 3. TF-XRD patterns of surfaces of unetched and etched Ti substrates subjected to electrodeposition at various T_{ON} .

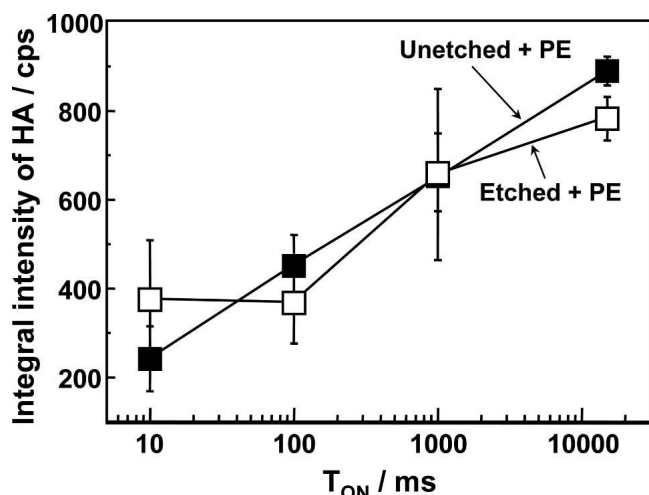


FIG. 4. Thickness of products deposited on Ti substrates as a function of T_{ON} .

T_{ON} . The decrease in intensity indicates a decrease in the weight fraction of apatite in the crystals on the surface comprising Ti metal, apatite crystals, and calcium phosphate [$Ca_3(PO_4)_2$, C-P] crystals.²⁶ Figure 5 shows the thickness of the products, comprising apatite and/or C-P,¹⁷ formed on Ti substrates by PE at various T_{ON} . The figure shows that there is no correlation between the thickness of the products deposited on the Ti substrates and T_{ON} . Therefore, it is considered that the pulse width affects the weight fraction of apatite in the products deposited on the Ti substrates, but not the total amount of products deposited by PE. According to Figs. 3 and 4, it is considered that the weight fraction of apatite crystals in the products deposited on the Ti substrates kept decreasing with T_{ON} . This indicates that the crystal growth of apatite was suppressed with a decrease in T_{ON} .

Figure 6 shows the FTIR spectra of the unetched and etched Ti substrates, which were subjected to PE at vari-

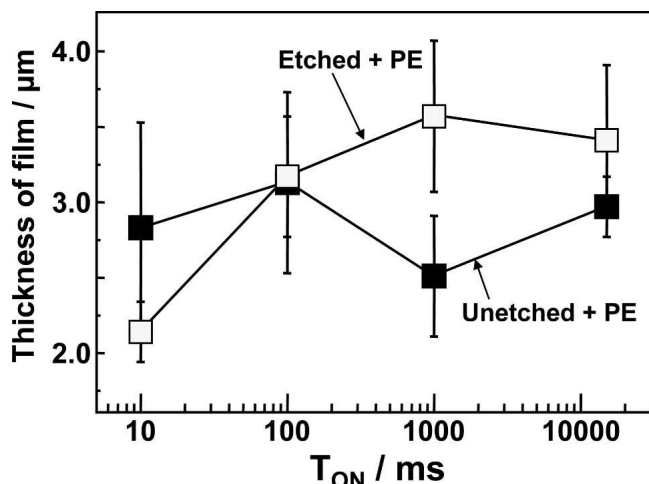


FIG. 5. Total intensities of diffraction peaks of apatite as a function of T_{ON} .

