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## Morphological and Surface Potential Characterization of Protein Nanobiofilm Formation on Magnesium Alloy Oxide: Their Role in Biodegradation

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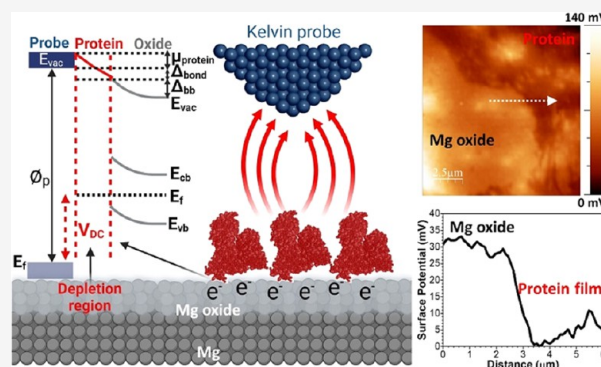
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**ABSTRACT:** The formation of a protein nanobiofilm on the surface of degradable biomaterials such as magnesium (Mg) and its alloys influences metal ion release, cell adhesion/spreading, and biocompatibility. During the early stage of human body implantation, competition and interaction between inorganic species and protein molecules result in a complex film containing Mg oxide and a protein layer. This film affects the electrochemical properties of the metal surface, the protein conformational arrangement, and the electronic properties of the protein/Mg oxide interface. In this study, we discuss the impact of various simulated body fluids, including sodium chloride (NaCl), phosphate-buffered saline (PBS), and Hanks' solutions on protein adsorption, electrochemical interactions, and electrical surface potential (ESP) distribution at the adsorbed protein/Mg oxide interface. After 10 min of immersion in NaCl, atomic force microscopy (AFM) and scanning Kelvin probe force microscopy (SKPFM) showed a higher surface roughness related to enhanced degradation and lower ESP distribution on a Mg-based alloy than those in other solutions. Furthermore, adding bovine serum albumin (BSA) to all solutions caused a decline in the total surface roughness and ESP magnitude on the Mg alloy surface, particularly in the NaCl electrolyte. Using SKPFM surface analysis, we detected a protein nanobiofilm (~10–20 nm) with an aggregated and/or fibrillary morphology only on the Mg surface exposed in Hanks' and PBS solutions; these surfaces had a lower ESP value than the oxide layer.



### 1. INTRODUCTION

Protein adsorption on the surface of all medical biomaterials occurs quickly during transplantation into human physiological media.<sup>1,2</sup> The distribution and conformational arrangement of protein nanobiofilms directly influence subsequent biological events, including cellular attachment, prefiltration and migration,<sup>3,4</sup> inflammatory responses,<sup>5</sup> and metal ion release (e.g., corrosion and biodegradation processes).<sup>6–8</sup> It is known that the adsorption of protein molecules on biomaterial surfaces is a complex process that involves different interactions at the protein molecule/solid interface, including hydrophobic, electrostatic, van der Waals, and hydrogen bonding forces.<sup>1,9</sup> These interactions depend on the physical and chemical properties of the biomaterial (electronic properties, chemical composition, wettability, surface defects, surface charge, roughness).<sup>10,11</sup> Furthermore, some physiological factors, including pH, ionic strength, temperature, and mass transfer,<sup>12</sup> can considerably affect the protein molecular structure (charge and hydrophobicity)<sup>13</sup> and then control its adsorption mechanism.

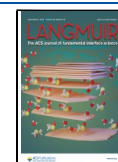
The adsorption of proteins onto Mg and its alloys is a complicated process because multichemical and electro-

chemical processes occur simultaneously. These include metal ion release,<sup>14</sup> adsorption of inorganic species ( $\text{Ca}^{2+}$ ,  $\text{HPO}_4^{2-}$ ,  $\text{H}_2\text{PO}_4^-$ , and  $\text{HCO}_3^-$ ),<sup>15,16</sup> and formation of complex corrosion products.<sup>17,18</sup> Mg and Mg alloys are promising materials for either macro- or miniscale medical devices<sup>19,20</sup> because they have superior biocompatibility<sup>21</sup> and bioresorbability or biodegradability,<sup>22,23</sup> alongside appropriate mechanical properties.<sup>24</sup> Nevertheless, these metals interact with protein molecules and require special attention when considering their corrosion and biodegradation behavior.<sup>25,26</sup> It is reported that depending on their type, proteins can inhibit or accelerate (detrimental impact) Mg degradation processes.<sup>22</sup> This interaction may also be time-dependent. For example, in a study on the corrosion of a Mg rare-earth (RE)

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**Table 1. Chemical Composition and Parameters of the NaCl, PBS, and Hanks' Solutions**

solution	CaCl <sub>2</sub> ·2H <sub>2</sub> O (g/L)	MgSO <sub>4</sub> (g/L)	KCl (g/L)	KH <sub>2</sub> PO <sub>4</sub> (g/L)	NaHCO <sub>3</sub> (g/L)	NaCl (g/L)	Na <sub>2</sub> HPO <sub>4</sub> (g/L)	pH	temperature (°C)
PBS			0.2	0.2		8	1.15	7.4	37
Hanks	0.185	0.097	0.4	0.06	0.35	8	0.047	7.4	37
NaCl						9		7.4	37

alloy, bovine serum albumin (BSA) initially inhibited corrosion and then accelerated the corrosion rate over longer-term immersion.<sup>27</sup> In addition, higher protein concentrations (1 g L<sup>-1</sup>) significantly inhibited the corrosion process of a Mg alloy when compared to lower concentrations (0.01 g L<sup>-1</sup>).<sup>28</sup> Indeed, the type of protein binding on the Mg oxide layer and the rearrangement of the adsorbed protein or even its detachment process in the form of metal–protein complexes impact Mg and Mg alloy biodegradation.<sup>29,30</sup>

Proteins are nanosized biological species. They are considered to be electrically conductive soft materials, depending on their molecular structure.<sup>31</sup> The specific electrical conductivity (EC) of adsorbed protein nanobiofilms on biomaterial surfaces can remarkably influence other biological events, particularly electrochemical interactions and metal ion release at the protein/oxide interface.<sup>10,32</sup> An extensive range of experimental approaches has been utilized to analyze the conductivity of biological species with a nanometer-scale structure, such as DNA and proteins. These techniques demonstrate valuable physical and chemical information regarding the EC of adsorbed DNA or protein molecules under both *ex situ* (in vacuum or air) and *in situ* (electrolyte media) conditions. The range of valuable techniques includes sandwiching proteins between two solid electrodes,<sup>33,34</sup> cyclic voltammetry,<sup>35,36</sup> electrochemical impedance spectroscopy (EIS),<sup>37,38</sup> scanning ion conductance microscopy (SICM),<sup>39</sup> scanning tunneling microscopy (STM),<sup>40,41</sup> current-sensing atomic force microscopy (CS-AFM),<sup>42,43</sup> and scanning Kelvin probe force microscopy (SKPFM).<sup>10,44</sup> Among the various localized-scale characterization procedures used to determine the EC of nanosized biological species, SKPFM is unique because it has high surface sensitivity.<sup>45</sup> This technique can analyze the local ( $\mu\text{m}$ – $\text{nm}$ ) electrical surface potential and/or surface charge distribution on small-scale simple or complex devices with diverse medical applications, including biosensors,<sup>46–48</sup> drug design,<sup>46,49,50</sup> medical implants,<sup>51</sup> and magnetic microrobots.<sup>52</sup>

The high spatial resolution and substantial surface sensitivity of SKPFM to any chemical and physical alterations provide essential information about the protein nanobiofilm/metal or oxide interface, including adsorption morphology, electrical surface potential, and predictive insights into electrochemical interactions.<sup>9,32</sup> This specific information supported by the SKPFM approach is vital in the case of Mg and Mg alloy bioactive surfaces because they have a high rate of degradation and form complex corrosion products. Furthermore, particularly at the early stages of immersion in human body media, the morphology of the adsorbed protein and its electrical surface potential distribution can directly influence cell adhesion/deformation,<sup>53</sup> the osseointegration process,<sup>54</sup> and the long-term durability of implanted biomaterials.<sup>26</sup>

In this study, we used SKPFM to visualize the adsorption and formation of a BSA protein nanobiofilm and its electrical surface potential on the surface of the WE43 Mg alloy in three different simulated solutions, including 0.9% NaCl, phosphate-buffered saline (PBS), and Hanks' solution. In addition,

