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Calcium-Phosphate Formation on Titanium Modified with Newly Developed Calcium-Hydroxide-Slurry Treatment

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Authors developed a new surface-modification method with calcium-hydroxide slurry, which make it possible to treat a titanium surface by contacting with an alkaline agent containing high concentration of calcium. The objective of the new surface modification method was to improve a bone conductivity of titanium with simple and low-cost processes. The calcium-hydroxide slurry was prepared by mixture of calcium-hydroxide reagent and deionized water. A titanium plate was completely buried in the calcium-hydroxide slurry, and the slurry including the titanium was heated in air at 873 K for 7.2 ks, followed by washing in deionized water, and drying in air. Characterization with X-ray photoelectron spectroscopy revealed that chemical state of the surface-modified-titanium surface was the same as that of calcium titanate. X-ray diffraction pattern showed that the perovskite-type calcium titanate was formed in the surface-modified layer, and depth profile by Auger electron spectroscopy titanium indicated that dioxide layer was formed under the calcium-titanate layer. When the surface-modified titanium was immersed in a Hanks' balanced saline solution for 9 d, hydroxyapatite was formed on the surface-modified-titanium surface, while was not formed on the unmodified-titanium surface without surface modification. However, after 18-d immersion, hydroxyapatite was also formed on unmodified-titanium surface. X-ray diffraction pattern showed that thickness of the hydroxyapatite layer formed on the surface-modified-titanium surface was thicker than that on the unmodified-titanium surface. These results indicated that the calcium-hydroxide-slurry treatment improves the performance of calcium-phosphate formation of titanium. Therefore, the new treatment technique is one of the promising methods for improvement of bone conductivity of titanium. [doi:10.2320/matertrans.48.105]

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Keywords: surface modification of titanium, calcium-hydroxide slurry, calcium titanate, calcium-phosphate formation, bone conductivity

1. Introduction

Titanium has been used as biomaterials, such as stem of artificial joints and dental implant, because of its excellent biocompatibility¹⁻⁴⁾ and mechanical strength,⁵⁻⁷⁾ whereas, it takes several months to obtain good fixation between the titanium and bone. Although bulk properties dictate the mechanical properties of biomaterials, tissue-biomaterial processes are surface phenomena and they are governed by surface properties. In order to activate bone conduction, therefore, various surface modification techniques have been attempted. Hydroxyapatite (HAP) coating is the most popular method, and plasma-splating process is employed for the coating.⁸⁻¹²⁾ With coating of HAP, good biomaterial-bone fixation is obtained.^{10,11)} However, a fracture at the HAP-titanium interface or in the HAP itself is often observed after a long-term use.¹³⁾ Therefore, it can be said that long-term stability of the plasma-sprayed HAP coating is insufficient.

Surface modifications improved the defects of the HAP coating are also studied by some researches. The most important requirement for the improvement is to obtain an unclear interface between the surface-modified layer and titanium substrate, yielding non-destructive interface. Hanawa *et al.* reported that implantation of calcium-ions with an energy of 18 keV modifies titanium surface to be bioactive.^{14,15)} The surface-modified layer of calcium-ion-implanted titanium consists of calcium oxide/hydroxide, titanium oxide, and calcium titanate,^{16,17)} and they concluded that calcium titanate causes of the bioactive surface.¹⁸⁾ Kokubo *et al.* showed that titanium bond to living bone directly if the titanium is treated by soaking in sodium-hydroxide solution

and the subsequent heating.¹⁹⁻²¹⁾ A surface of the treated titanium has sodium-titanate layer with gel-like structure.^{22,23)} After immersed in a simulated body fluid (SBF), sodium in the layer rapidly dissolved in the fluid, calcium in the SBF is taken in the layer instead of sodium, and then, calcium titanate is formed in the treated-titanium surface.^{24,25)} In the above two surface modification methods, calcium titanate is key material for the improvement of bone conductivity.

Recently, some of the authors (N. Ohtsu and T. Hanawa) attempted to improve surface of titanium by calcium-titanate coating with radiofrequency magnetron sputtering.²⁶⁻³⁰⁾ The deposited calcium-titanate layer without some post-treatment has an amorphous structure,^{28,29)} is dissolved in a physiological solution rapidly,²⁷⁾ and is not effective for the improvement.²⁹⁾ Whereas, if successive heating is carried out after the deposition, the calcium-titanate coating is crystallized to be a perovskite structure,^{28,29)} not dissolved easily, and facilitates new bone formation on titanium in hard rat tissue.³⁰⁾ These results indicate that the calcium-titanate coating with perovskite structure modifies titanium surface to be bioactive. However, sputtering technique requires a complex instrument equipped with an ultra-high vacuum system and further is inapplicable to complex-shaped biomaterials. Therefore, the sputtering technique is sometimes not suitable for commercialization, and development of new calcium-titanate coating technique with simple and low-cost is required.

Hanawa *et al.* attempted to form a calcium-titanate on titanium by soaking in a calcium-oxide solution.¹⁸⁾ After soaking in the solution, calcium is taken in the surface, however, the atomic ratio of calcium to titanium (Ca/Ti) is

about 0.1 in the surface,¹⁸⁾ and this ratio much smaller than that of stoichiometric calcium titanate. Hamada *et al.* attempted by hydrothermal modification in calcium-oxide solution.³¹⁾ Although the hydrothermal modification enhances the synthesis of calcium titanate, the atomic ratio of calcium to titanium is still lower than that of stoichiometric calcium titanate, and the calcium-titanate layer is not crystallized to be a perovskite structure. One simple method to increase the concentration of calcium in the modified surface is to treat titanium with an alkaline solution containing high concentration of calcium. However, since the water solubility of calcium oxide and hydroxide is extremely low, it is impossible to prepare the alkaline solution with high calcium concentration.

Authors developed a new surface modification method using calcium-hydroxide slurry. The new method makes it possible to modify titanium surface by contacting with alkaline agent containing high concentration of calcium. The objectives of this study were to characterize a surface of titanium modified with the newly developed calcium-hydroxide-slurry treatment and to estimate performance of calcium-phosphate formation in a SBF. The surface of surface-modified titanium was characterized with grazing incident angle X-ray diffractometry (GI-XRD), X-ray photoelectron spectroscopy (XPS), and depth analysis by Auger electron spectrometry (AES). The surface-modified titanium was immersed in Hanks' balanced saline solution (HBSS), and thereafter, the calcium phosphate formed in the solution were observed with scanning electron microscopy (SEM), and phase of the calcium phosphates were identified with GI-XRD.

2. Experimental Procedures

2.1 Surface modification with calcium-hydroxide-slurry treatment

Commercially available grade 1 pure Ti (cpTi; Furuuchi Chemical Co, Japan) with a disk shape (8 mm in diameter \times 1 mm in thickness), was mechanically polished with SiC paper (#1500) to obtain a rough surface. The titanium plate was soaked in 6 mol L⁻¹ HCl solution at 353 K for 60 s to etch the surface oxide, followed by gently rinsed in deionized water, and dried at 333 K in air. Calcium-hydroxide slurry for the treatment was prepared by a mixture of 1 g of reagent-grade calcium-hydroxide reagent (Nacalai Tesque, Inc) with 1 mL of deionized water (Millipore). Immediately after the mixture, the etched titanium plate was completely buried in the slurry. Then, the slurry containing the titanium plate was heated at 873 K for 7.2 ks in air with an electric furnace. After the heating, the slurry changes to a dry solid with many cracks due to vaporization of water. The titanium plate was retrieved from the slurry, washed ultrasonically in distilled water, and dried at 333 K in air.

2.2 Characterization of surface for the modified-titanium surface.

Chemical state and its composition in the surface were characterized with XPS. The photoelectrons were excited by Al K α radiation ($h\nu = 1486.6$ eV) from a monochromatized X-ray source (SSX-100, Surface Science Inc, U.S.A). The

Table 1 Ion concentrations of Hanks' balanced saline solution (HBSS).

	Concentration (mol L ⁻¹)
Na ⁺	1.42×10^{-1}
K ⁺	5.81×10^{-3}
Mg ⁺	8.11×10^{-4}
Ca ²⁺	1.26×10^{-3}
Cl ⁻	1.45×10^{-1}
HPO ₄ ²⁻	7.78×10^{-4}
SO ₄ ²⁻	8.11×10^{-4}
CO ₃ ²⁻	4.17×10^{-3}

spot size of the X-ray was about 300 \times 500 μ m. The binding energy of Au 4f_{7/2}, Ag 3d_{5/2} and Cu 2p_{3/2} were 84.0, 368.3 and 932.6 eV, respectively and the FWHM of the Au 4f_{7/2} peak was 1.1 eV. Atomic ratio of calcium to titanium (Ca/Ti) was evaluated from the integrated intensities of Ca 2p_{3/2} and Ti 2p_{3/2}, where the sensitive factors installed in the standard SSI software were used for the calculation.

Depth profiles of the elements were determined with AES (JAMP-7100E, JEOL, Japan) with Argon-ion sputtering. The acceleration voltage of electron probe was fixed to 10 kV and that of Ar ions for ion etching was fixed to 3.0 kV. The etching rate estimated with SiO₂ layer was 0.3 nm s⁻¹. The profiles were determined by monitoring of differential spectra for Ti L₃M₂₃M₂₃, Ca L₃M₂M₃, and O KVV transition lines.

Crystallinity of the surface was identified by GI-XRD with a step-scanning mode at 0.8 deg min⁻¹ and the X-ray incident angle $\alpha = 1.0$ degree against the specimen surface (Rotaflex RU-200B, Rigaku, Japan).

2.3 Evaluation of performance for calcium-phosphate formation in HBSS

HBSS was employed as a SBF to evaluate the performance of calcium phosphate formation, whose inorganic contents was similar to that of human blood plasma. The ion concentrations of the HBSS are listed in Table 1. The HBSS was prepared by dissolving prescribed amount of reagent-grade NaCl, KCl, MgSO₄·7H₂O, Na₂HPO₄, KH₂PO₄, and NaHCO₃ (Nacalai Tesque, Inc) in deionized water (Millipore). The pH value of the HBSS immediately after preparation was about 7.4. Four plates of the surface-modified titanium were immersed in 12.6 mL of the HBSS at 310 K. Also, four unmodified-titanium plates, which mechanically polished with SiC paper (#1500), was immersed in HBSS as a control material. Avoiding Si contamination eluted from the vessels,³²⁾ specimens were immersed in the vessels made of perfluoroalkoxy fluoroplastic (Teflon® PFA) and the vessel was completely sealed. In order to avoid the increase of the pH, HBSS were changed once every three days. Two specimens were retrieved from the HBSS after immersion for 9 d; the other two specimens were retrieved after immersion for 18 d. After retrieval, the specimens were gently rinsed with distilled water, and dried at 313 K in air.