ous T_{ON} . In Fig. 5, all spectra showed PO_4^{3-} absorption bands²⁷ at 1100–1000, 960, 600, and 580 cm^{-1} , an OH^- band at 1650 cm^{-1} , and a CO_3^{2-} band²⁸ or HPO_4^{2-} band²⁹ at 865 cm^{-1} . Furthermore, the spectra of the unetched Ti substrates subjected to PE at T_{ON} above 100 ms and those of the etched Ti substrates subjected to PE at T_{ON} above 1 s showed CO_3^{2-} bands at 1540–1350 cm^{-1} . The existence of this band indicates that carbonate-containing apatite crystals and/or $CaCO_3$ were deposited onto the substrate by PE.¹⁸ The intensity of CO_3^{2-} adsorption bands at 1540–1350 cm^{-1} , PO_4^{3-} absorption band at 960 cm^{-1} , and CO_3^{2-} band²⁸ or HPO_4^{2-} band at 865 cm^{-1} decreased with T_{ON} . This indicates that the amount of CO_3^{2-} in apatite and/or $CaCO_3$ and the fraction of apatite crystals in the deposited products on the Ti substrates decrease with T_{ON} . However, it is considered that the acid etching hardly affects the crystal structure of apatite because similar TF-XRD patterns and FTIR spectra were obtained for both the unetched and etched Ti substrates.

Figure 7 shows the FE-SEM photographs of the unetched and etched Ti substrates subjected to PE at various T_{ON} . For each specimen, the left side is the FE-SEM photograph of the unetched Ti substrates subjected to PE at various T_{ON} , and the right side is that of the etched Ti substrates subjected to PE at various T_{ON} . These photographs show that dense uniform layers were formed on the surfaces of all specimens. Figures 3 and 6 show these layers might be apatite. It was found that the apatite crystals on all specimens had cracks. These cracks might be formed by vaporization of moisture in apatite crystals, when the substrates were dried in air at room temperature after the electrodeposition. The width of the cracks decreased with T_{ON} . During plating, it is known that amorphous thin films have no cracks on its surface.³⁰ Similarly in this study, it is considered that the increase in the amorphous component of the products decreased the crack width.

Figure 8 shows the FE-SEM photographs of the unetched and etched Ti substrates subjected to PE at various T_{ON} followed by the peeling test using Scotch tape. The deposited apatite was easily peeled off by the Scotch tape for the etched and unetched Ti substrates subjected to PE at $T_{ON} = 1$ s and 15 s (see Fig. 7). However, it did not peel off (with the glue of the Scotch tape remaining on the surface) from the etched Ti substrates subjected to PE at $T_{ON} = 10$ ms and 100 ms and the unetched Ti substrates subjected to PE at $T_{ON} = 10$ ms. The glue of the Scotch tape that remained on the surface indicates that the adhesive strength of the apatite coating of these specimens was relatively high. The cracks did not extend outside the scratch trace, and the apatite films deposited on the etched Ti substrates subjected to PE at $T_{ON} = 10$ ms and 100 ms did not come off even though a high load of 200 g was applied to the scratch needle. The results for

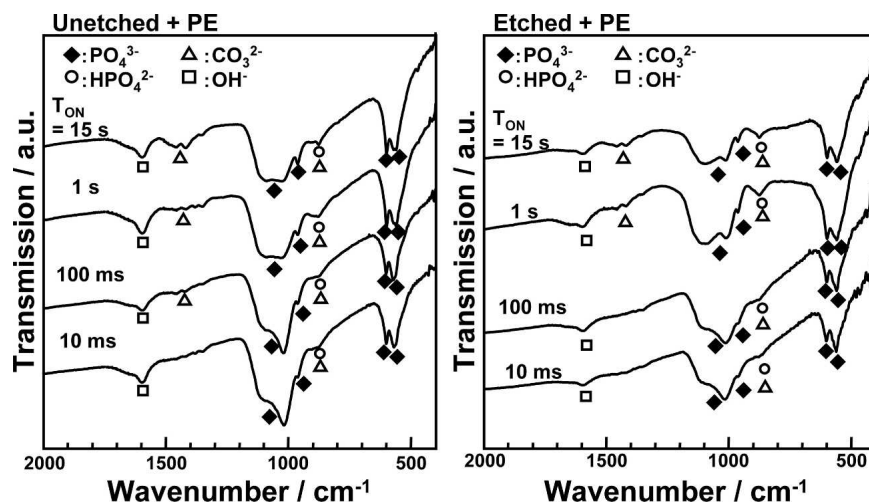


FIG. 6. FTIR spectra of unetched and etched Ti substrates subjected to electrodeposition at various T_{ON} .