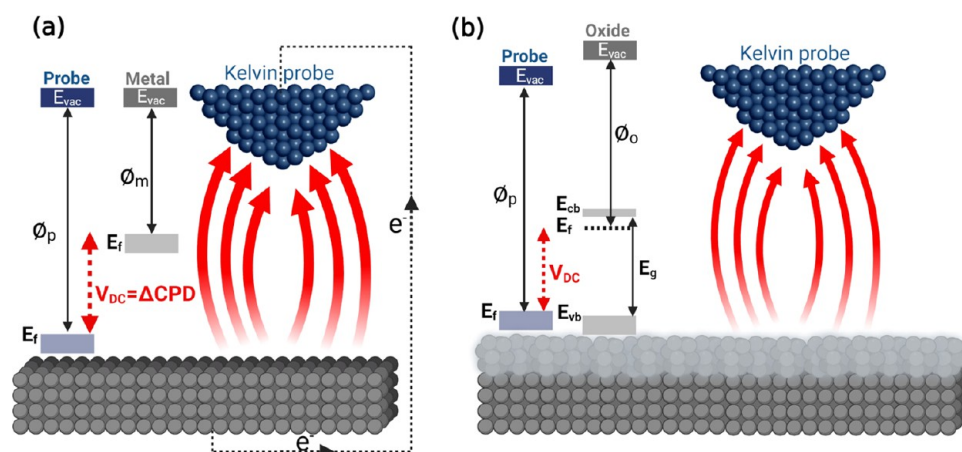
complementary electrochemical measurements, X-ray photoelectron spectroscopy (XPS), and scanning electron microscopy (SEM) were used to reveal further information regarding the role of the BSA protein on electrochemical interactions and surface chemical and microstructural evolutions on the Mg alloy in various simulated human body environments.

## 2. EXPERIMENTAL PROCEDURE

**2.1. Sample Preparation.** Specimens with surface areas of 1 cm<sup>2</sup> were cut from a bar of WE43 magnesium (Mg). Mg alloys containing the rare-earth (RE) elements, such as WE43 alloy, represent improved mechanical properties (tensile strength and creep) at ambient and high temperatures alongside adequate corrosion resistance. Both properties are indispensable factors for an appropriate biomedical material. The chemical composition (atom %) of the WE43 Mg alloy (91.67 Mg, 3.87 Y, 2.18 Nd, 0.91 Zr, and 1.37 RE) was determined using inductively coupled plasma optical emission spectrometry (ICP-OES) after multi-acid (HCl, HNO<sub>3</sub>, and HF in a molar ratio of 30:10:1) digestion. All samples were mechanically ground and polished to a mirror-like surface, washed with ethanol, ultrasonicated in acetone for 20 min, and dried by air blowing before surface characterization.

**2.2. Electrolyte and Electrochemical Measurements.** The alloy was cut into 10 × 10 × 10 mm cubes. Each cube was glued to a copper wire using conductive epoxy and then mounted in non-conductive transparent epoxy resin such that only a flat surface of known area (1 cm<sup>2</sup>) would be exposed to the three different simulated body fluids, including 0.9% NaCl, phosphate-buffered saline (PBS, ASTM Standard (F2129)<sup>55</sup>), and Hanks' (according to H 8264 (without glucose), Sigma-Aldrich) solutions, as shown in Table 1. To investigate the BSA protein's role in electrochemical interactions, surface roughness, and electrical surface potential events on the WE43 Mg alloy, 4 g L<sup>-1</sup> BSA protein (lyophilized powder; ≥96% agarose gel electrophoresis, Sigma-Aldrich) was added to the examined solutions by adjusting the pH (pH meter, GLP 21 CRISON) to 7.4 ± 1 and temperature to 37 ± 1 °C. Albumin proteins are commonly found in blood plasma and are unique among the major plasma proteins in containing no carbohydrate residues.<sup>6</sup> Human serum albumin (HSA) is the most abundant protein in human blood plasma, which is mainly responsible for the maintenance of blood pH and osmotic pressure.<sup>6,56,57</sup> Since BSA and HSA are homologous proteins having a similar sequence and conformation (heart-shaped that formed by three homologous domains (I, II, and III)), we used BSA for this study, which is consistent with numerous previous investigations.<sup>11,53,58,59</sup> The electrochemical measurements were carried out using an AUTOLAB PGSTAT302 potentiostat in a conventional three-electrode electrochemical cell with Ag/AgCl/KCl<sub>sat</sub> (+219 mV vs SHE), a platinum wire, and the specimen as the reference, counter, and working electrodes, respectively. All electrochemical analyses were performed after 10 min of immersion in the solution to stabilize the open circuit potential (OCP) and reach the steady-state condition. The potentiodynamic polarization (PDP) measurements were performed at a scan rate of 1 mV s<sup>-1</sup> from -1.75 V (cathodic branch) to -1.05 V (anodic branch).

**2.3. Chemical Surface Characterization by XPS.** The chemical composition of the WE43 surface layer was analyzed using a Kratos Analytical Axis ULTRA spectrometer containing a DLD spectrometer using a monochromatic aluminum source (Al K $\alpha$ , 1486.6 eV) operating at 150 W (10 mA emission current and 15 kV HT). Analysis was conducted on a 700 × 300  $\mu\text{m}^2$  area of the sample. Survey scans were obtained at a 1 eV step size and a pass energy of



**Figure 1.** Schematic illustration of the SKPFM principle alongside the energy level diagram during the electrostatic interaction between a conductive AFM tip apex and (a) metal or (b) semiconductor oxide bulk materials at the atomic scale.

160 eV and averaged over two scans using Vision Processing software by Kratos Analytical. The kinetic energy of the photoelectrons was measured at a 90° take-off, and the vacuum in the analysis chamber was approximately  $5 \times 10^{-10}$  torr. In addition, the proportion of protein adsorption on the surface of the WE43 Mg alloy in different environments was evaluated by comparing the relative atomic ratio between N (N 1s) and the oxidized carbon C 1s peaks.<sup>56</sup>

**2.4. Microstructure Characterization by SEM/EDXS and AFM/SKPFM.** Protein adsorption and its impact on surface topography and the electrical surface potential distribution of the WE43 Mg alloy were studied using combined SEM, AFM, and SKPFM measurements. The microscopy observations were conducted on as-polished (as reference sample) and also immersed samples (for 10 min) in the different simulated physiological solutions (NaCl, PBS, and Hanks') with or without the addition of the BSA protein. SEM was performed on a JSM-7610FPlus instrument (JEOL) equipped with an Oxford X-MAX20 energy-dispersive X-ray spectrometer (EDXS). All SEM images were recorded at a working distance of 15 mm, an accelerating voltage of 5 kV, and secondary electron (SE) mode. The AFM/SKPFM surface analyses were performed by a Nanoscope IIIa Multimode device with an n-type doped silicon pyramid single-crystal tip coated with PtIr5 (SCM-Pit probe, tip radius, and heights were 20 nm and 10–15 μm, respectively). The surface potential maps were recorded in a dual-scan mode. In the first scan, surface topography maps were captured using tapping mode. The tip was lifted to 100 nm in the second scan, and the potential signal was recorded by following the topography contour registered in the first scan. All topographic and surface potential maps were obtained in an air atmosphere at 27 °C with an approximate relative humidity (RH) of 28%, a pixel resolution of  $512 \times 512$ , zero-bias voltage, and a scan frequency rate of 0.2 Hz.

**2.5. Energy Levels and SKPFM Surface Potential Signal at the Metal and Metal/Oxide Interface.** The SKPFM is capable of measuring the local contact potential difference (ΔCPD) between a conductive AFM tip and the studied sample, thereby imaging the electrical surface potential or work function energy (WFE) of the sample with high lateral resolution (micrometer to nano or sub-nanometer scale).<sup>60</sup> Therefore, any chemical and physical variations on solid surfaces (e.g., exposed metal, oxide, or soft matter) are strongly affected by WFE or local surface potential.<sup>61</sup> Figure 1a,b shows the fundamental principle of SKPFM analysis alongside energy diagrams between an AFM tip apex and the metal or oxide film/metal interface with relevant energy parameters including valence and conduction bands ( $E_{vb}$  and  $E_{cb}$ ), Fermi level ( $E_f$ ), band-gap energy ( $E_g$ ), and vacuum level ( $E_{vac}$ ). With a constant bias voltage and tip-substrate distance, the recorded ΔCPD value directly corresponds to WFE values between the probe ( $\phi_p$ ) and homogeneous or heterogeneous surfaces (metal ( $\phi_m$ ) or oxide ( $\phi_o$ )) under study. This relationship is given as follows<sup>10</sup>

$$\Delta CPD = \frac{(\phi_p - \phi_s)}{e} \quad (1)$$

where  $e$  is the elementary charge. As defined in eq 1 and Figure 1a, an appropriate value of the applied external bias ( $V_{DC}$ ) should be exerted to nullify the electrostatic force ( $F_{EF}$ ) during the electrical connection ( $V_{DC} = \Delta CPD$ ), since ΔCPD is equal to the WFE difference between the tip and the sample. The electrostatic force between the AFM tip and the sample surface is defined as follows<sup>60</sup>