Performance for calcium-phosphate formation of the each specimen was estimated by observations of surface using scanning electron microscopy (SEM), where the acceleration voltage of SEM was 10 kV. Phase of the calcium phosphates

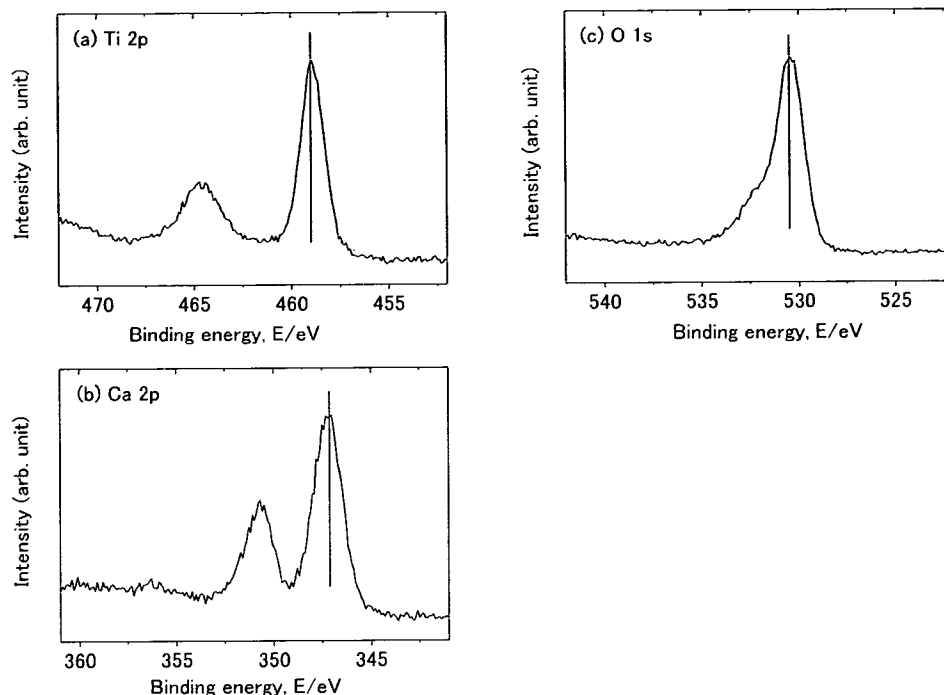


Fig. 1 XPS spectra for the surface-modified-titanium surface: (a) Ti 2p, (b) Ca 2p, and (c) O 1s regions.

were identified with GI-XRD. The measurement conditions of the GI-XRD measurement were the same as the conditions mentioned in the above.

3. Results and Discussions

3.1 Characteristics of the surface-modified-titanium surface

In the XPS survey spectrum for the surface-modified-titanium surface, peaks originated from calcium, titanium, oxygen, and carbon were observed. The binding energy of C 1s peak was 284.8 eV, and origin of carbon was surface contamination adsorbed by exposure to atmosphere. Figures 1(a) and 1(b) show the XPS spectra for Ti 2p and Ca 2p regions, respectively. FWHM of Ti 2p and Ca 2p peaks are 1.4 and 1.8 eV, respectively, and shapes of the peaks are almost symmetry. The results indicate that the chemical states of Ti and Ca consist of one state. Figures 1(c) shows the XPS spectrum for O 1s region. The O 1s peak has small tail in high energy side due to water adsorbed in atmosphere and/or hydroxyl group existing at the outermost surface. It is impossible to estimate which states are dominant in the tail. Binding energies for the modified surface and a calcium titanate of single-crystal are shown in Table 2. The binding energies for the modified surface almost agree with those of the calcium titanate of single-crystal.³³⁾ Furthermore, results of quantitative calculation showed that the atomic ratio of calcium to titanium (Ca/Ti) was 1.0. According to these results, it is concluded that chemical state of the modified surface within the XPS effective depth is calcium titanate.

Depth profile of elements for the modified surface is shown in Fig. 2. The profile of calcium shows that calcium is taken into the surface oxide layer up to ca. 10 nm from the

Table 2 Binding energies of Ti 2p_{3/2}, Ca 2p_{3/2}, and O 1s peaks for the surface-modified-titanium surface and single-crystal calcium titanate.

Electron orbital	Binding energy (eV)	
	Modified surface	Calcium titanate ³²⁾
Ti 2p _{3/2}	459.1	459.1
Ca 2p _{3/2}	347.2	347.3
O 1s	530.5	530.5

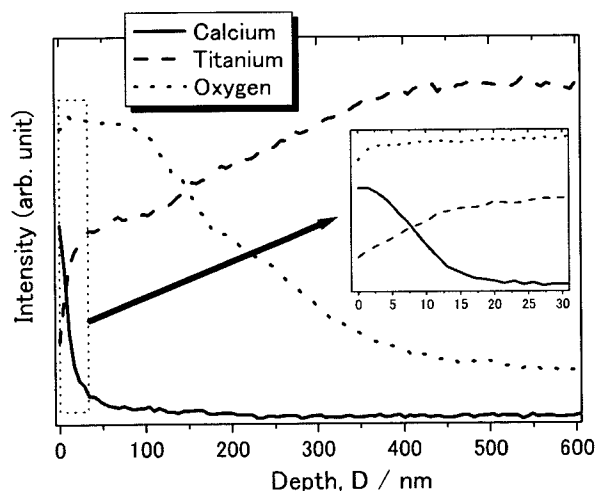


Fig. 2 Depth profile of calcium, titanium, and oxygen for the surface-modified-titanium surface. Inserted figure is enlargement of outermost surface region, which is enclosed by dash line.

outermost surface by the calcium-hydroxide-slurry treatment and the concentration of calcium gradually decreases towards to interior. The profile of oxygen shows that concentration of

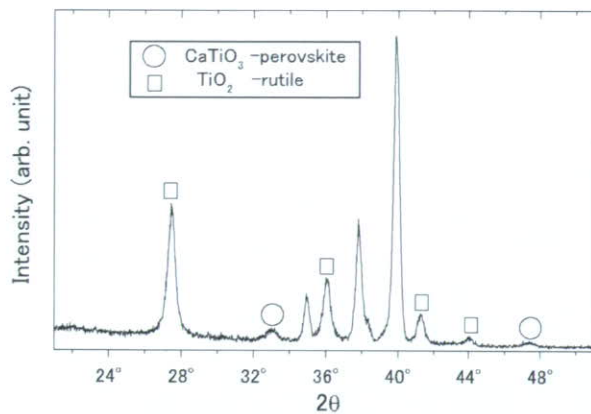


Fig. 3 GI-XRD pattern for the surface-modified-titanium surface.

oxygen diffused into the surface-modified titanium is almost constant up to 120 nm from the outermost surface. Eventually, the surface oxide layer was thickened. The depth range is much larger than that of calcium. These results indicate that surface of the modified titanium has double-layered structure; the outer layer consists of calcium, oxygen, and titanium, and the inner layer consists of oxygen and titanium. Furthermore, in the interface region between the inner layer and titanium substrate, oxygen gradually decreases towards to the substrate. The results indicate that the interface of the modified surface is unclear, that is expected to give non-destructive interface.

Figure 3 shows GI-XRD patterns of the modified surface. Detected peaks except peaks originated from titanium-substrate are identified as those of calcium titanate with perovskite structure and titanium dioxide with rutile structure. It is clear that calcium titanate and titanium dioxide existed at the outer layer and the inner layer, respectively, in the double-layered modified surface.

Some of authors discussed in the previous reports on the characteristics of calcium-titanate film prepared with radio-frequency magnetron sputtering.²⁸⁻³⁰ In the reports, it is concluded that calcium-titanate film which can activate the bone formation should have following two properties: chemical state of calcium is calcium titanate and the film crystallized to be a perovskite-type calcium titanate.^{29,30} Results of the characterization in this study indicated that the calcium-hydroxide-slurry treatment changed titanium surface to the calcium-titanate layer whose characteristics satisfied the requirement for the activation.

3.2 Performance of calcium-phosphate formation in HBSS

SEM images for surfaces of the unmodified and surface-modified titanium after immersion in HBSS for 9 d are shown in Figs. 4(a) and 4(b), respectively. On the surface-modified-titanium surface, many sphere-like precipitates with about 1- μ m diameter are observed. The precipitates are piled up and completely covered the modified surface. On the other hand, no precipitates are observed on the unmodified-titanium surface.

Figure 5 shows the SEM images of the surfaces of the unmodified and surface-modified titanium after immersion in HBSS for 18 d. Also, reduced images are inserted in the upper quarter part of the figures. Both the unmodified and modified surfaces were covered with the precipitate layer. Cracks observed in the layer are due to the evacuation of the water in an ultra-high vacuum. It is clear that the layer formed on the modified surface was thicker than that on the unmodified surface. A part of the layer formed on the modified surface is peeled off. The phenomenon is explained by the effect of stress concentration due to the thick precipitate layer. Peeling off is not observed on the unmodified surface because the layer is not so thick.

Figure 6 shows the GI-XRD patterns for the unmodified-

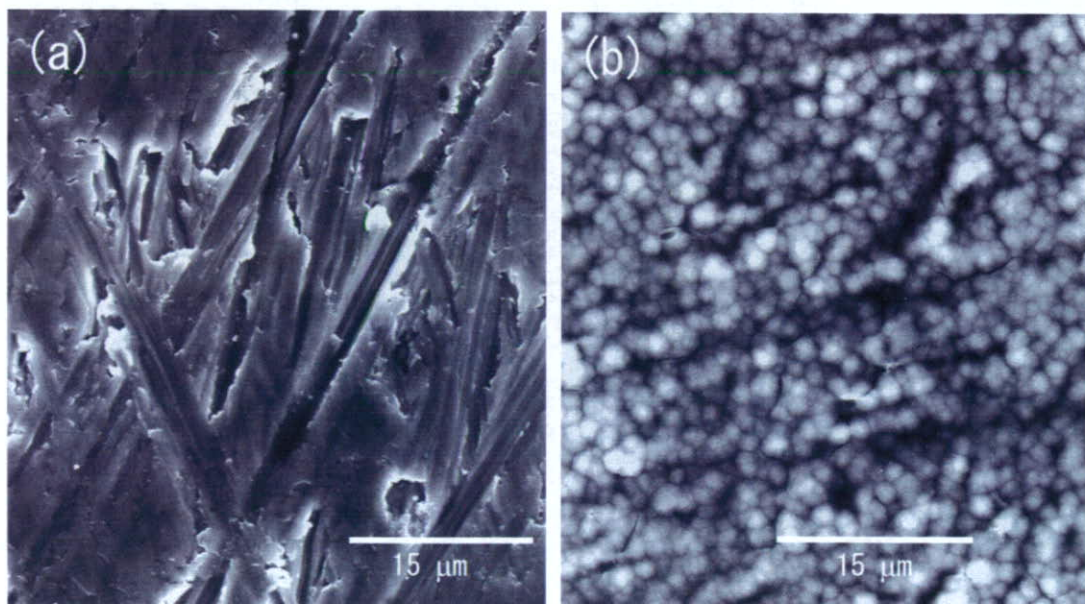


Fig. 4 SEM images for the surface after immersion in HBSS for 9 d: (a) unmodified-titanium surface, and (b) surface-modified-titanium surface.