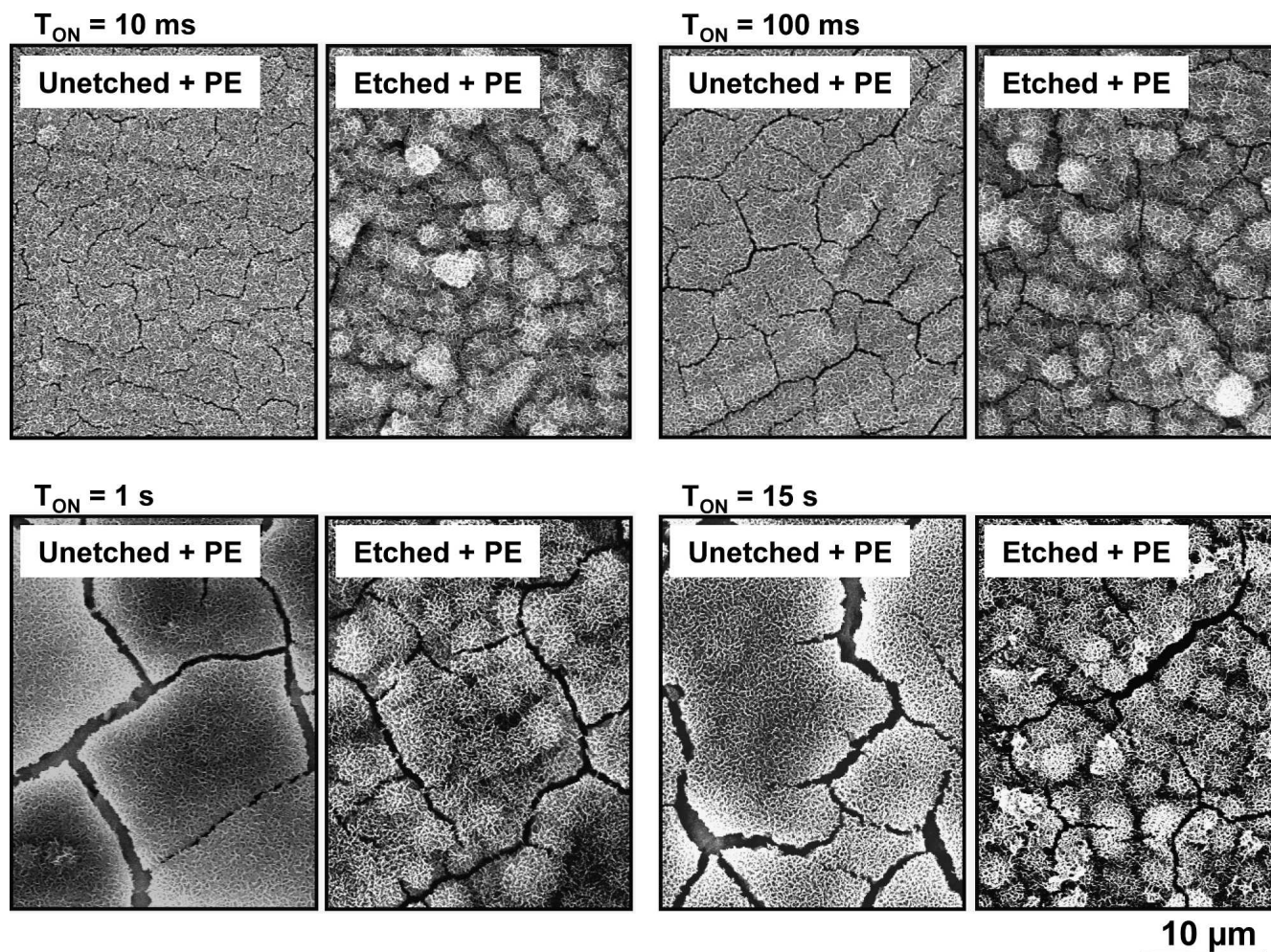


FIG. 7. FE-SEM photographs of unetched and etched Ti substrates subjected to electrodeposition at various T_{ON} .

the above peeling and scratch tests were summarized in Table II.