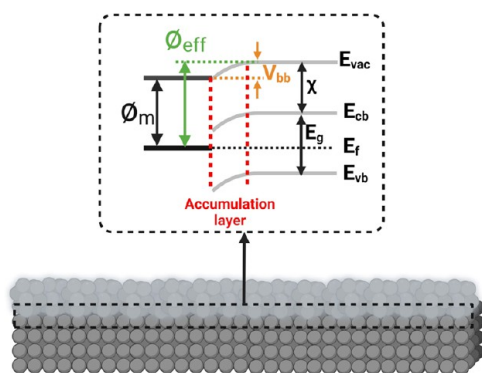
$$F_{EF}(z) = -\frac{1}{2} \Delta V^2 \frac{dC(z)}{dz} \quad (2)$$

where  $z$  is the direction normal to the surface of the studied sample,  $\Delta V$  is  $V_{DC} - \Delta CPD$ , and  $dC/dz$  is the capacitance gradient between the tip and the sample surface. The electrical forces between the AFM tip and the substrate in SKPFM analysis can be divided into capacitance forces due to surface potential and dielectric screening and Coulombic forces due to static charges and multipoles.<sup>62</sup> All of these factors directly influence the total WFE difference or ΔCPD values measured by SKPFM. In the case of complex systems, including metallic alloys and oxide films with various oxide compounds and/or heterogeneous distribution of n- or p-type semiconductors, the electrostatic interaction and local surface potential represent the concentration-weighted sum of the WFEs of the elements or oxide constituents ( $\phi_{total}$ ).<sup>10</sup> For example, in this study, we used the Mg–Y–Nd–Gd–Zr alloy for which  $\phi_{total} = \phi_{Mg} + \phi_Y + \phi_{Nd} + \phi_{Gd} + \phi_{Zr}$ . Additionally, in the case of a metallic surface or one covered by an oxide film (light gray color in Figure 1b), the electronic properties of the oxide film (e.g., n- or p-type semiconductor, band gap, dipoles, the position of occupied and unoccupied states, oxygen vacancies, cation interstitials, etc.) influence the magnitude of the local surface potential. Hence, we should consider the WFE of oxidation states that are higher than those obtained for the pure metal.<sup>63</sup> Notably, at the oxide/metal interface,  $E_{vac}$  on the oxide side increases to a higher value with slightly misaligning behavior (Figure 2). The degree of this misalignment (e.g., band bending value ( $V_{bb}$ ) at the metal/oxide interface) is dependent on the magnitude of the interfacial dipole moment. Consequently, a new term for WFE can be defined at metal/oxide interfaces, which is the effective work function energy ( $\phi_{eff}$ )<sup>64</sup>

$$\phi_{eff} = \chi + E_g + (E_f - E_{vb}) \quad (3)$$

where  $\chi$  is the electron affinity. Therefore, the total amount of ΔCPD measured by SKPFM on a metallic surface covered by a native oxide film is the mean value of  $\phi_o$  and  $\phi_{eff}$  (which, in turn, depends on the bias voltage and oxide thickness). This ΔCPD will be comparatively lower than that measured on a metallic surface, which will typically have a lower charge resistance value.





**Figure 2.** Schematic of the energy level alignment (band bending) in a metal/oxide interface.

**2.6. Protein Impact on Energy Levels and Surface Potential Signal on the Oxide Film.** When a solid surface is covered by a thin layer of external organic or inorganic materials, the WFE can alter to a new value due to electron transfer and structural relaxation at the interface.<sup>65</sup> Similar changes can occur in doped semiconductor surfaces due to band bending and the formation of depletion regions in the semiconductor subsurface.<sup>66</sup> With the adsorption and then the formation of a monolayer or multilayers of soft biological matter such as protein and DNA molecules on the surface of biomaterials, a new arrangement of the energy level is established at the biological molecule/oxide film interface (Figure 3a). As shown in Figure 3a, a monolayer of BSA protein molecules (in this study) adsorbed on a metal oxide film will affect the electrostatic interaction and capacitance magnitude due to changes in local WFE or ΔCPD between the AFM tip and the BSA-covered oxide layer. Indeed, the BSA protein–oxide film interactions meaningfully alter the electrostatic forces and ΔCPD owing to band bending (Δ<sub>bb</sub>, due to the formation of a depletion region), effective protein molecular dipole (μ<sub>BSA</sub>), and interfacial bond (Δ<sub>bond</sub>, due to the new arrangement of electron density at the protein/oxide film interface).<sup>67</sup> Therefore, the new local surface potential on the adsorbed protein–oxide complex (SP<sub>BSA–oxide</sub>) can be described as follows<sup>10,67</sup>

$$SP_{BSA-oxide} = SP_{oxide} + \mu_{BSA}/e + \Delta_{bond} \quad (4)$$

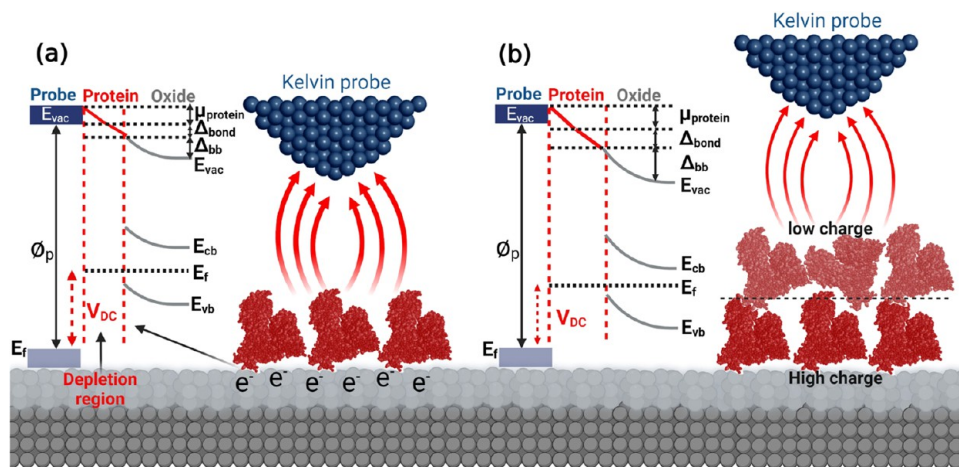
However, with increasing thickness of the adsorbed protein layer from a monolayer to multilayers on the oxide film surface (Figure 3b), the degree of the misalignment in energy levels increases (more band bending), resulting in a reduction of the intensity of electrostatic

forces and the ΔCPD value between the tip and the protein–oxide surface. The protein monolayer on the oxide film surface results in both μ<sub>BSA</sub> and Δ<sub>bond</sub> being charge transfer controlled and distributed across the protein/oxide film interface and through the protein nanobiofilm. Therefore, in the first layer or monolayer of the BSA protein, the charge distribution at both the protein/oxide film interface and on the surface of the protein monolayer (dark red color in Figure 3b) is more significant than for the second or subsequent layers of the adsorbed protein (light red color in Figure 3b).<sup>13</sup> It is worth noticing that the electrical charge transport function in the BSA molecule structure is lower than those for other proteins such as Azuring and bacteriorhodopsin, which considerably influences the electron transfer process in protein–protein interactions.<sup>59</sup> As reported in the literature,<sup>67,68</sup> for adsorbed organic films with thicknesses higher than 100 nm, the impact of the substrate (metal or oxide film in this study) can generally be neglected in the total amount of SP<sub>BSA–oxide</sub> because of limited electrostatic interactions between the tip and the studied surface.