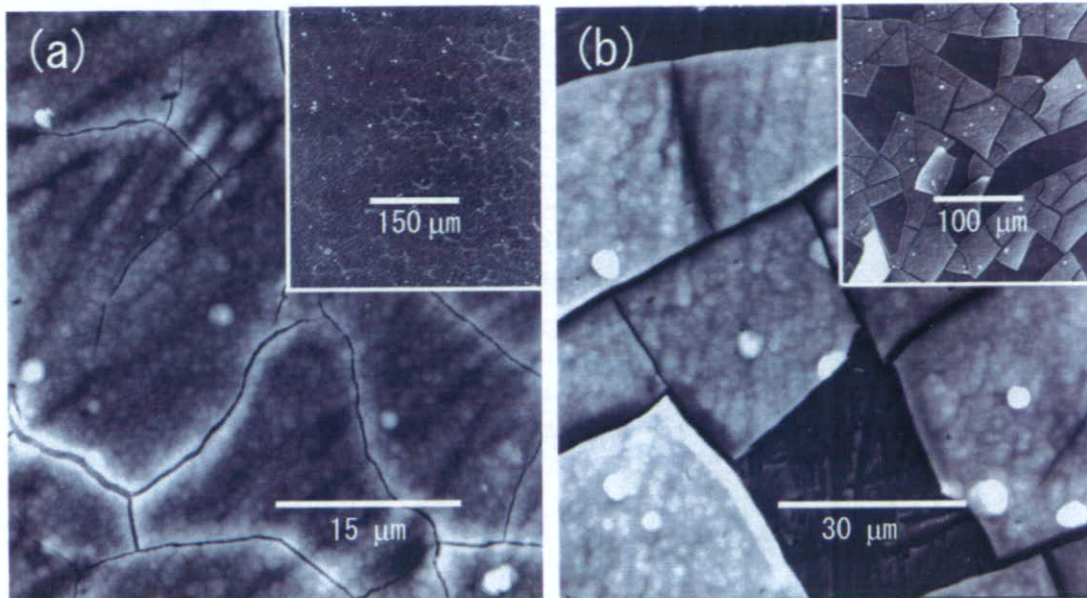


Fig. 5 SEM images for the surface after immersion in HBSS for 18 d: (a) unmodified-titanium surface, and (b) surface-modified-titanium surface. Inserted figures in upper quarter part are reduced SEM images.

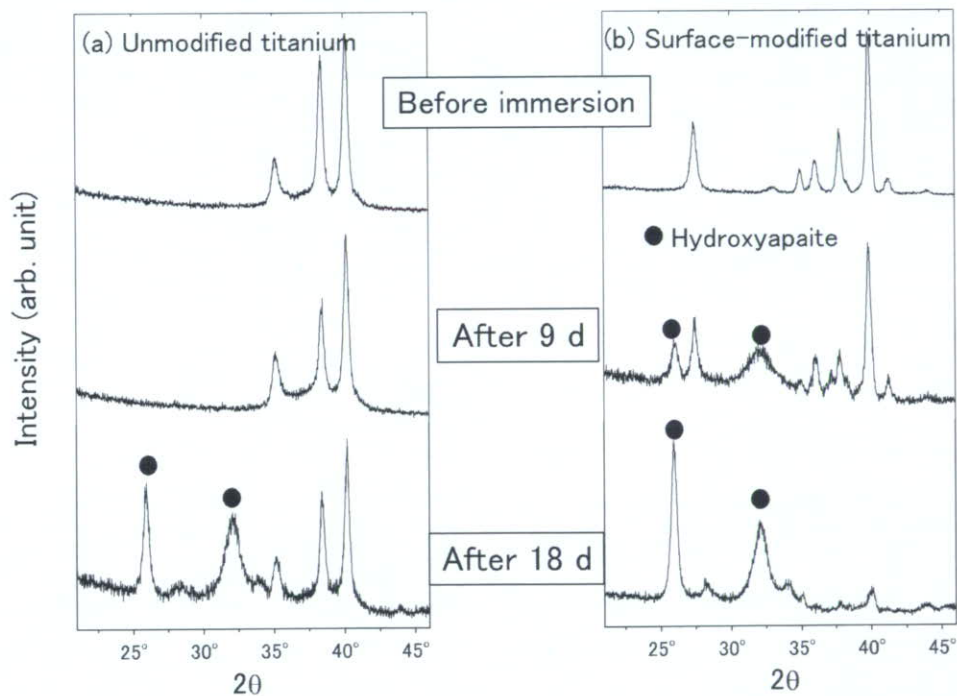


Fig. 6 GI-XRD patterns for (a) unmodified- and (b) surface-modified-titanium surfaces before and after immersion in HBSS for 9 and 18 d.

and surface-modified-titanium surface before and after immersion in HBSS for 9 and 18 d, respectively. New peaks appearing after the immersion are assigned to those of HAP. This result shows that the precipitates formed on the surfaces after the immersion are HAP. At 9 d, the peaks of HAP are observed only in the XRD patterns of the modified surface. This result corresponds to that of the SEM observation. At 18 d, the peaks of HAP are observed in the both modified and unmodified surface. On the other hand, the peaks of titanium

are disappeared in the modified-surface XRD pattern because the thickness of the precipitate layer formed on the modified surface exceeds the effective depth of the GI-XRD. The result indicates that the thickness of the HAP layer formed on the modified surface is thicker than that on the unmodified surface and corresponds to the result of the SEM observation.

The results of the SEM and GI-XRD show that the formation speed of HAP layer on the surface-modified-titanium surface is much faster than that on the unmodified-

titanium surface. Thus, it is concluded that the performance of the calcium-phosphate formation in HBSS is improved by the calcium-hydroxide-slurry treatment. In general, surface which can form calcium phosphates rapidly is bioactive. Therefore, the calcium-hydroxide-slurry treatment is one of the promising treatments for improvement of bone conductivity.

4. Conclusion

By treating with newly developed calcium-hydroxide-slurry treatment, the surface of titanium changes to calcium-titanate layer with perovskite structure. Also, titanium-dioxide layer is formed under the calcium-titanate layer. The characteristics of the surface-modified titanium surface correspond to those of the calcium-titanate film prepared with radiofrequency magnetron sputtering and post-heating which can facilitate the bone formation. On the other hand, the interface between the modified layer and substrate is unclear, expected to give non-destructive interface. Speed of the hydroxyapatite formation on titanium in Hanks' balanced saline solution becomes faster by modifying with the calcium-hydroxide-slurry treatment, and thus, it is concluded that the treatment improve the performance of calcium-phosphate formation on titanium. The results show that the calcium-hydroxide-slurry treatment can modify titanium surface to a bioactive calcium-titanate layer with simple and low-cost, and it is indicated that the new technique is one of the promising treatments for improvement of bone conductivity.

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Determination of the Immobilization Manner of Amine-Terminated Poly(Ethylene Glycol) Electrodeposited on a Titanium Surface with XPS and GD-OES

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Poly(ethylene glycol), PEG, is a bifunctional molecule that inhibits the adsorption of proteins. Therefore, the immobilization of PEG on a metal surface is an important step in making metal surfaces biofunctional. The bonding manner of PEG to a titanium surface is significant for the design of PEG-immobilized materials; however, there are few characterization techniques for the determination of the immobilization manner of PEG. In this study, PEG terminated at one or both terminals with amine bases was immobilized on a titanium surface with electrodeposition and immersion. The electrodeposition was carried out with -5 V for 300 s. The immobilization manner of PEG was characterized using X-ray photoelectron spectroscopy (XPS) with an angle-resolved technique and glow discharge optical emission spectroscopy (GD-OES). As a result, not only electrodeposition but also immersion led to the immobilization of PEG onto a titanium surface. However, more terminated amines combined with titanium oxide as an ionic NH-O with electrodeposition, while more amines randomly existed as NH_3^+ in the PEG molecule with immersion. Moreover, the difference in the amine termination resulted in a different manner of bonding. The PEG terminated at both terminals immobilized in a U shape, and the PEG terminated at one terminal immobilized a brush. Characterization with XPS and GD-OES is useful to determine the immobilization mode of PEG to a solid surface. [doi:10.2320/matertrans.48.287]

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Keywords: titanium, polyethylene glycol, immobilization, electrodeposition, surface analysis

1. Introduction

Recently, biofunctions are required for metals. For example, stents are placed at stenotic blood vessels for dilatation, and blood compatibility or prevention of adhesion of platelets is necessary. Sliding lubrication in blood vessels is important when inserting guide wires and guiding catheters. In addition, if metals were to be used as sensing devices, the control of cell adhesion would be necessary. For these purposes, the fundamental property is the inhibition of protein adsorption. Poly(ethylene glycol), PEG, is a bifunctional molecule that inhibits the adsorption of proteins. Therefore, the immobilization of PEG to a metal surface is an important event to bio-functionalize a metal surface.

There are several stages in the process of immobilizing PEG on solid surfaces that have been recently studied. For example, the immobilization of PEG on silica or titanium surfaces consists of a two-stage reaction.¹⁾ In the first stage, an aminosilane or a thiosilane is reacted with a clean oxide surface to form a surface that is rich in amine or thiol functionalities. In the second stage, a PEG polymer that contains a functional group that is reactive with the modified surface is coupled to the surface. On the other hand, a certain amount of PEG is immobilized on titanium with electrodeposition when PEG is terminated by amine bases.²⁾ The adsorption of albumin on a titanium surface is inhibited by immobilization. The technique is simple and universal for electroconductive materials and those with a complex morphology. However, the immobilization manner of PEG to a titanium surface with electrodeposition is not completely elucidated in the study referred to above.

In this study, PEG with both terminals or one terminal modified with amine bases was immobilized onto a titanium

surface with electrodeposition. The immobilization manner and chemical bonding state were characterized using X-ray photoelectron spectroscopy (XPS) with an angle-resolved technique and glow discharge optical emission spectroscopy (GD-OES). This study will enhance the understanding of the electrodeposition of PEG on metals.

2. Materials and Methods

2.1 Electrodeposition

Both terminals of PEG were terminated with $-\text{NH}_2$ (PEG1000 Diamine, NOF Corporation, Japan), and only one terminal was terminated with $-\text{NH}_2$ (SUNBRIGHT MEPA-10H, NOF Corporation, Japan). The chemical structures of the PEGs are shown in Fig. 1. The molecular weights of both PEGs were about 1,000. These terminated PEGs were dissolved in a 0.3-mol L^{-1} NaCl solution as a concentration of 2 mass%. In the solution, the $-\text{NH}_2$ terminal was dissociated and positively charged as $-\text{NH}_3^+$. The pH of

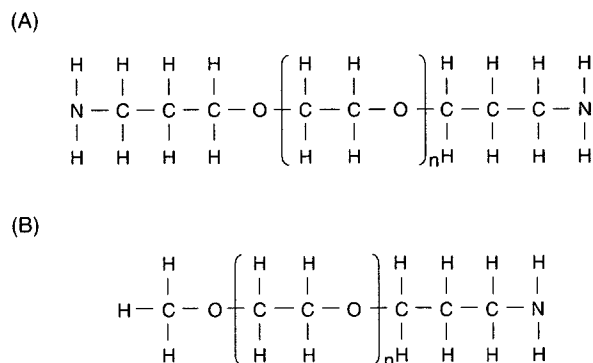
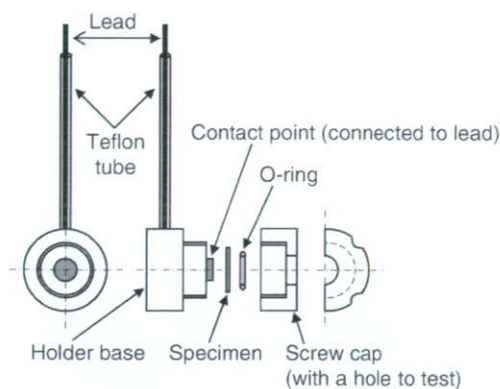


Fig. 1 Chemical structures of PEGs in which both terminals (A) and one terminal (B) were terminated with amine.