The mechanism for apatite formation and good adhesion on specimens electrodeposited under a short pulse

current at T_{ON} values, such as 10 ms, can be interpreted as follows. When the potential is loaded at the cathode (Ti), cations (H^+ , Na^+ , K^+ , and Ca^{2+}) in the electrolyte move to the surface of the cathode within 10–30 ms, and

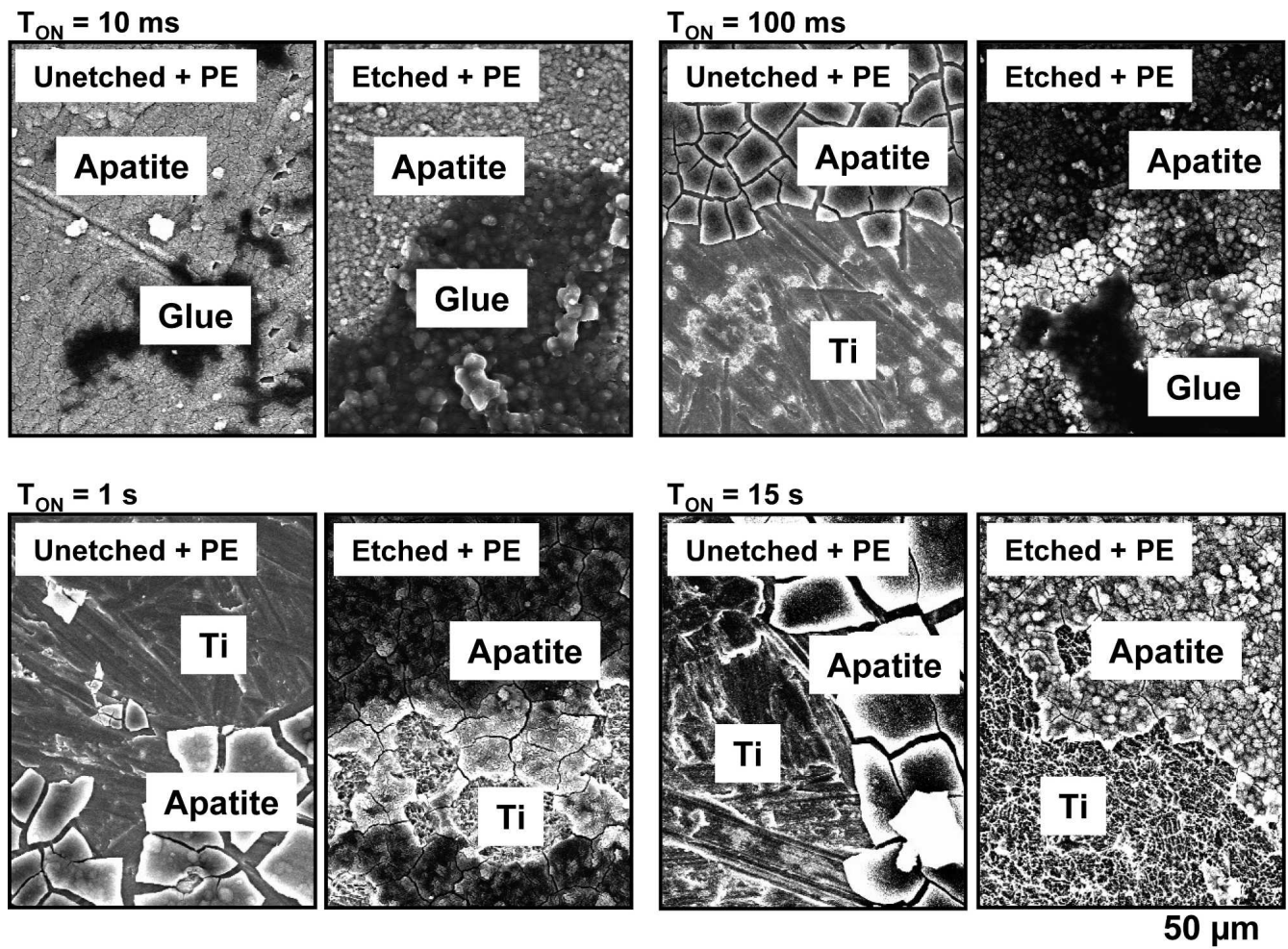
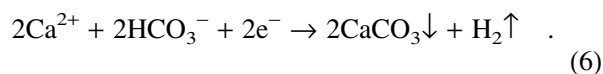
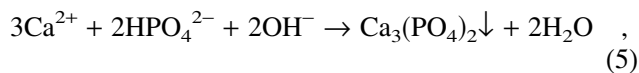
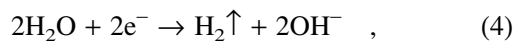