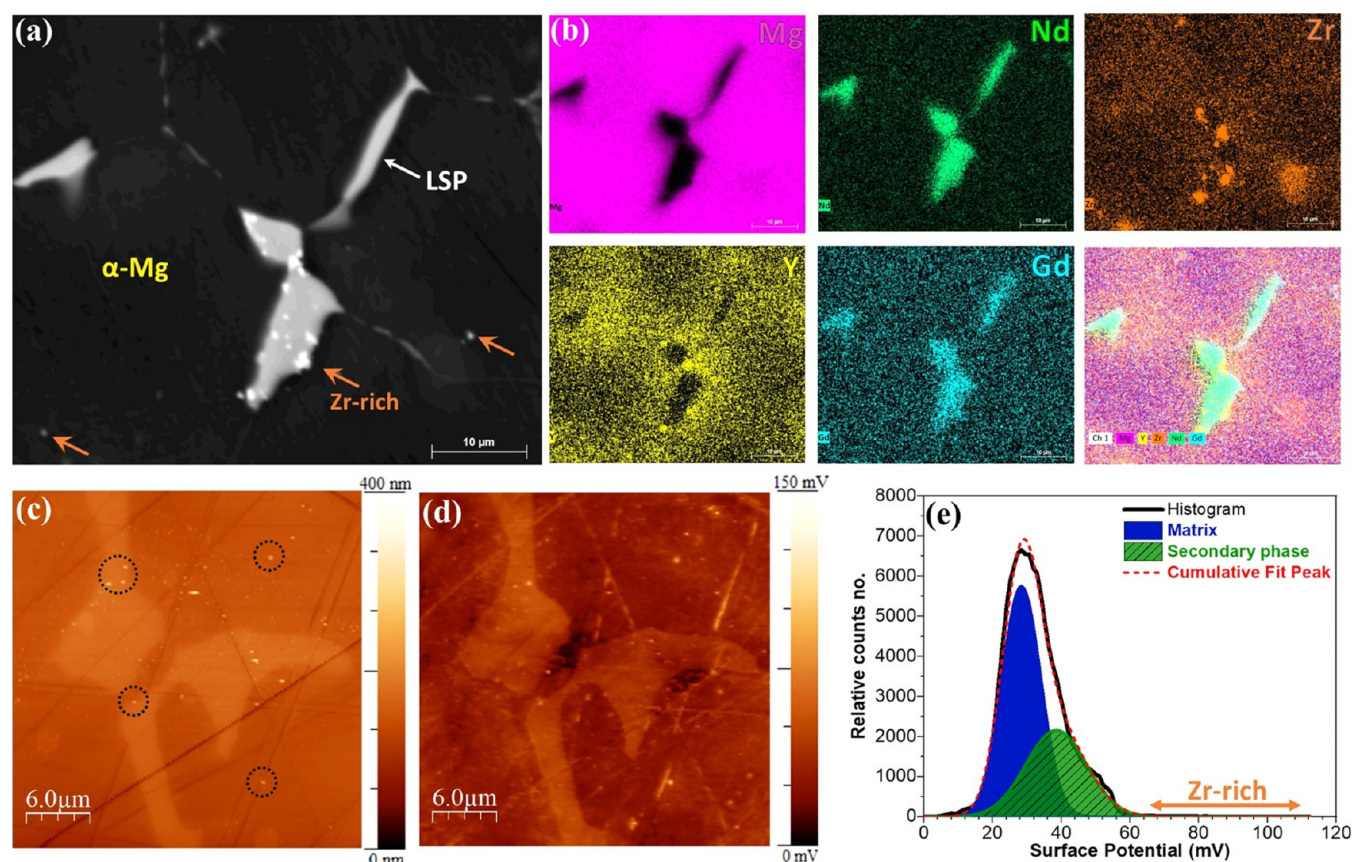
### 3. RESULTS AND DISCUSSION

**3.1. Microstructural and Surface Potential Characterization of the Mg Alloy.** SEM–EDXS measurements were used to reveal microstructural and elemental surface distribution on the different phases of Mg WE43. As presented in the SEM image and its EDXS corresponding elemental maps in Figure 4a,b, Mg WE43 is composed of three individual regions, including the matrix or α-Mg, a large secondary phase (LSP), and a Zr-rich phase (the chemical composition of the individual phases is reported in Table 2). The LSPs are discretely distributed around the matrix grain boundaries, which are more enriched in Nd and Gd. The Zr-rich phases are mainly precipitated at the LSP/α-Mg boundary and slightly dispersed in α-Mg. Y is more distributed around the LSPs than in their bulk region and is precipitated primarily in Zr-rich phases. These nonhomogeneous elemental distributions in the individual phases and their boundaries can directly influence the material's electronic properties, such as its electrical surface potential signal.<sup>10</sup>

The topography and surface potential maps of the three previously mentioned regions were obtained by AFM and SKPFM measurements (Figure 4c,d). Notably, the surface potential image shows different WFEs for separate phases. The LPS phases present a higher electrical surface potential than



**Figure 3.** Schematic representation of the SKPFM principle alongside the energy level diagram during the electrostatic interaction between a conductive AFM tip apex and (a) a monolayer or (b) multilayers of protein molecules on semiconductor oxide bulk materials at the atomic scale. The adsorbed protein molecules strongly influence the total surface potential difference.



**Figure 4.** (a) SEM image, (b) EDXS elemental maps, (c) AFM, and (d) SKPFM images of the as-polished Mg-based alloy. (e) The histogram of surface potential distribution on the surface of the as-polished Mg-based alloy obtained from (d).

**Table 2. Chemical Composition of the Different Individual Phases of the WE43 Mg Alloy**

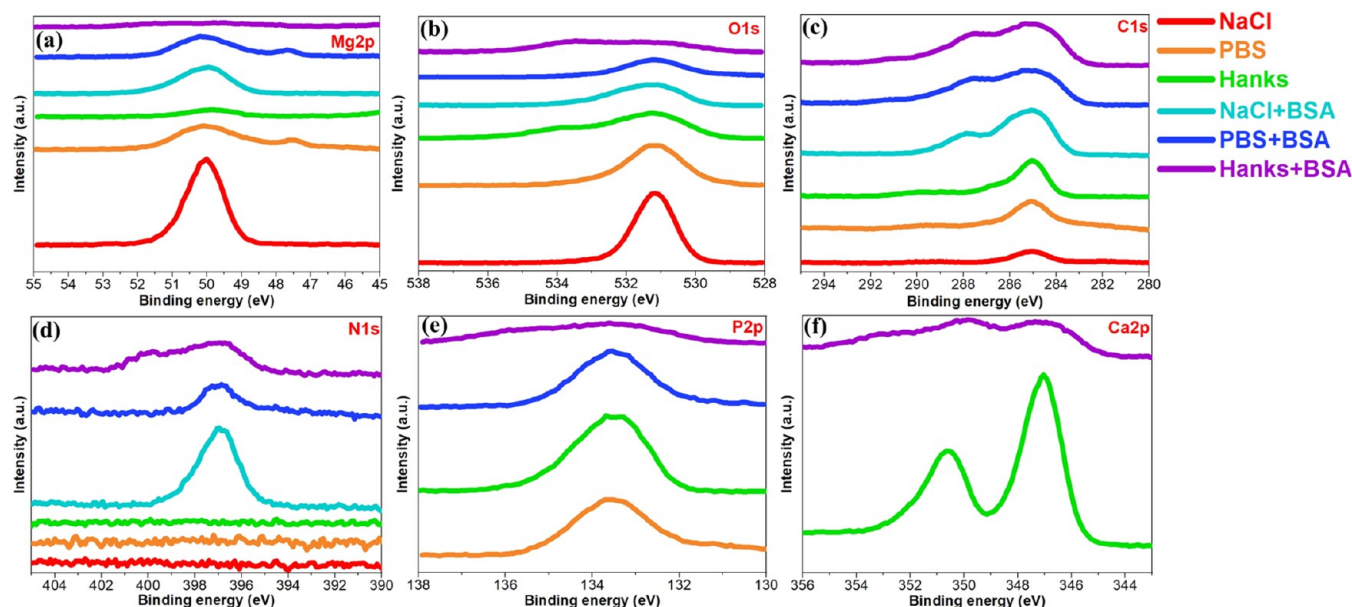
phase	chemical composition (atom %)				
	Mg	Y	Nd	Zr	Gd
$\alpha$ -Mg	96.77	1.43	0.49	1.04	0.27
$\beta$ -Mg	89.08	1.61	5.78	1.67	1.86
Zr-rich	74.16	6.13	2.64	15.83	1.24

the  $\alpha$ -Mg phases. Moreover, some bright spots with the highest potential signal are seen in the various regions, such as LPSs, LSP/ $\alpha$ -Mg, and  $\alpha$ -Mg. These are believed to be Zr-rich regions<sup>69</sup> (marked by dash circles). According to the literature<sup>10,63,70</sup> and Table 2, the various metal elements and their concentration in these individual phases can create different surface potential or WFE values. From the corrosion point of view, all these phases with unique electrical surface potentials, or WFEs, have different tendencies to transfer valence electrons for participation in electrochemical reactions at the metal or oxide film/electrolyte interface.<sup>71</sup> The microgalvanic driving force for corrosion can be forecasted based on the electrical surface potential for the various phases measured by SKPFM at the solid/air interface (histogram analysis in Figure 4e). Although the quick and spontaneous formation and then time-dependent structural alternation of the oxide film on the metal or alloy surface in aqueous media alongside the complexity of the environment<sup>6</sup> strongly affect the corrosion and biodegradation behavior of a simple or complex metallic system, in turn, they can make some