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(A)



(B)

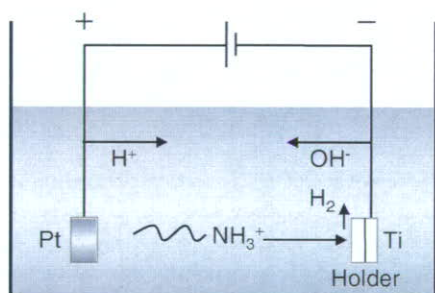


Fig. 2 Design of a polytetrafluoroethylene holder of titanium for electro-deposition (A) and schematic illustration of electro-deposition (B).

the PEG solution with both terminals modified was 11.2, and that of the solution with one terminal modified was 11.0. The resultant solution was used as an electrolyte for electro-deposition at 310 K.

A commercially pure titanium disk (8 mm ϕ \times 2 mm in thickness) with grade 2 was metallographically polished and ultrasonically rinsed in acetone and deionized water (Millipore). The titanium disk was fixed in a polytetrafluoroethylene holder that was completely insulated from the electrolyte with the exception of an open window for electro-deposition (6.0 mm), as shown in Fig. 2(A). Therefore, the exposed area for electro-deposition was 28.3 mm². The open circuit potential of titanium, E_{open} , vs. a saturated calomel electrode, SCE, before electro-deposition was measured as ca. -0.5 V. Thereafter, the cathodic potential was charged from E_{open} to -5 V vs. SCE with a sweep rate of 0.1 V s^{-1} and maintained at this potential for 300 s. During charging, the terminated PEGs migrated electrically to a titanium cathode, where they were deposited as shown in Fig. 2(B). After electro-deposition, specimens were rinsed in deionized water and dried with a stream of nitrogen gas (99.9%). For comparison, titanium was immersed in a solution containing PEG terminated at both terminals for 24 h without any electric charge.

2.2 Ellipsometry

The thickness of the PEG layer deposited on titanium was

Table 1 Photoionization cross sections of level j of element i relative to that of the O 1s, $\sigma_{ij}/\sigma_{\text{O}1s}$.

Level	Photoionization cross section		
	Ti 2p _{3/2}	C 1s	N 1s
σ_{ij}	1.28	0.34	0.62
Reference	6	7	7

determined with an ellipsometer (DVA-36Ls, Mizojiri Optical Co., Ltd.) in air. The use of an ellipsometer resulted in the underestimation of the thickness compared to that in solution. The light source was a He-Ne laser with a wavelength of 632.8 nm, and the incident angle to the titanium surface was 70°. The thickness was calculated by optical constants: the refractive index and absorption coefficient of titanium oxide with the titanium substrate were 2.209 and 3.079,^{3,4} and those of the titanium substrate were 2.22 and 2.99,⁵ respectively.

2.3 XPS

The immobilization manner of PEG to a titanium surface and the chemical bonding state were characterized using XPS (SSX100, SSI, UK). All binding energies given in this paper are relative to the Fermi level, and all spectra were excited with the monochromatized Al K α line (1486.61 eV). The spectrometer was calibrated against Au 4f_{7/2} (binding energy, 84.07 eV) and Au 4f_{5/2} (87.74 eV) of pure gold and Cu 2p_{3/2} (932.53 eV), Cu 2p_{1/2} (952.35 eV), and Cu Auger L₃M_{4,5}M_{4,5} line (kinetic energy, 918.65 eV) of pure copper. The energy values were based on published data.⁶ In order to estimate the photoelectron peak intensities, the background was subtracted from the measured spectrum according to Shirley's method.⁷ The composition and thickness of the surface oxide and the composition of the substrate were simultaneously calculated according to previous studies.^{8,9} Empirical data¹⁰ and theoretically calculated data¹¹ of the relative photoionization cross sections were used for the quantification. The relative photoionization cross sections used in this study are summarized in Table 1, where $\sigma_{ij}/\sigma_{\text{O}1s}$ represents the relative photoionization cross section of a level j electron of an element i to that of O 1s electrons.

An angle-resolved technique for XPS was applied to specimens at photoelectron take-off angles of 12°, 24°, 35°, 53°, and 90°, where the take-off angle is defined as the angle between the direction of the photoelectron path to the electron spectrometer and the specimen surface.

2.4 GD-OES

The depth profiles of elements from a PEG-electrodeposited layer and titanium substrate were determined using a GD-OES instrument (Jovin Yvon RF 5000GD-OES, HORIBA, Japan) with a power of 40 W in 600-Pa argon atmosphere. The spectral wavelengths (nm) employed for detecting various elements were as follows: H, 121.57; C, 156.14; N, 149.26; Ti, 365.35; and O, 130.00.^{12,13} The sputtering time was converted directly to the depth, assuming that the sputtering rate was constant throughout the analysis of the PEG-electrodeposited layer and titanium substrate.

Table 2 Thicknesses of the PEG layers determined by ellipsometry. B-PEG: PEG terminated at both terminals. O-PEG: PEG terminated at one terminal.

The type of PEG/ immobilization methods	Thickness of PEG layers [nm]
B-PEG/Electrodeposition	2.37
O-PEG/Electrodeposition	1.85
B-PEG/Immersion	3.87
Mean (standard deviation)	

3. Results and Discussion

3.1 Thickness of PEG layers by ellipsometry

Table 2 shows the thicknesses of the PEG deposition layers determined by ellipsometry. These thicknesses are measured in air; therefore, the real thickness in solutions is larger than these values. The thickness of the deposition layer, in other words, the amount of deposited PEG, is the largest in this order: B-PEG immobilized with immersion for 24 h, B-PEG immobilized with electrodeposition for 300 s, O-PEG immobilized with electrodeposition for 300 s (B-PEG: PEG terminated at both terminals. O-PEG: PEG terminated at one terminal). This indicated that electrodeposition was more effective than immersion for the deposition of PEG on the titanium surface.

3.2 Chemical state of titanium and oxygen by XPS

Carbon, oxygen, titanium, and nitrogen were detected using XPS. Sodium and chlorine were not detected. Carbon and nitrogen originate from PEG molecules.

The typical Ti 2p electron energy region peak on the XPS spectrum is shown in Fig. 3. The Ti 2p peak gave four doublets according to the valences, Ti^0 , Ti^{2+} , Ti^{3+} , and Ti^{4+} . Published data¹⁴⁾ were used for the binding energy of titanium at each valence. Figure 3 shows an example of the decomposition of the Ti 2p spectrum. Ti^0 originating from a titanium substrate was detected at a large take-off angle but not at a small take-off angle. Ti^{2+} , Ti^{3+} , and Ti^{4+} peaks were detected at all take-off angles and originated from titanium oxide, and the integrated intensity of Ti^{4+} was the largest. The titanium surface oxide mainly consisted of TiO_2 containing a small amount of TiO and Ti_2O_3 . In other words, the titanium surface was covered mainly by TiO_2 .¹⁵⁾ Therefore, the oxygen detected by XPS originated from PEG molecules and titanium oxide covering the titanium substrate, and titanium originated from both titanium oxide and the titanium substrate.

The O 1s electron energy region peak is deconvoluted to peaks originated from O^{2-} , hydroxide, or hydroxyl groups, OH^- , and hydrate and/or adsorbed water.¹⁶⁾ Naturally, a titanium surface oxide contains hydroxyl groups at the surface and inside the oxide.

3.3 Depth profiles of elements obtained by the XPS angular-resolved technique

The entire PEG layer was detected by XPS because PEG was immobilized on titanium that was covered with titanium oxide. Surface oxide was detected by XPS at all take-off angles, and the titanium substrate was detected only at high

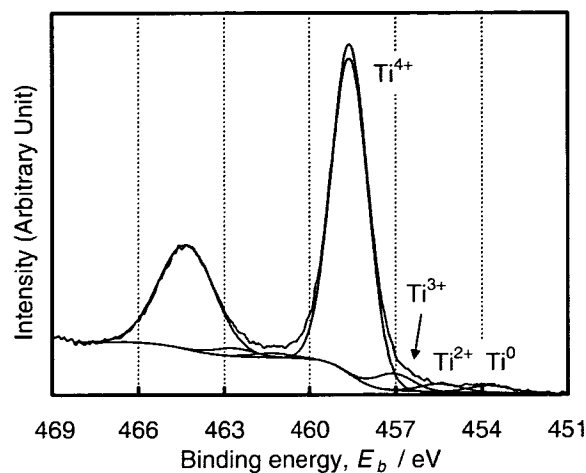


Fig. 3 Typical Ti 2p energy electron region XPS peak and decomposition into eight peaks ($2p_{3/2}$ and $2p_{1/2}$ electron peaks in four valences). The numbers with arrows are valence numbers.

take-off angles. When the PEG layer and/or titanium oxide layer is thick, the intensity of Ti^0 decreases because the signal from the titanium substrate is small.

The relative concentrations of carbon, oxygen, titanium, and nitrogen were calculated assuming that the gross amount of these elements, as detected using XPS, was 100 mole percent. The apparent concentrations of elements calculated under the assumption of a homogeneous depth distribution within the PEG-immobilized layer were dependent on the take-off angle, generally suggesting the existence of concentration gradients. The concentrations of carbon, oxygen, titanium, and nitrogen obtained by the angle-resolved technique are plotted in Fig. 4 against the average escape depth of photoelectrons. In the average effective escape depth of photoelectrons for angle-resolved XPS measurements, λ is the average mean-free-path of C 1s, O 1s, Ti 2p, and N 1s photoelectrons, and the effective escape depth is estimated as λ times \sin (take-off angle).

The concentrations of carbon at small take-off angles were high, indicating that carbon atoms distribute more in the outer layer than in the inner layer in the PEG layer (Fig. 4(A)). The concentrations of oxygen at small take-off angles were low, indicating that oxygen distributes more in the inner layer than in the outer layer in the PEG/titanium oxide combination layer (Fig. 4(B)). The concentrations of titanium at small take-off angles were low, indicating that titanium distributes more in the inner layer than in the outer layer, which is natural because the PEG layer is formed on titanium oxide covering a titanium substrate (Fig. 4(C)). In these elements, no difference between electrodeposition and immersion was observed.