FIG. 8. FE-SEM photographs of unetched and etched Ti substrates subjected to electrodeposition at various T_{ON} followed by peeling test using Scotch tape.

then an electric double layer is formed.³¹ In the electric double layer near the cathode, anions (OH^- , Cl^- , HCO_3^- , and HPO_4^{2-}) can exist due to diffusion. After the electric double layer is formed, the following reaction occurs at the surface of the cathode.¹⁷



Therefore, it is considered that the electric charge is consumed for the formation of the electric double layer and the reduction of H_2O during T_{ON} . Due to reaction (4), the pH value increases at the surface of the cathode. The sudden increase in pH triggers the crystal nucleation of C-P crystals.³² The C-P crystals react with H_2O and HCO_3^- in the surrounding fluid to form apatite, including CO_3^{2-} . The decrease in T_{ON} causes an increase in the

TABLE II. Results for peeling and scratch tests.

T_{ON}	Unetched or etched	Peeling test	Scratch test
10 ms	Unetched	○	—
	Etched	○	○
100 ms	Unetched	X	—
	Etched	○	○
1 s	Unetched	X	—
	Etched	X	—
15 s	Unetched	X	—
	Etched	X	—

○, not peeled off or not coming off; X, peeled off; —, not measured.

consumption rate of the electric charge for the formation of the electric double layer per period. This means that both the reduction of H_2O and the amount of C-P crystal nuclei at the cathode surface per period decrease. As a result, it is considered that the nucleation of C-P crystals with low crystallinity occurred on the Ti surface without sufficiently reacting with other C-P crystals, H_2O and HCO_3^- , and hence small apatite crystals were deposited on the Ti surface. It is known that a residual stress in the

electrodeposited layer gives a large effect on their adhesive strength.³³ In this study, we speculate that the deposition of small apatite crystals increase the number of crystal grain boundary and hence decrease the residual stress that originated from the lattice mismatch between apatite crystals and Ti.³³ Therefore, the adhesive strength between apatite and the Ti substrates was enhanced for the specimen with $T_{ON} = 10$ ms. As shown in Fig. 7, the adhesive strength was also enhanced at $T_{ON} = 100$ ms for the etched Ti substrates subjected to PE. This can be attributed to the anchoring effect of the microstructural roughness formed during acid etching as well as the deposition of small apatite crystals. In conclusion, the electrodeposition technique described in this study is useful for coating bioactive apatite onto metallic materials.

IV. SUMMARY

Ti metal substrates etched in 75% H_2SO_4 were roughened uniformly. Apatite films were deposited onto unetched substrates and substrates etched in 75% H_2SO_4 by electrodeposition under a pulse current. For the specimens electrodeposited at $T_{ON} = 10$ ms, the adhesive strength between apatite and the Ti substrates was greatly improved. In these specimens, we consider that the mismatch relaxation, which resulted from the difference in the lattice constant between apatite crystals and Ti, occurred due to the restraint of the growth of calcium phosphate into apatite.

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