contradictions with SKPFM predication at the air/solid interface.<sup>72</sup>

**3.2. Evaluating Interactions of Inorganic Species and Protein with the Mg Alloy Surface by XPS.** A systematic surface chemical analysis of the Mg oxide film or corrosion products and BSA protein adsorption was carried out by XPS to reveal the role of the different simulated body fluids, including 0.9% NaCl, PBS, and Hanks' solutions with various inorganic species (mainly phosphate and calcium). The results of individual high-resolution spectra of elements are displayed in Figure 5. In Figure 5a, Mg 2p peaks occur from 46 to 52 eV. These are assigned to Mg (OH)<sub>2</sub> at 49.5 eV, MgO at 50.8 eV, and MgCO<sub>3</sub> at 51.8 eV.<sup>73</sup> A single peak at around 47.5 eV in PBS solution is attributed to the Mg metal. Normally, the MgO oxide peak appears on all Mg samples due to the formation of a thin MgO/Mg(OH)<sub>2</sub> film.<sup>74</sup> The O 1s spectrum has three individual peaks at 531.2, 531.9, and 533.8 eV, attributed to MgO, chemisorbed OH<sup>-</sup>, and H<sub>2</sub>O, respectively.<sup>75,76</sup> The main content of the C signal on all samples (more visible on surfaces without protein contact) originated from air contamination. According to the literature,<sup>2,77</sup> the molecular structure of the albumin protein is included in carboxyl (–COOH) groups, CO–NH peptides, and amino groups (–NH<sub>2</sub>). Therefore, three individual peaks can be deconvoluted in C 1s, including 284.8, 286.2, and 288.1 eV, which are attributed mainly to C–H and C–C bonds, C–O or peptidic residues, C–N bonds, and N–C=O bonds, respectively.<sup>2,30</sup> Moreover, C 1s peaks at around 289 to 290 eV reveal the presence of CO<sub>3</sub><sup>2-</sup> on the Mg alloy surface due to the formation of MgCO<sub>3</sub> and CaCO<sub>3</sub> in the corrosion/biodegradation

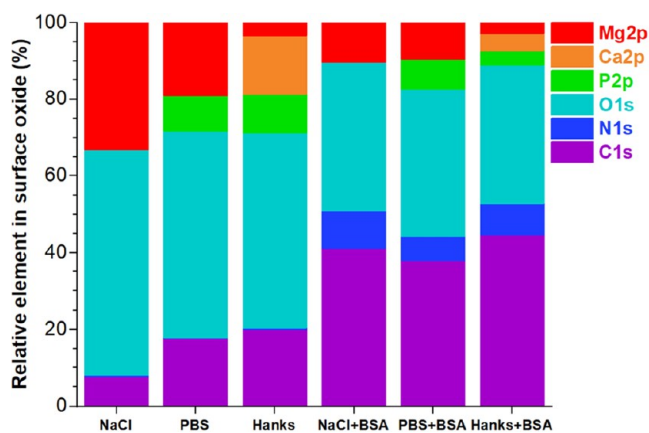




**Figure 5.** XPS spectra of (a) Mg 2p, (b) O 1s, (c) C 1s, (d) N 1s, (e) P 2p, and (f) Ca 2p electron energy regions on the WE43 Mg alloy after 10 min of immersion in 0.9% NaCl, PBS, and Hanks' solutions with or without the BSA protein ( $4 \text{ g L}^{-1}$ ) at pH 7.4 and  $37^\circ\text{C}$ .

products.<sup>74</sup> Hence, the higher intensity of C 1s peaks alongside the appearance of N 1s peaks (Figure 5d) in the oxide layer or corrosion products of the Mg alloy exposed to the albumin protein are related to protein adsorption and complex formation. The P 2p spectrum at around 133.5 eV is attributed to the formation of  $\text{Mg}_3(\text{PO}_4)_2$  in both PBS and Hanks' solutions and  $\text{Ca}_{10}(\text{PO}_4)_6(\text{OH})_2$  at 133.2 eV (Hanks' solution).<sup>73</sup> The Ca spectrum exhibits two individual peaks only in Hanks' solutions for  $\text{CaCO}_3$  at 347.1 eV and  $\text{Ca}_{10}(\text{PO}_4)_6(\text{OH})_2$  at 350.8 eV.<sup>73</sup>

Figure 6 presents the elemental distribution in the Mg surface oxide calculated from the XPS spectra. By changing the

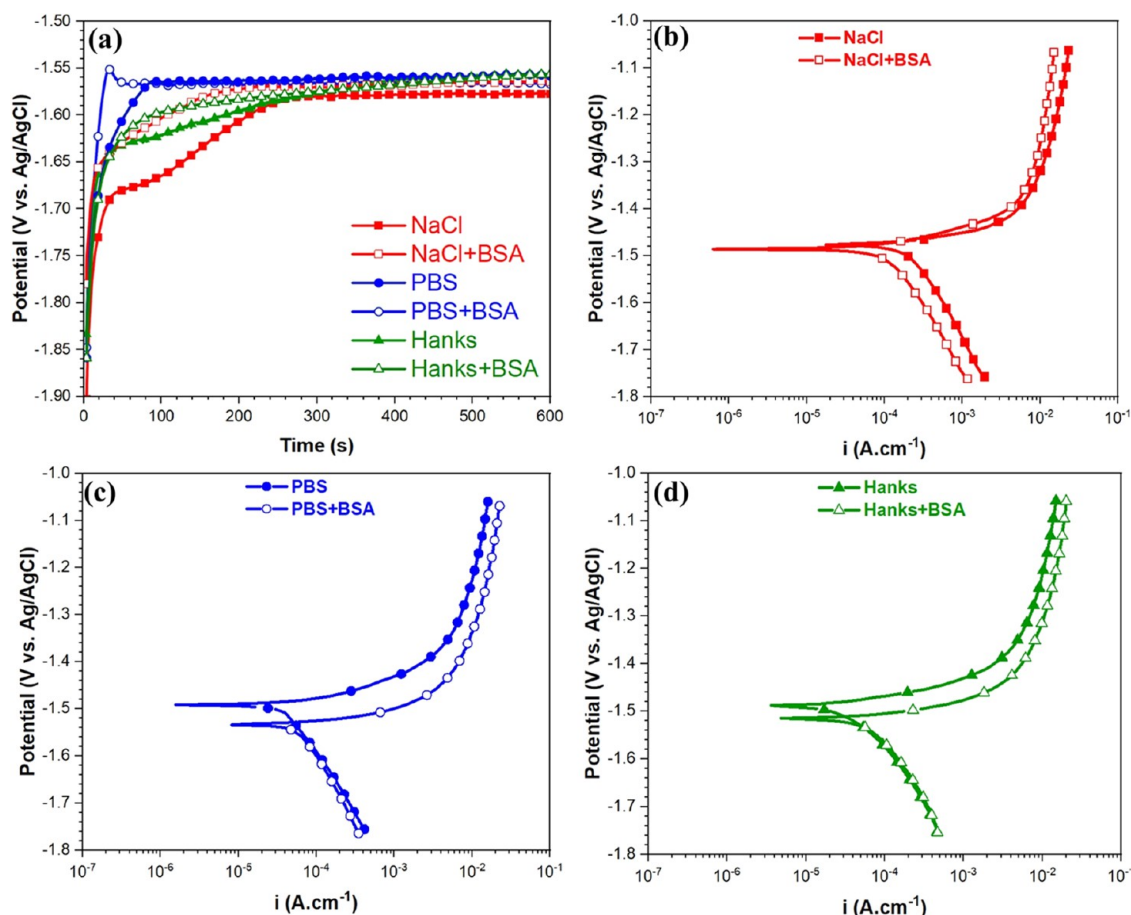


**Figure 6.** Relative percentage of elements in the surface oxide of the WE43 Mg alloy calculated from the XPS spectra in Figure 5.