The concentration of nitrogen is relatively low when compared with those of carbon, oxygen, and titanium (Fig. 4(D)) because the concentration of nitrogen in the PEG molecule is very small (see Fig. 1). At all take-off angles, nitrogen in electrodeposited PEG terminated at both terminals and PEG terminated at one terminal were much smaller than both-terminated deposited with immersion. Nitrogen originates from terminal amines in PEG. The

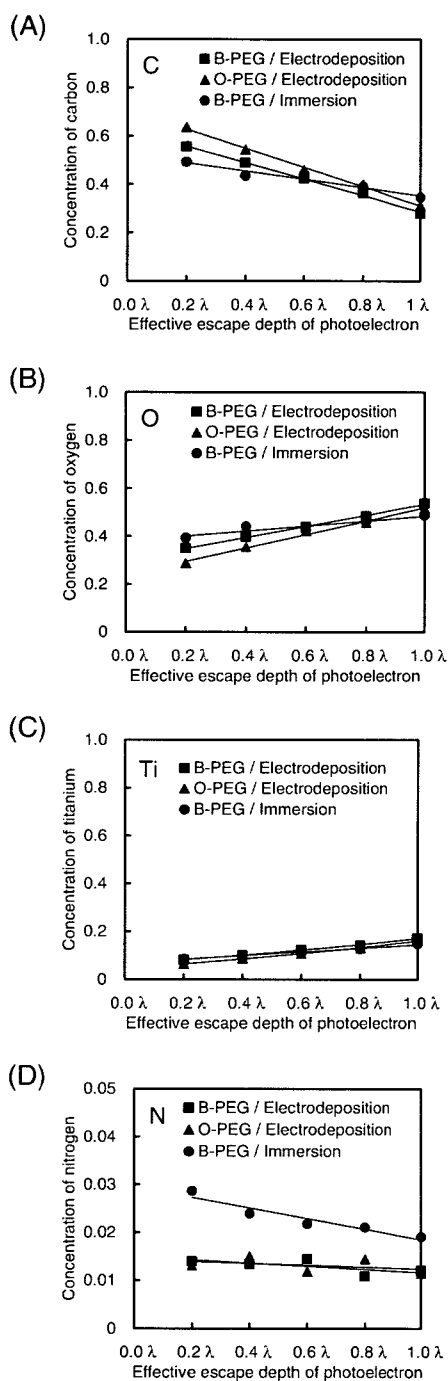


Fig. 4 Concentrations of carbon (A), oxygen (B), titanium (C), and nitrogen (D) against the average effective escape depth of photoelectrons for angle-resolved XPS measurements. Lambda (λ) is the average escape depth of C 1s, O 1s, Ti 2p_{3/2}, and N 1s photoelectrons, and the effective escape depth is the escape depth times sin (take-off angle). B-PEG: PEG terminated at both terminals. O-PEG: PEG terminated at one terminal.

photoelectron signals detected by XPS from a deep site are weak because they decay as they pass through molecules and solids. Therefore, more nitrogen existed inside the PEG-immobilized layer. The results above show that the PEG layer exists on the titanium surface oxide and the oxide exists on the titanium substrate. In addition, more nitrogen atoms in PEG molecules are located inside the PEG-immobilized

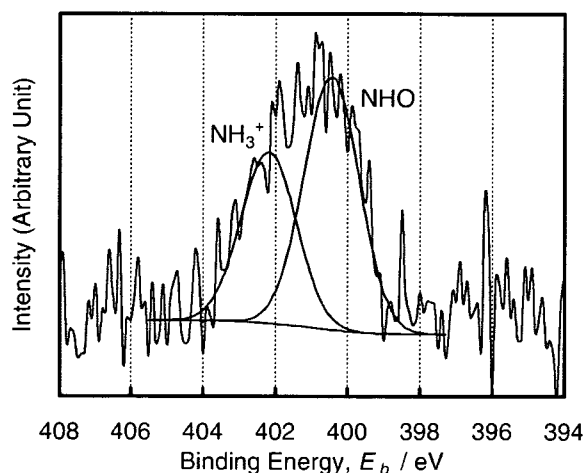


Fig. 5 N 1s electron energy region XPS peak and decomposition of the peak into two component peaks originating from the chemical state of NH_3^+ and NH-O.

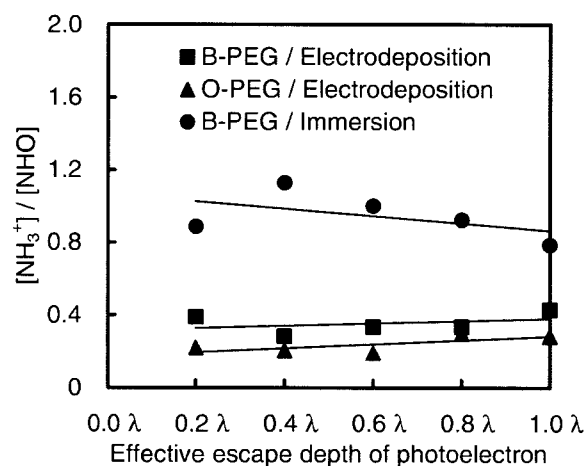


Fig. 6 Ratio of the integrated intensity of a peak originating from NH_3^+ to that from N-HO, $[\text{NH}_3^+]/[\text{N-HO}]$, against the average effective escape depth of photoelectrons for angle-resolved XPS measurements. Lambda (λ) is the average escape depth of N 1s photoelectrons, and the effective escape depth is the escape depth times sin (take-off angle). B-PEG: PEG terminated at both terminals. O-PEG: PEG terminated at one terminal.

layer as a result of electrodeposition than they are with immersion. PEG immobilized on titanium does not always form a single layer; but it forms a multi-layer. Even in this case, nitrogen in each PEG layer exists inside the layer.

3.4 Chemical bonding of amine to a titanium surface as determined by XPS

Typical XPS spectra of an N 1s electron energy region and the deconvolutions of these peaks are shown in Fig. 5 according to published data.^{17,18)} A change in the ratios, $[\text{NH}_3^+]/[\text{N-HO}]$, in the N 1s peak is shown in Fig. 6. This ratio is much smaller in electrodeposited specimens than it is in immersed specimens. Nitrogen atoms in electrodeposited specimens existed as a N-HO bond rather than as NH_3^+ . The N-HO bond is preferential at the interface between an amine and a titanium surface, *i.e.*, in the first layer of PEG

deposited, according to the chemical structure of PEG. The NH_3^+ bond is preferential in the second and third layers. The N-HO bond is an ionic bond formed between NH_3^+ and OH^- on titanium oxide. This indicated that a strong ionic bond was formed between the amine and titanium oxide with electrodeposition, while a weaker bond was formed with immersion.

A certain amount of PEG is immobilized on titanium not only with electrodeposition but also with immersion when PEG is terminated by amine and charged in an aqueous solution. However, more terminated amines exist at the surface of titanium and combine with titanium surface oxide as a N-HO bond with electrodeposition, while more amines randomly exist in the PEG layer and fewer amines form an ionic bond with titanium oxide with immersion. In PEG terminated at both terminals, the PEG deposited in a U-shape because both terminals combined with the titanium surface; in PEG terminated at one terminal, the PEG deposited as a brush because only one terminal combined with the surface in the first immobilized layer.

3.5 Depth profiles of elements by GD-OES

The GD-OES results are expressed with the emission lines plotted against the acquisition time. The depth profiles of elements of a PEG layer immobilized with electrodeposition on titanium and that with immersion obtained with GD-OES are shown in Fig. 7. The dashed line represents the interface between the PEG layer and titanium surface oxide, which is determined by the following element profiles: the minimum value in hydrogen and change in the slope of titanium. Hydrogen is absorbed to titanium by a specimen preparation process that includes polishing and electrodeposition. Hydrogen evolution during electrodeposition increases with a cathodic potential according to the Pourvaix diagram;¹⁹⁾ therefore, the amount of absorbed hydrogen increased after electrodeposition.

The nitrogen signal outside the interface originates from PEG molecules, while that inside originates from the nitrogen-absorbed titanium oxide and substrate. In the electrodeposition of PEG terminated at both terminals, nitrogen in the PEG molecules increased toward the interface (Fig. 7(A)); however, in immersion, it decreased (Fig. 7(B)). That is, more nitrogen atoms in PEG molecules are located inside the PEG-immobilized layer formed with electrodeposition than inside that formed with immersion. In this sense, the XPS result shown in Fig. 4(D) is completely in accordance with this GD-OES result.

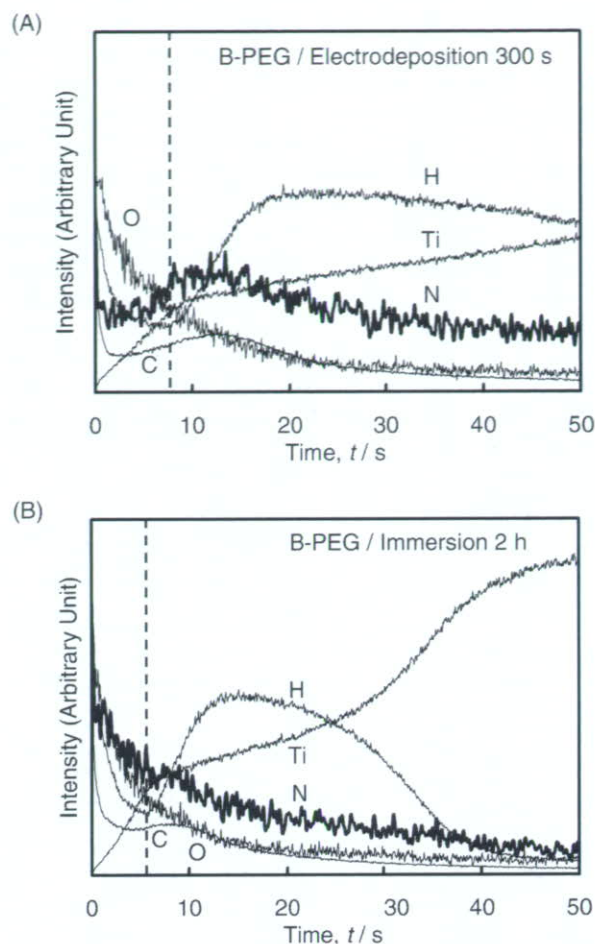


Fig. 7 Depth profiles of elements from PEG-immobilized titanium with electrodeposition (A) and immersion (B) determined by GD-OES. The dashed lines represent the interface between the PEG molecule and titanium.

According to the results above, the immobilization manner of the first PEG layer determined with XPS and GD-OES is illustrated in Fig. 8.

4. Conclusions

In this study, the immobilization manner of PEG terminated with amines at both terminals and at one terminal was characterized with an XPS angular-resolved technique and

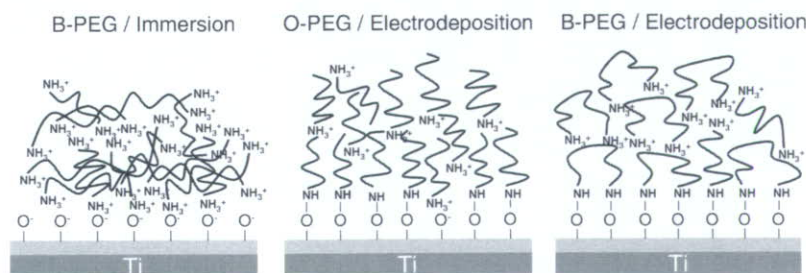


Fig. 8 Schematic model of the deposition manner and chemical bonding state of PEG with immersion and electrodeposition. B-PEG: PEG terminated at both terminals. O-PEG: PEG terminated at one terminal.