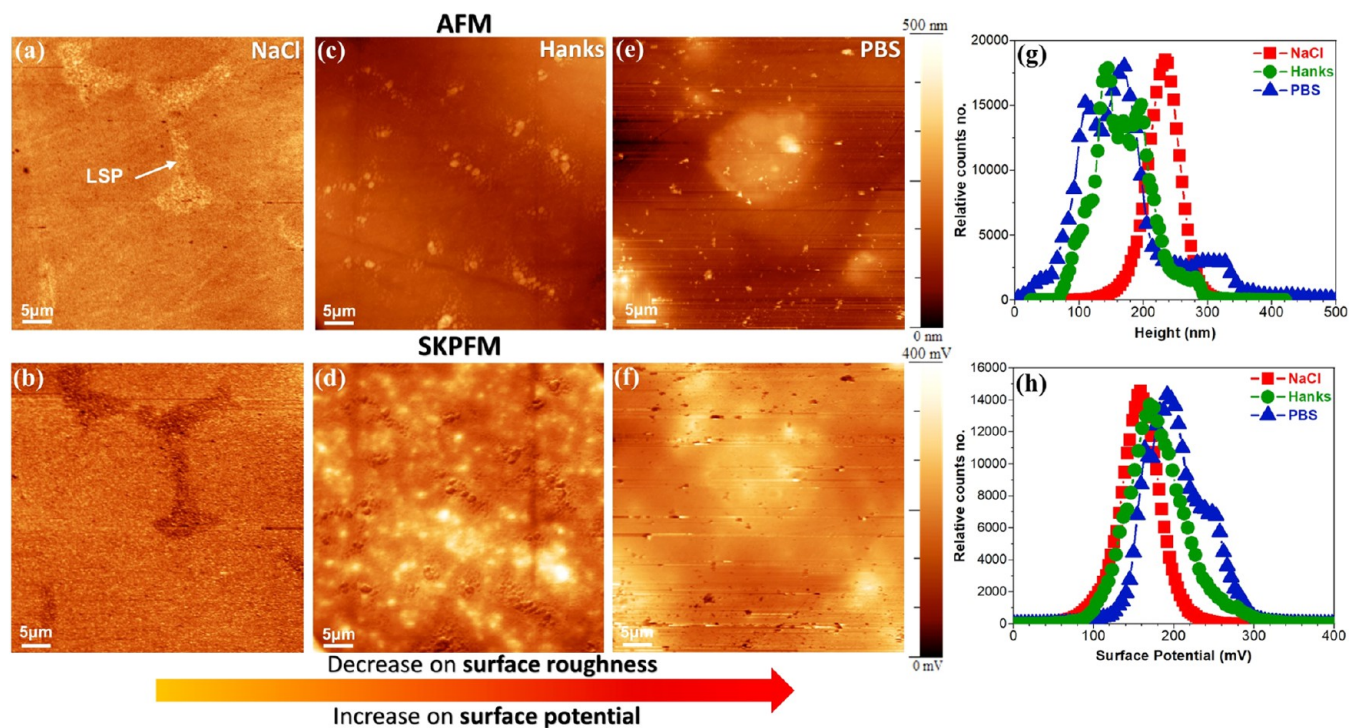
solution composition from NaCl to PBS and then Hanks' solutions, the Mg and O signal intensity slightly decreased in the oxide layer or corrosion products of the Mg alloy (Figures 5 and 6). Moreover, the oxide formed in PBS and Hanks' solutions comprises a high amount of P and Ca/P elements. The addition of the BSA protein in all solutions caused a decrease in the amount of Mg in the corrosion products. This was also true for the P content and the Ca/P ratio. These

parameters were lowest in the Hanks'+BSA solution. In the PBS media,  $\text{HPO}_4^{2-}$  and  $\text{H}_2\text{PO}_4^-$  have a high tendency to interact with the Mg oxide film to form a thin phosphate–magnesium complex ( $\text{Mg}_3(\text{PO}_4)_2$ ) as detected by XPS. It is reported that the acceleration/promotion of the formation of  $\text{Mg}(\text{OH})_2$  and  $\text{Mg}_3(\text{PO}_4)_2$  can strongly inhibit metal ion release and, in turn, enhance the corrosion resistance of Mg alloys,<sup>78</sup> which agrees with our electrochemical measurements in Figure 7. In Hanks' media, the preferential interaction of phosphate species with  $\text{Ca}^{2+}$  at near-neutral pH initially triggers the formation of  $\text{Ca}_{10}(\text{PO}_4)_6(\text{OH})_2$  (hydroxyapatite), which also inhibits corrosion (Figure 7).<sup>26</sup> Likewise, the unique interaction of  $\text{Ca}^{2+}$  and  $\text{HCO}_3^-$  species in the Hanks' environment can lead to the formation of  $\text{CaCO}_3$  products in the electrolyte, which may cover the surface of Mg WE43, causing the inhibition of the corrosion or biodegradation processes.<sup>78</sup> Nevertheless,  $\text{CaCO}_3$  products (and similarly  $\text{MgCO}_3$ ) have no significant inhibitory action due to their porous nature.<sup>78</sup>

A comparison of the atomic ratio of N (N 1s) to the oxidized carbon C 1s peaks  $[\text{N}/(\text{C}_2 + \text{C}_3)]$ <sup>56</sup> in all solutions containing the BSA protein indicates the amount of protein adsorption on the Mg oxide layer. This ratio increases in the order NaCl (0.238) > Hanks' (0.181) > PBS (0.163). The higher adsorption of the BSA protein on the Mg surface in the NaCl solution than that in the complex PBS and Hanks' environments is related to the competition between inorganic species and protein molecules.<sup>2</sup> Indeed, the tiny ionic size of inorganic species compared to the nanosize of the BSA protein makes them more mobile for interaction with the Mg oxide layer. In addition, the low rate of protein adsorption in PBS and Hanks' environments is due to the shielding or repulsing interaction of protein molecules (negative  $\zeta$  potential at pH 7.4) with negatively charged inorganic species such as  $\text{HPO}_4^{2-}$ ,  $\text{H}_2\text{PO}_4^-$ , and  $\text{HCO}_3^{2-}$ . Considering a very similar ionic strength of the different simulated body fluids (0.15–0.17 M), the relatively minor variation in ionic strength that occurs in response to changes in ion concentration can be considered negligible. Indeed, the complex film of inorganic species

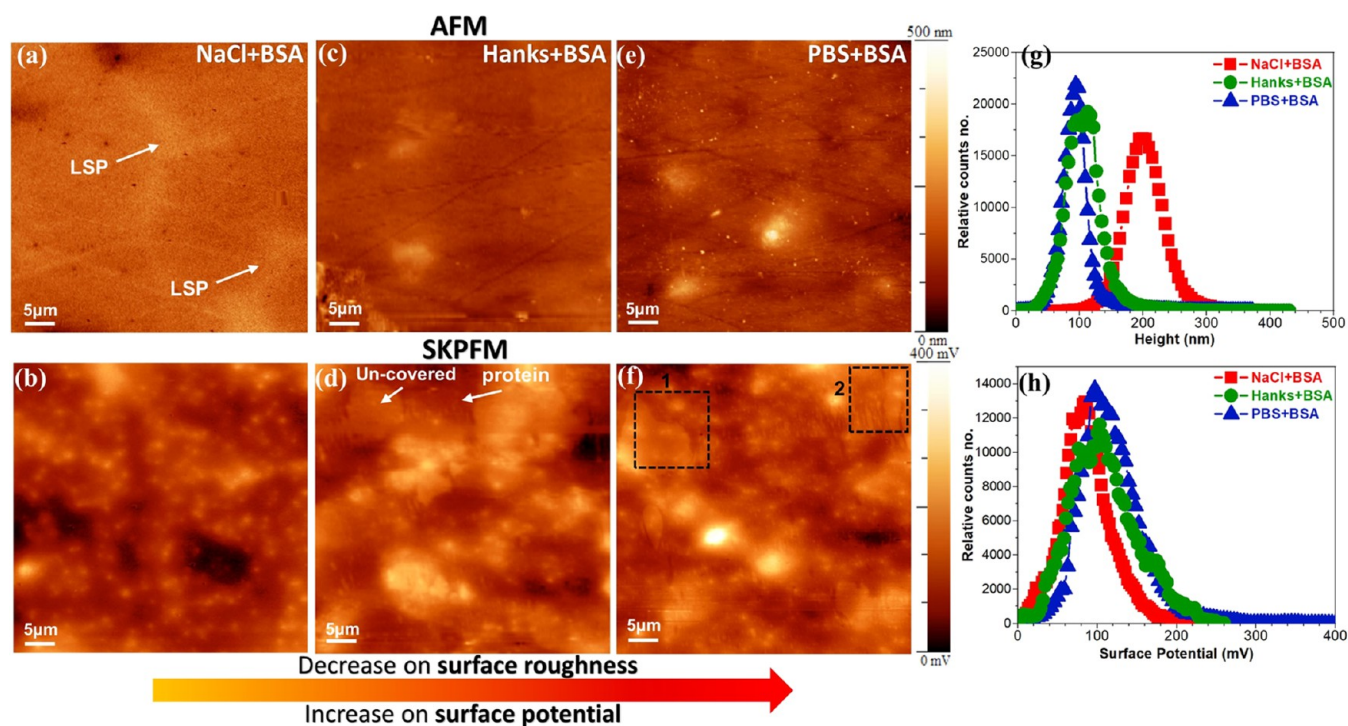


**Figure 7.** (a) Open circuit potential (OCP) and potentiodynamic polarization curves of the WE43 Mg alloy after 10 min of immersion in (b) 0.9% NaCl, (c) PBS, and (d) Hank's solutions with and without the BSA protein at 37 °C, pH 7.4, and aerated conditions.



**Figure 8.** AFM (first row) and SKPFM (second row) images of the WE43 Mg-based alloy after 10 min of immersion in (a, b) 0.9% NaCl, (c, d) Hanks', and (e, f) PBS physiological solutions at 37 °C, pH 7.4, and aerated conditions. (g) Height and (h) surface potential histograms that are related to SKPFM images in (b), (d), and (f).





**Figure 9.** AFM (first row) and SKPFM (second row) images of the WE43 Mg alloy after 10 min of immersion in (a, b) 0.9% NaCl + 4 g L<sup>-1</sup> BSA, (c, d) Hanks + 4 g L<sup>-1</sup> BSA, and (e, f) PBS + 4 g L<sup>-1</sup> BSA physiological solutions at 37 °C, pH 7.4, and aerated conditions. (g) Height and (h) surface potential histograms that are related to SKPFM images in (b), (d), and (f).

formed at the Mg oxide layer strongly affects the adsorption mechanism of the BSA protein as well as its migration and conformational arrangement, particularly concerning the electrical surface potential distribution.<sup>2</sup> Electrochemical measurements in Figure 7 and XPS results show that the high adsorption of the protein on Mg in the NaCl solution is accompanied by improvement in the corrosion resistance (lower corrosion current density). This event can be explained by the formation of a thick or multilayer of the BSA protein (a strong metal–protein complex), with lower surface potential or electronic conductivity than the substrate, that strongly controls the whole charge transfer for the electrochemical interaction at the solid/protein/electrolyte interfaces (as fully described in Section 2.6 and is discussed in the next section).<sup>10,32</sup> In PBS and Hanks' environments containing the BSA protein, the corrosion resistance of the Mg alloy slightly decreased owing to the decline in the P and Ca/P intensity signals and particularly enhancing metal–protein complex formation. The self-protecting action of phosphate and calcium phosphate species against the corrosion and biodegradation processes diminished in BSA protein media due to the imperfect, thin protective film and nonhomogeneous distribution of phosphate and calcium phosphate products.<sup>78</sup>

**3.3. Morphological and Surface Potential Evolutions in Different Simulated Fluids Containing the Albumin Protein.** The solution chemistry, pH, and type and concentration of ions/inorganic species can strongly affect the rate of metal ion release or degradation, the type of degradation products, and particularly the formation of a protective layer on the surface of Mg and its alloys.<sup>26</sup> Additionally, these parameters remarkably modify the distribution of charged and polar residues in the protein molecular structure and its isoelectric point (IEP), which

directly controls the type of protein interaction and its adsorption mechanism.<sup>10</sup> Herein, we visualize the topography and surface potential on Mg WE43 during the early stages of immersion and when an adsorbed protein nanobiofilm is present. Figure 8 presents the topography and surface potential maps after 10 min of immersion in three different solutions, including 0.9% NaCl, PBS, and Hanks' solutions at 37 °C and pH 7.4. The AFM topography image and its corresponding surface potential map in the NaCl solution (Figure 8a,b) show uniform corrosion on both  $\alpha$ -Mg and LSPs, which is slightly higher corroded in  $\alpha$ -Mg, as further proved by SEM images in Figure S1. From Figure 8a, it can be seen that the surface potential of the Mg oxide film or corrosion products for LSP is lower than that for  $\alpha$ -Mg, and its magnitude is the opposite of the fresh surface's (Figure 4). Therefore, the formation of complex corrosion products of rare-earth elements in LSP with unique electronic properties (e.g., WFE, n- or p-type semiconductor characters) significantly alters the surface potential magnitude.<sup>2</sup> However, in Hanks' and PBS solutions, AFM and SKPFM maps show heterogeneous topographies and surface potential distributions, with no obvious evidence of LSPs. The histogram analysis of topography in Figure 8g shows that the overall surface roughness distribution decreases in both Hanks' and especially PBS solutions. This reduction in the overall surface roughness is accompanied by increasing or shifting the surface potential histogram of the Mg alloy to a higher value in Hanks' and especially PBS solution (Figure 8h). This occurrence confirms the impact of inorganic species on forming the complex and protective thin film with Mg and its corrosion products, as previously established by XPS (Section 3.2) and electrochemical analysis (Figure 7).

The addition of the BSA protein to all solutions resulted in both topography and surface potential peak shifts in the histograms (Figure 9g,h) to lower values (Figure 10). This

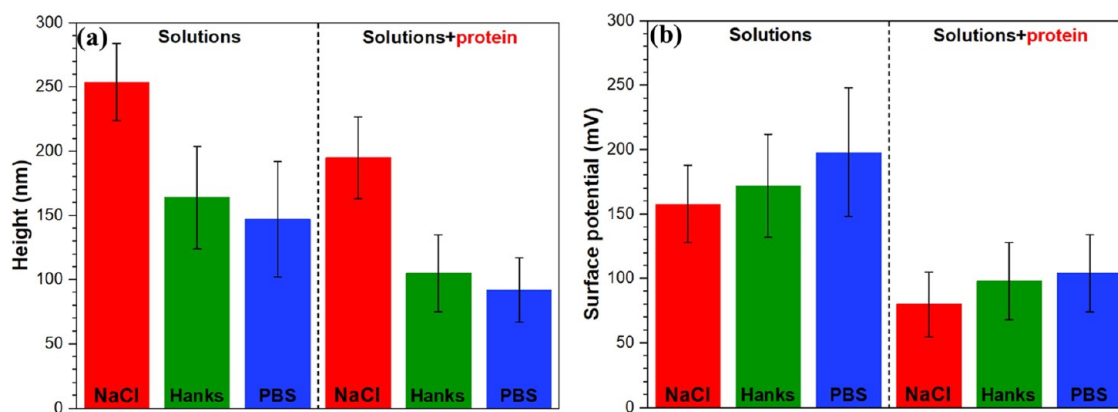


Figure 10. Extracted Gaussian distribution from the topography and surface potential histograms in Figures 8g,h and 9g,h.

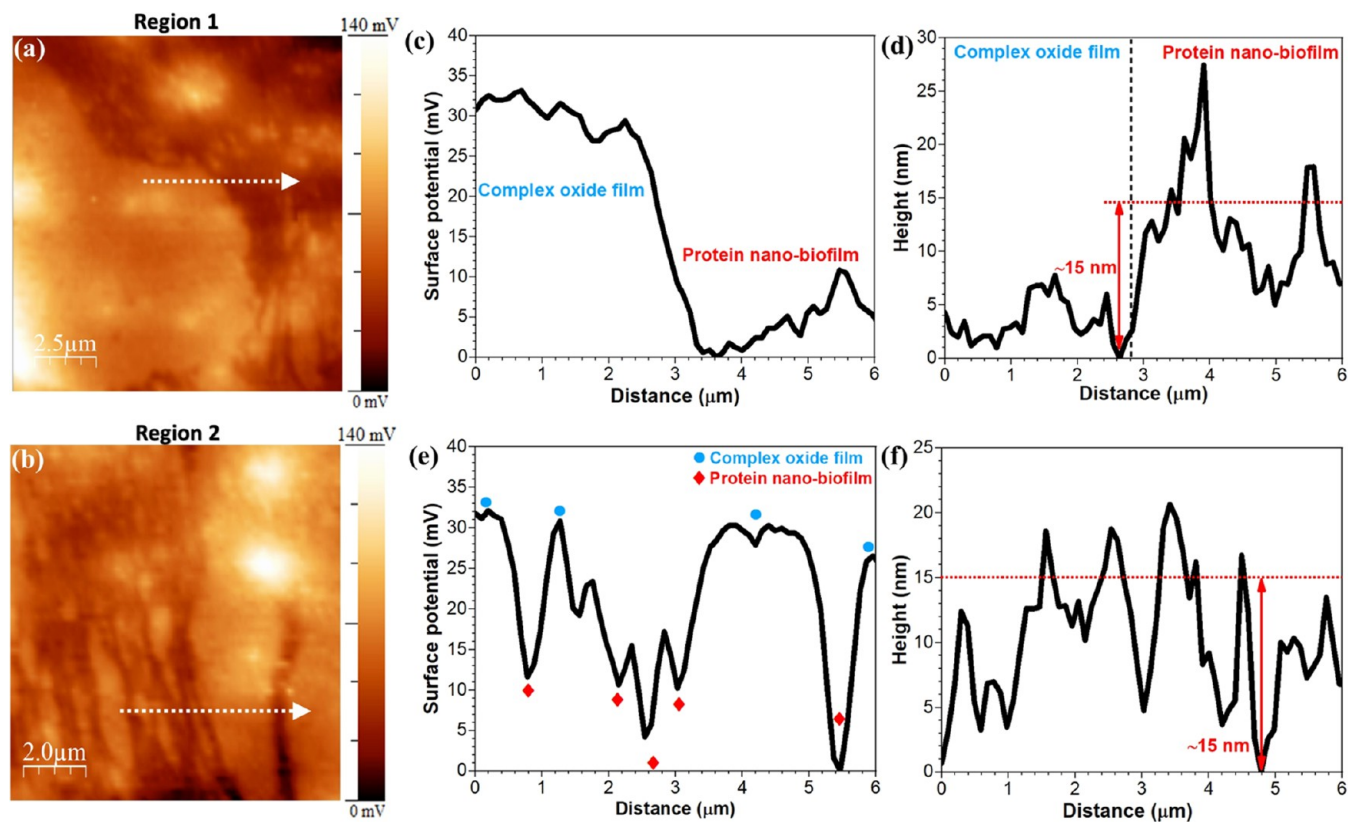


Figure 11. SKPFM high-magnified surface potential maps of (a, b) adsorbed BSA nanobiofilm alongside uncovered regions on the WE43 Mg-based alloy corresponding to regions 1 and 2 indicated in Figure 9f. (c, d) and (e, f) Surface potential and topography line profiles from (a) and (b), respectively. Topography line profiles are obtained from the same marked region in Figure 9e.