GD-OES. In the first layer of immobilized PEG formed with electrodeposition, more terminated amines exist at the interface between the PEG layer and titanium oxide and combine with titanium oxide as a N-HO ionic bond formed between NH_3^+ and OH^- , while more amines randomly exist as NH_3^+ in the PEG layer formed with immersion. In the PEG terminated at both terminals, the PEG immobilized as a U-shape because both terminals combine with the titanium surface; in the PEG terminated at one terminal, the PEG was immobilized as a brush because only one terminal combines with the surface. Over second layers, nitrogen atoms mainly exist as NH_3^+ , while nitrogen is located deeper within the PEG layer. The characterization with XPS and GD-OES is useful to determine the immobilization mode of PEG relative to a solid surface.

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Inhibition Effect of Zirconium Coating on Calcium Phosphate Precipitation of Titanium to Avoid Assimilation with Bone

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In order to control of the calcium phosphate precipitation of Ti in body fluid, which might result in assimilated bone re-fracture during removal operation of implanted devices such as femoral nails and bone screws after healing, vacuum vapor deposition of Zr on Ti Substrate was carried out. The calcium phosphate precipitation was evaluated through the immersion test into the Hanks' solution. Scanning electron microscopy (SEM) and X-ray photoelectron spectroscopy (XPS) were conducted to evaluate inhibition effect of coated Zr layer.

Calcium phosphate particle were observed on the Ti with and without 90-nm thick Zr coated layer by SEM. On the other hand, on the specimens with 130-nm thick Zr coated layer and those of Zr sputter-coated, no calcium phosphate particle was observed. By the surface analysis of the specimens Zr sputter-coated, Ca was not identified. It was considered to form zirconium phosphate instead of calcium phosphate by immersion in the Hanks' solution. It was suggested that Zr coating technique can be applied to orthopedic devices to avoid re-fracture of bone by calcium phosphate precipitation during removal operation. [doi:10.2320/matertrans.48.301]

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Keywords: biomedical titanium alloys, zirconium coating, surface modification, calcium phosphate, biocompatibility

1. Introduction

Due to its less toxicity and superior biocompatibility, Ti and its alloys are used as biomaterials widely in the field of orthopedics. It is well known that Ti is thermodynamically active against to maintain a stable passive oxide film. This passive film acts as a protective film for corrosion. In a macroscopic view, the film is stable, namely inactive. However the partial dissolution and precipitation are repeated constantly in a microscopic view.¹⁾ Therefore, when Ti is implanted in living body, the composition of the oxide film is gradually changed by influence of the environment during repeated dissolution and precipitation. Elements contained in body fluid, such as P, Ca and S are involved in oxide film. Then calcium phosphate is formed on the surface of Ti. This phenomenon is confirmed both *in vivo* and *in vitro*.²⁻⁴⁾ In general, bone conductivity of a material could be evaluated with the formation of calcium phosphate on the material in a simulated body fluid.

For the fast fixation of artificial hip joints to hard tissue after surgical operation, the enhancement of this calcium phosphate precipitation is a key technology and several kinds of surface modification have been developed, such as electrolysis techniques,^{5,6)} immersion in alkaline solution,⁷⁾ immersion in hydrogen peroxide,⁸⁾ hydro-thermal treatment⁹⁾ and calcium ion implantation^{10,11)} to modify the surface inducing active interactions between the modified surface and hard tissue. Some of them have been already applied to the commercial artificial hip joints to shorten the fixation period.

However, for some kinds of internal fixation devices implanted into marrow, such as femoral, tibial and humeral nails and screws for bone plate fixation, titanium's property forming fastly calcium phosphate is not ideal. For these

removal implant devices, assimilation with bone by the calcium phosphate precipitation might cause in re-fracture of the bone during removal operation. Thus, for the removal implant devices into marrow, the inhibition of the calcium phosphate precipitation is required. The requirement is completely opposite against the purpose of a large portion of surface treatments in Ti materials for the acceleration of bone formation.

Zirconium is known to have less toxicity and good corrosion resistance similar to Ti. Ti-13Nb-13Zr alloy has been developed in early 1990s as a beta-type Ti orthopedic alloy. In the previous paper on alloys' surface analysis after immersion into Hanks' solution, an artificial body fluid simulating inorganic components of the extracellular body fluid, zirconium phosphate precipitation was observed on pure Zr and Ti-Zr binary alloys containing more than 34.4 mol%³⁾ instead of calcium phosphate precipitation seen, while calcium phosphate was precipitated on Ti and Ti-14.9 mol%Zr alloy. It was concluded that zirconium oxide didn't involve Ca ion during repeated dissolution and precipitation of oxide. In other words, Zr has an ability to prevent the precipitation of calcium phosphate.

In the present study, vacuum vapor deposition of Zr onto the Ti disk substrates was carried out in order to control of the calcium phosphate precipitation of Ti. The calcium phosphate precipitation was evaluated through the immersion test into the Hanks' solution, followed by scanning electron microscopy, SEM, and X-ray photoelectron spectroscopy, XPS.

2. Experimental

2.1 Specimens

Disks of Ti (10 mm in diameter and 1.5 mm in thickness) were cut out from pure Ti rods (99.9 mass% in purity, Nilaco) by an abrasive wheel cutter. Top surface for Zr deposition was mirror-polished with diamond paste (9 μm) and alumina

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Table 1 Ion concentration of Hanks' solution.

Ion	Na ⁺	K ⁺	Mg ⁺	Ca ⁺	Cl ⁻	HCO ₃ ⁻	HPO ₄ ²⁻	SO ₄ ²⁻
Concentration (mol%)	141.9	5.8	1.7	1.3	144.8	4.2	0.9	1.7

paste (0.05 μm). Bottom surface and round surfaces were finished with 320 grid SiC papers.

Vacuum vapor deposition of Zr onto the Ti disk substrates was carried out using a commercial vacuum evaporator (JEE-420, JEOL) with Zr ingots on tungsten baskets. The basket (Zr ingot) height was fixed 80.0 mm above of the Ti disks. Five or six disks were deposited same time of deposition. Zirconium ingots were made from sponge Zr (99.6 mass% in purity, Toho Titanium) by argon arc melting on a water-chilled copper hearth. Half-masked disks were used to measure Zr coated layer thickness. Coated layer thickness was measured using a laser profile microscope (VF-7500, Keyence) by step height of coated part and uncoated part of a specimen which was half-masked tightly with a small piece of thin paper.

Reference materials, such as Ti without Zr and sputter-coated Zr on Ti substrates (Hokuto Denko) were also tested. The thickness of the coated layer on the latter specimens was approximately 1 μm which was measured by the cross-sectional optical microscopy of a specimen. The surface of sputter-coated Zr showed slight roughness ($R_a = 0.26 \mu\text{m}$).

2.2 Immersion test in Hanks' solution

The immersion test was conducted with Hanks' solution (pH 7.4), an artificial body fluid simulating inorganic components of the extracellular body fluid, at 310 K. The ion contents of Hanks' solution are shown in Table 1. Specimens were immersed into 50 mL of Hanks' solution in sealed containers made of polytetrafluoroethylene (PTFE) for 1 d (specimens for XPS) or 7 d (those for SEM). The PTFE containers were kept in an incubator to keep the temperature at 310 K. For the immersion test for SEM, Hanks' solution was changed once in three days. The pH alteration during immersion tests was detected using a pH meter (HM-60V, DKK-Toa).

2.3 Surface observation and analysis

Surface observation was carried out with specimens after immersion test for 7 d with an SEM (JSM-6400, JEOL) at 7 to 15 kV of acceleration voltage. Surface analysis for 1 d immersed specimens was conducted with an XPS (SSX-100, SSI). An X-ray source was monochromatized Al K α (1486.61 eV) accelerated with 15 kV. The photoelectrons were detected through 58.7 eV in FAT pass energy, with 35° of detection angle under a vacuum atmosphere at 10⁻⁷ Pa or less. Binding energies of photoelectrons were calibrated by binding energy peak of C 1s (binding energy, 285.0 eV) originating from so-called contaminated carbon. The spectrometer was calibrated against Au 4f_{7/2} (84.07 eV) and Au 4f_{5/2} (87.74 eV) of pure gold and Cu 2p_{3/2} (932.53 eV), Cu 2p_{1/2} (952.35 eV) and Cu Auger L₃M_{4,5}M_{4,5} line (kinetic energy, 918.65 eV) of pure copper. The energy values were based on the published data.¹²⁾ Analysis was conducted at least two points of a specimen. The composition and

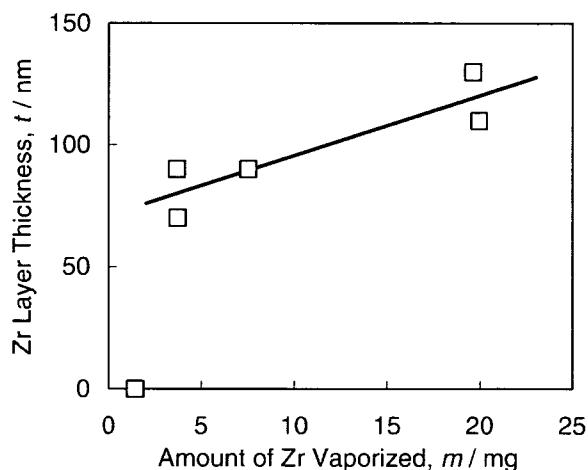


Fig. 1 Relation between amount of Zr vaporized and layer thickness after vapor deposition.

thickness were determined with an inductively calculating method presented by Asami *et al.*¹³⁾

3. Results and Discussion

3.1 Thickness of Zr coated layer

Mass loss of Zr ingot after each vacuum vapor deposition was varied from 3.69 mg to 19.95 mg. Relation between mass loss and thickness of Zr coated layer measured by the laser profile microscope is shown in Fig. 1. It shows good linear relation at higher than 3.69 mg vapor deposition. It is suggested that the thickness of Zr coated layer can be controlled by the vaporized amount of Zr.

3.2 SEM observation after immersion test

Scanning electron micrographs of surface of Ti without Zr coating, Ti with Zr vapor deposited with coating thickness as 90 nm and 130 nm, and Ti with Zr sputter-coated are shown in Fig. 2. On the surface of Ti without Zr coating, precipitates with 1 μm in diameter are observed. In contrast, no precipitates were observed on Ti specimens with Zr vapor-coated layer. On the specimen with sputter-coated layer showed very small amount of precipitates. It is considered formed on the sputter-coated layer due to its rough surface.

3.3 XPS surface analysis before and after immersion test

Specimens immersed into the Hanks' solution for 1 d were used for the XPS analysis. Figure 3 shows the survey spectrum of the specimens (a) before and (b) after immersion. From the Ti without and with Zr vapor deposited (90 nm thickness) after immersion into Hanks' solution, peaks corresponding to binding energy of P 2p clearly seen.

Figure 4 shows narrow area spectrum of P, Ca, Ti, Zr and O. Examples of peak of P, Ca, Ti and O are taken from Ti without Zr coating after 1-d immersion in the Hanks' solution, while that of Zr from Ti with Zr vapor deposited with 130-nm coating thickness.

Spectra of P 2p was obtained as combined peak at around 134 eV. According to the previous report, P 2p binding

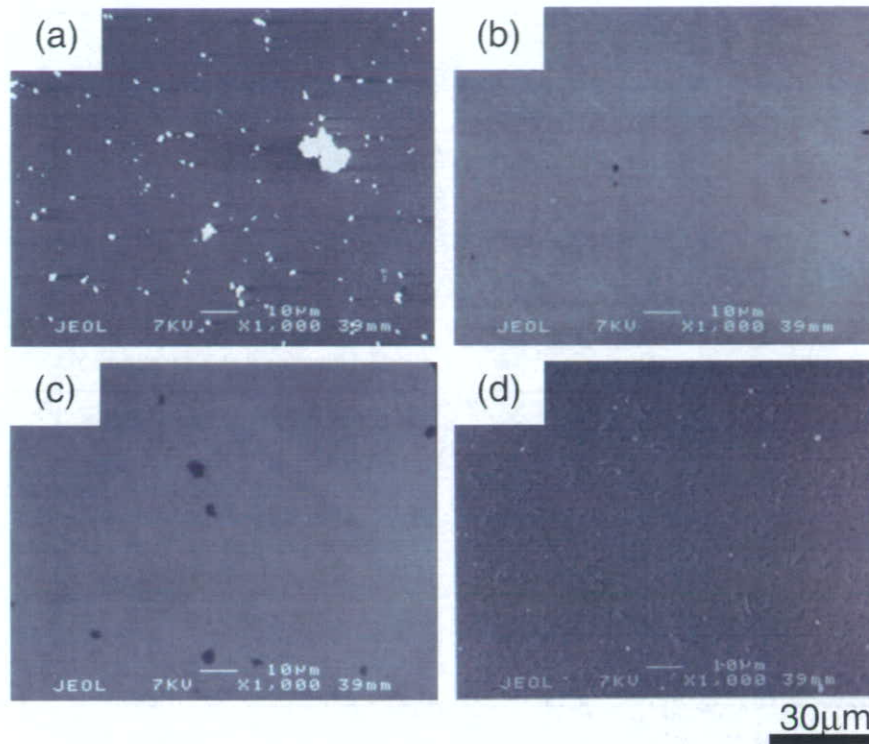


Fig. 2 Specimens' surface after 7 d immersion into Hanks' solution at 310 K, (a) Ti without Zr coating, (b) Ti with Zr vapor-coated (90 nm), (c) Ti with Zr vapor-coated (130 nm), and (d) Ti with Zr Spatter-coated.

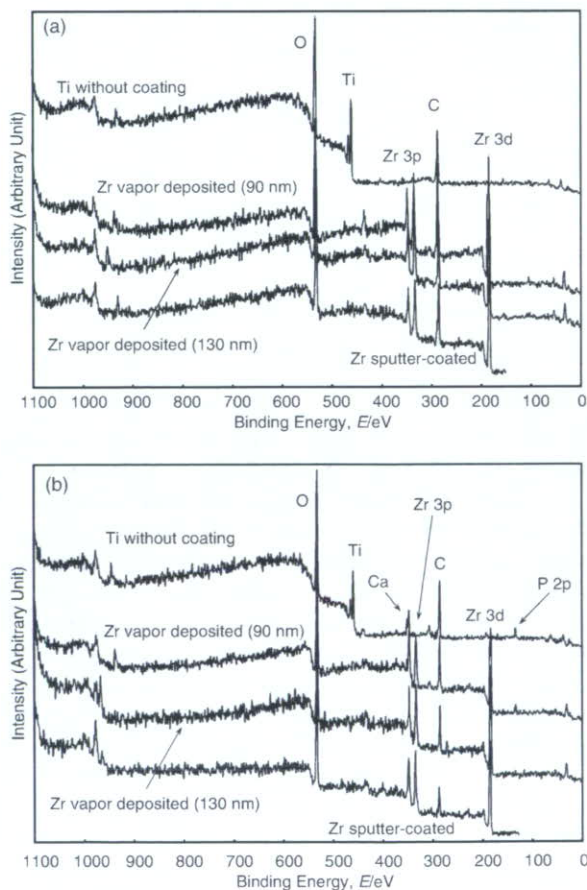


Fig. 3 Survey spectrum of specimens, (a) before and (b) after immersion into Hanks' solution for 1 d.

energies of potassium phosphates and sodium phosphates are varied from 132.7 eV to 133.9 eV, while those of several calcium phosphates, such as $\text{Ca}_{10}(\text{PO}_4)(\text{OH})_2$, $\text{CaHPO}_4 \cdot \text{H}_2\text{O}$ and $\text{Ca}(\text{H}_2\text{PO}_4)_2 \cdot \text{H}_2\text{O}$ are 133.2 eV to 134.3 eV.²⁾ Although it was difficult to deconvolute the spectra of P 2p into H_2PO_4^- , HPO_4^{2-} and/or PO_4^{3-} in the present study, the combined peak around 134 eV shows good correspondence to P 2p binding energies of calcium phosphates in the previous report. Doublet peaks of Ca 2p was identified as Ca 2p_{1/2} and Ca 2p_{3/2} referring to previous report.^{2,3)}

The peak of Ti was deconvoluted into Ti^0 from metallic substrate and cations having different valence (Ti^{2+} , Ti^{3+} , Ti^{4+}). On the other hand, only tetra-valent oxide was detected for Zr in the present study. According to previous report, peak from O 1s can be deconvoluted into O^{2-} , OH^- and H_2O .¹³⁾ They correspond to binding energy of hydroxyl, bound water or adsorbed water in the oxide layer and oxygen consisting oxide, respectively. Because phosphate groups were detected in Ti without Zr coating specimen together with only cation Ca^{2+} , the particles seen in SEM observation were concluded as calcium phosphates.

Quantitative analysis based on the peak deconvolution shown in Fig. 4 was carried out. Concentration of P and Ca taken from specimen after 1-d immersion in Hanks' solution is shown in Fig. 5. In the Ti without Zr coating and the 90-nm thick Zr deposited specimen, approximately 3 mol% of P and Ca was detected, while 1.5 mol% of P and no Ca was exist on the specimen the 90 nm thick Zr deposited and that Zr sputter-coated. It is considered that this difference was presumably caused from imperfection of coated layer.

The ratio, $[\text{Ca}]/[\text{P}]$, on the Ti without Zr specimen,

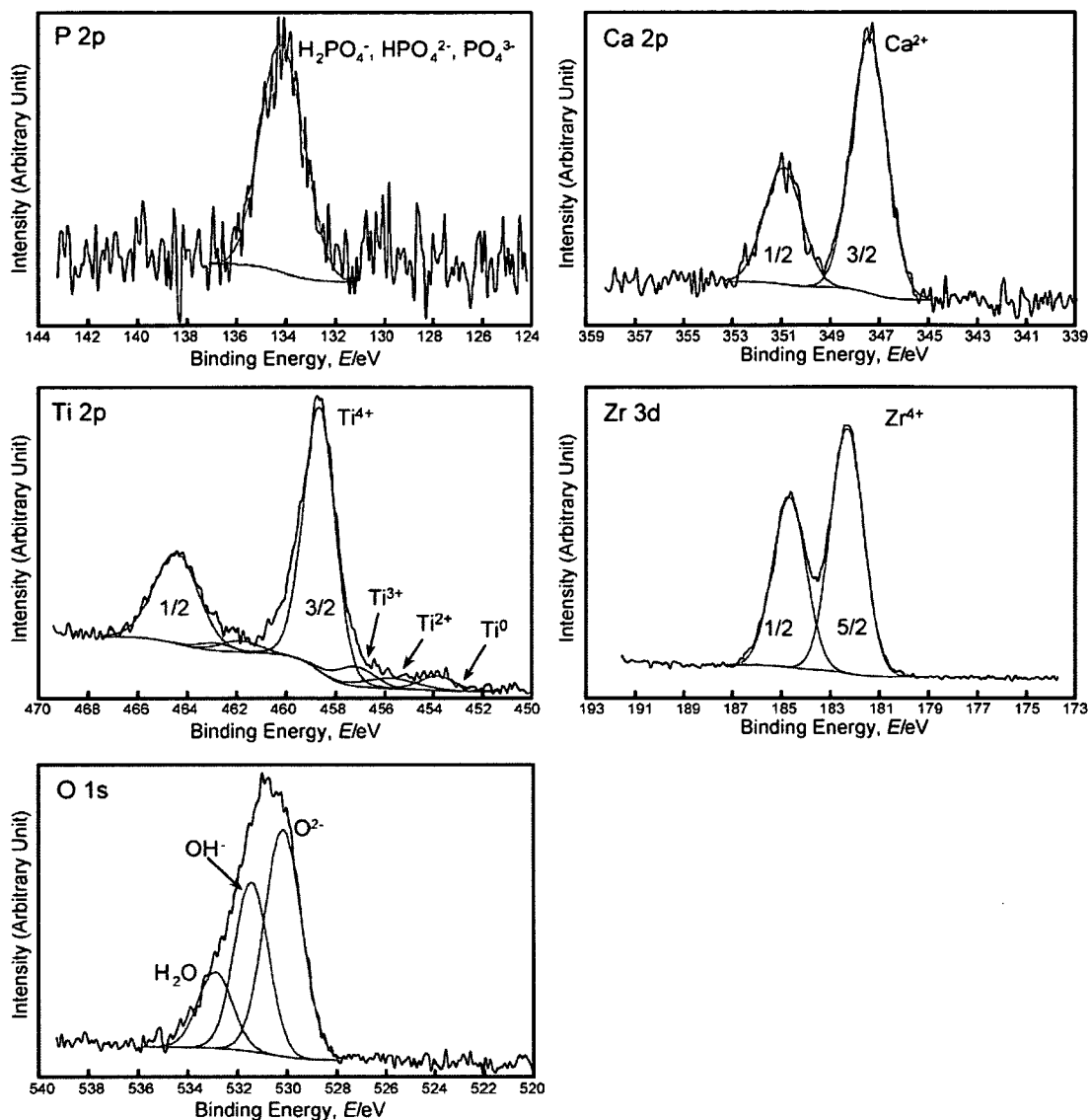
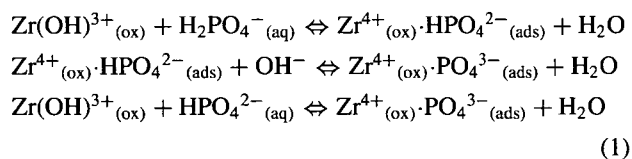


Fig. 4 Examples of peak deconvolution of narrow area spectrum of XPS analysis for P, Ca, Ti, Zr and O. All of the example spectrum are taken from specimen of Ti without Zr coating after immersion, except Zr spectra which is taken from that of Ti with Zr vapor deposited with 130-nm coating thickness.

calculated from above results was about 1.2. It is slightly smaller than that of hydroxyapatite ($\text{Ca}_{10}(\text{PO}_4)_6(\text{OH})_2$), which is known to 1.67, and rather nearer to that of $\text{Ca}_8(\text{HPO}_4)_2(\text{PO}_4)_4$, which is 1.33.³⁾

According to the previous report, zirconium phosphate was observed on the pure Zr (purity > 99.9%) surface instead of calcium phosphate.¹⁴⁾ It is reported that phosphate ion adhesion was occurred by dehydration and oxidation as follows,



Suffixes of ox, aq and ads correspond to ions in oxide, aqueous solution and adhesion respectively. After these reactions, these phosphate ions were involved in oxide layers

as forms of PO_4^{3-} , HPO_4^{2-} and $\text{H}_2\text{PO}_4^{-}$. According to the report, on the Ti surface, the adhesion of phosphate ions and Ca ion occurs alternately resulting in calcium phosphate precipitation. On the other hand, on the Zr surface, Ca ion adhesion is obstructed due to an interaction between the oxide and Ca ion, zirconium phosphate can be precipitated. In the present study, spectra of P was observed on the Zr sputter-coated specimens, which are considered to maintain the most fixed Zr layers, no spectra of Ca was detected.

As shown in Fig. 5, approximately 80% of atoms were O in the general results of quantity analysis by XPS. Ratio of the P and Ca is already stated above. N, Na and Si are considered as surface contamination.

According to the results of peak deconvolution of O spectra into O^{2-} , OH^{-} , and H_2O , which correspond to hydroxyl adsorbed on surface oxide, water molecules mainly adsorbed on oxide surface, and oxygen in the titanium oxide (TiO_2) and/or zirconium oxide (ZrO_2) respectively, state

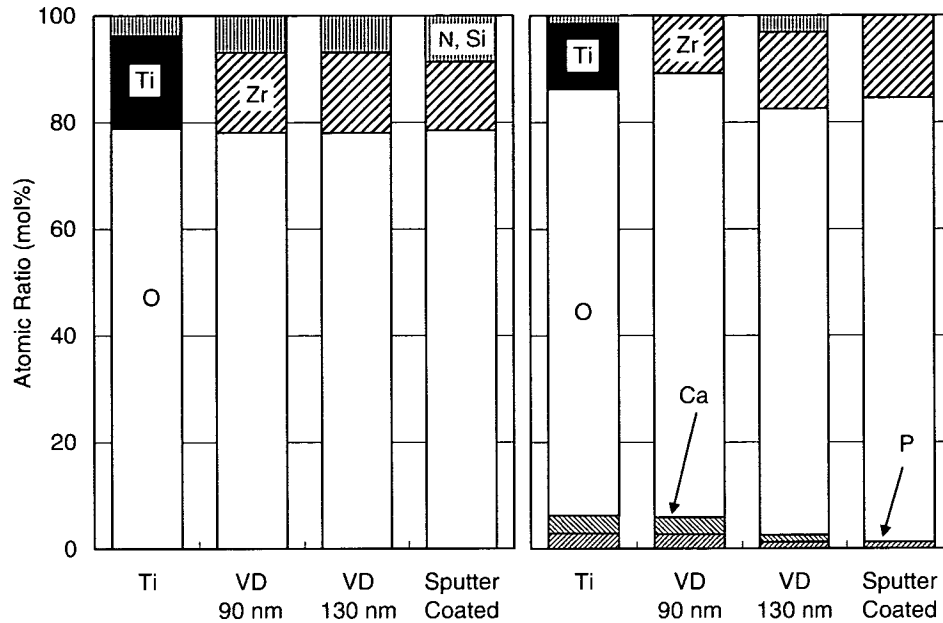


Fig. 5 Atomic ratio of specimens (left) before and (right) after immersion in Hanks' solution for 1 d. Ti: Ti without Zr coating, VD: vapor deposited. N and Si were contamination atoms.

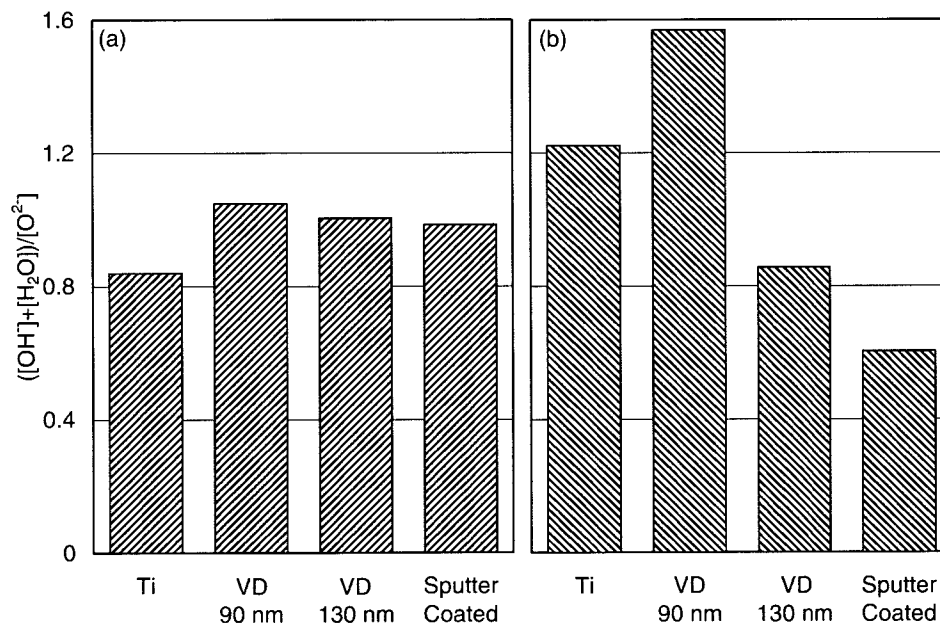
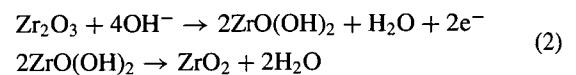


Fig. 6 $([\text{OH}^-] + [\text{H}_2\text{O}])/[\text{O}^{2-}]$ ratio of specimens (a) before and (b) after immersion in Hanks' solution for 1 d. Ti: Ti without Zr coating, VD: vapor deposited.

analysis was carried out. Figure 6 indicates ratio of oxygen on the oxides surface (OH^- and H_2O) for that in the oxides (O^{2-}). From the surface analysis before Hanks' solution immersion, the specimens Zr vapor deposited and that Zr sputter-coated showed higher $([\text{OH}^-] + [\text{H}_2\text{O}])/[\text{O}^{2-}]$ value than Ti without Zr. Comparing the $([\text{OH}^-] + [\text{H}_2\text{O}])/[\text{O}^{2-}]$ value before and after immersion, $([\text{OH}^-] + [\text{H}_2\text{O}])/[\text{O}^{2-}]$ value increased for the Ti specimen without Zr and the specimens Zr vapor deposited (90-nm thick), while the value decreased for the specimens Zr vapor deposited (130-nm

thick) and Zr sputter-coated.

If once Zr is immersed into aqueous solution, following reactions might occur on the surface.³⁾



According to these reactions, oxides film grows (which results in increasing of $[\text{O}^{2-}]$) and the $([\text{OH}^-] + [\text{H}_2\text{O}])/[\text{O}^{2-}]$ value will be decreased. Decreasing in the specimens Zr deposited (130-nm thick) and Zr sputter-coated

can be understood by this description. On the other hand, the $([\text{OH}^-] + [\text{H}_2\text{O}])/[\text{O}^{2-}]$ value in the Ti specimen without Zr and the specimen Zr deposited (90-nm thick) was increased after immersion into Hanks' solution. It is considered due to calcium phosphate formation on these specimens. As shown in equations (1), hydroxyl group contributes to reactions for calcium phosphates. In the recent reports, it was also reported that there is a certain relation between quantity of hydroxyl and crystallization of calcium phosphate.⁶⁾

4. Conclusion

On the surface of Ti without Zr, particle precipitates were observed and they were analyzed as calcium phosphate. Calcium phosphate was also identified on the specimens with 90-nm thick Zr coated layer. On the other hand, the specimens with 130-nm thick Zr coated layer exhibited control of calcium phosphate precipitation on the surface. By the surface analysis of the specimens Zr sputter-coated, Ca was not identified. It was considered to form zirconium phosphate instead of calcium phosphate.

From above results, efficiency of vapor deposited coating of Zr on inhibiting calcium phosphate in the body fluid was confirmed. It will be more effective when the thickness of the coated layer becoming thicker. Within the XPS results in the present study, it was confirmed that the sputter coating which provides thicker layer can prevent calcium phosphate precipitation. It was suggested that this technique can be applied to orthopedic devices such as femoral, tibial and

humeral nails and bone screws to avoid re-fracture of bone by calcium phosphate precipitation during removal operation.

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Characterization of calcium titanate thin films deposited on titanium with reactive sputtering and pulsed laser depositions

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Abstract

In the present study, characteristics of calcium titanate thin films deposited on titanium by reactive sputtering and pulsed laser deposition techniques were investigated. In both techniques, a calcium titanate target was used as a deposition source, and the titanium substrate was heated at 873 K during the deposition. The oxygen flow for the reactive sputtering was in the range of 1 to 10 sccm, and the oxygen pressure for the pulsed laser deposition was in the range of 0.13 to 13 Pa. The deposited films were crystallized into perovskite-type calcium titanate; furthermore, a titanium-dioxide layer formed in the interface between the film and substrate. In the film deposited by reactive sputtering with low oxygen flow, titanium-to-calcium ratio ([Ti]/[Ca]) is lower than that of stoichiometric calcium titanate due to the formation of calcium hydroxide. The ratio increases with an increase of oxygen flow, and the ratio of the film deposited with a 10-sccm oxygen flow was almost in accordance with that of stoichiometric calcium titanate. On the other hand, in the pulsed laser deposition, [Ti]/[Ca] ratios of the deposited film were almost in accordance with that of stoichiometric calcium titanate at the oxygen pressure under the present experimental condition. In both deposition techniques, the thickness of the titanium-oxide layer increased with an increase of the amount of oxygen gases. The results indicate that the pulsed laser deposition has an advantage for the preparation of the stoichiometric calcium titanate film without formation of a thick titanium-oxide layer.

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Keywords: Film characterization; Calcium titanate film; Reactive sputtering deposition; Pulsed laser deposition

1. Introduction

One of the purposes of surface modifications for biomaterials is to improve tissue responses in a living body because tissue-biomaterial reactions are interfacial phenomena which are governed by surface properties of the biomaterial. Ceramic coatings are often applied to facilitate osteogenesis on metallic biomaterials. Among ceramics, hydroxyapatite (HAP) is the most popular coating material [1–5]. Many researchers have demonstrated good osteogenesis on HAP-coated metals [6–9], and HAP-coated titanium prepared with a plasma-spraying process has been used clinically [10–12]. However, fractures at the HAP-

titanium interface and in the HAP layer itself are often observed after long-term use in the human body [13]. Accidents caused by these fractures result in a loss of the biomaterial-bone fixation. Consequently, clinical use of the HAP-coated titanium has decreased in recent years.

Recently, some of the present authors succeeded in developing a bioactive calcium titanate (CaTiO_3) film which can activate osteogenesis on titanium [14–17]. The bioactive CaTiO_3 film was prepared by radiofrequency (RF) magnetron sputtering with a CaTiO_3 target in an argon atmosphere and post-annealing at 873 K in air [16,17]. The prepared film was crystallized into perovskite-type CaTiO_3 , and the chemical composition of the film was almost in accordance with that of stoichiometric CaTiO_3 . A remarkable feature of the bioactive CaTiO_3 film was that the thickness was about 50 nm [17]. The thickness was 1/1000 that of plasma-sprayed HAP coating. This thickness made it possible to improve the mechanical strength

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