reduction in the topography value is due to the significant role of the protein-covered layer and its thin film behavior, i.e., it covers the rough Mg surface. Likewise, the lowest surface potential value of the Mg alloy in the NaCl + protein media than those in other solutions is related to the high tendency of the BSA protein for adsorption on the Mg oxide surface in the NaCl solution, which is consistent with XPS analysis (Figure 6). Therefore, the lower electrical surface potential (as a proxy for the conductivity of protein molecules) or surface charge of BSA multilayers significantly hindered the electrostatic interaction between the AFM tip apex and the Mg oxide film surface. The topography maps of Mg alloys in all simulated solutions containing the BSA protein (Figure 9a,c,e) are remarkably different from those visualized in the absence of the

protein (Figure 8). Particularly in the NaCl solution, the LSPs are not easily identifiable in the topography image, and its corresponding SKPFM map only displays a heterogeneous distribution of surface potential due to various Mg–protein complexes and corrosion (Figure 9b). The surface potential map in PBS and Hanks' solutions containing the BSA protein shows a heterogeneous surface potential distribution with new surface features with lower electrical surface potentials than the Mg oxide layer. These new surface features are nanolayers of the adsorbed protein in the form of aggregated and/or fibrillar morphologies (white arrows and black rectangles in Figure 9d,f). Typically, the electronic surface potential of soft biological matters such as DNA, protein, and peptides is controlled by the total surface charge distribution and IEP in

their molecular structure.<sup>79</sup> A protein molecule can display an overall neutral, negative, or positive charge, depending on the ionization state of protein amino acid groups.<sup>10</sup> The ionization state of a protein molecule strongly depends on environmental conditions, such as the pH of the electrolyte.<sup>80</sup> The IEP of the BSA protein has been calculated both theoretically and experimentally in the range of 4.7–5.4.<sup>2</sup> Hence, at pH 7.4, used in this study, BSA proteins are negatively charged, which strongly affects their surface potential value, electrochemical interactions, and particularly protein conformational arrangement during interaction with Mg or Mg oxides.

As mentioned above, the nanolayer of the adsorbed protein with lower surface potential (dark regions in Figure 9d,f) than the substrate (bright regions in Figure 9d,f) is only visualized in PBS and Hanks' environments. In the NaCl solution, no evidence of the adsorbed protein layer was detected. The conformational or structural pattern alterations of the adsorbed protein layer directly correlate with the physical and chemical properties of the substrate and the protein's migratory parameters.<sup>81</sup> The complex film of inorganic species formed at the Mg alloy surface, which includes  $\text{Mg}_3(\text{PO}_4)_2$ ,  $\text{Ca}_{10}(\text{PO}_4)_6(\text{OH})_2$  (only in Hanks'), and  $\text{CaCO}_3$  (only in Hanks'), strongly affects the adsorption mechanism of the BSA protein as well as its migration and conformational arrangement.<sup>82</sup> Based on our previous study<sup>2</sup> and SKPFM results, the BSA adsorption mechanism on the Mg alloy surface is under a mixed adsorption mechanism involving Langmuir isotherm and cooperative adsorption mechanisms. Langmuir's adsorption model illustrates the tendency of the protein molecules to fill the available unoccupied surface sites on solid surfaces. In the cooperative adsorption model, the protein molecules tend to adsorb in the vicinity of preadsorbed proteins that finally trigger the formation of a cluster domain and/or network-like morphology on solid surfaces,<sup>83</sup> which is consistent with the surface potential maps shown in Figure 9d,f. Indeed, these SKPFM maps demonstrate a semi-homogeneous distribution of the protein nanofilm alongside some protein-dense regions into the cluster and fibrillar morphologies with the lowest surface potential values (due to the formation of a thick protein film). This type of adsorption mechanism, which depends on the system conditions, was observed previously using in situ atomic force microscopy (in situ AFM),<sup>84</sup> supercritical angle fluorescence (SAF) microscopy,<sup>85</sup> and Monte Carlo simulation.<sup>83</sup> These studies showed the growth of two-dimensional surface clusters or protein-dense domains alongside some uncovered regions.

To better visualize the protein adsorption morphology, surface potential distribution, and thickness of the protein nanofilm on the Mg oxide film surface, higher-magnification SKPFM maps were obtained and are presented in Figure 11. These maps show that the BSA protein covered the Mg oxide surface by a dense fibrillar morphology with surface potential and height (thickness) difference values of  $\Delta V = \sim 30$  mV and  $\Delta H = \sim 15$  nm, respectively. It is worth noticing that the size of the BSA protein ( $14 \text{ nm} \times 4 \text{ nm} \times 4 \text{ nm}$ <sup>86</sup>) is consistent with the thickness of the adsorbed protein layer (nanofilm) on very smooth surfaces such as silica<sup>59,86</sup> or highly oriented pyrolytic graphite (HOPG) substrates.<sup>87</sup> However, in our study on bioactive materials, and because of the slight porosity of Mg oxide as well as its surface roughness, we were only able to assess the thickness of the adsorbed protein layer and could not resolve the size of BSA. Figure 11 proves that the chemisorption of the BSA protein on the Mg oxide film surface

induces new potential steps and band bending in Mg oxide energy diagrams that eventually reduce the total surface potential at the protein/Mg oxide film interface ( $\Delta V = \sim 30$  mV). Consequently, the multilayers of the adsorbed protein have both lower surface potential and surface charge. Thus, they present a more significant barrier to charge transfer (lower electronic conductivity) than Mg and Mg oxide surfaces. As a result, these films exact substantial control on the electrochemical interactions and rate of mass transport at protein/oxide/electrolyte interfaces.<sup>9</sup> However, since the surface potential or surface charge on the covered protein region (cluster domains or fibrillar regions) is lower than the uncovered region (Mg oxide film surface), the protein regions represent a localized active behavior with anodic reactions due to the formation of galvanic coupling. Therefore, in the covered protein regions, the deterioration reactions are dominantly localized. However, in the uncovered regions, due to better access to free electrons for reduction reactions, the cathodic reactions without lower detrimental impact will be established.<sup>88</sup> The evolutions of these chemical and physical properties at the protein/solid/electrolyte interfaces play a crucial role in metal ion release, biocompatibility, and long-term durability of Mg and its alloys in humans and simulated human fluids.

## 4. CONCLUSIONS

This research studied the electrical surface potential of a protein nanobiofilm on the surface of Mg oxide using theoretical and experimental approaches. SKPFM was used in different simulated body fluids containing the BSA protein. XPS demonstrated that the BSA protein in 0.9% NaCl has a higher tendency to cover Mg oxide than Hanks' and PBS (lowest protein adsorption) environments. In addition, the role of inorganic species in PBS and Hanks' solutions was predominant in protein adsorption, conformational arrangement, and electrical surface potential distribution on the surface of the Mg oxide film, resulting in different electrochemical responses and biodegradation mechanisms. In the early stages of immersion, SKPFM identified the adsorbed BSA protein in cluster domains and fibrillar morphology (film thickness  $\sim 15$  nm) with lower surface potential than the Mg oxide film ( $\Delta V = \sim 30$  mV). The protein cluster domains were only observed in PBS and Hanks' solutions, although no evidence of the protein nanobiofilm was detected in NaCl media. The overall surface potential of Mg oxide in all blank solutions was higher than that in the presence of the BSA protein, confirming the lower electrical surface potential or surface charge of the adsorbed protein nanofilm that reduces the electrostatic interactions between the tip apex and the substrate. Our SKPFM analysis and interpretation highlight more detailed information about protein adsorption, conformational arrangement, and electrical surface potential on Mg and Mg alloys, which directly affect their biocompatibility, biodegradability, cell adhesion, and inflammatory reactions during application in human body media.

## ■ ASSOCIATED CONTENT

### Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acs.langmuir.2c01540>.

SEM images of the immersed Mg alloy in different environments (PDF)



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### Notes

The authors declare no competing financial interest.

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## REFERENCES

- (1) Hedberg, Y. S. Role of proteins in the degradation of relatively inert alloys in the human body. *npj Mater. Degrad.* **2018**, 2, No. 26.
- (2) Rahimi, E.; Offioach, R.; Baert, K.; Terryn, H.; Lekka, M.; Fedrizzi, L. Role of phosphate, calcium species and hydrogen peroxide on albumin protein adsorption on surface oxide of Ti6Al4V alloy. *Materialia* **2021**, 15, No. 100988.
- (3) Lee, W.-H.; Loo, C.-Y.; Rohanizadeh, R. A review of chemical surface modification of bioceramics: Effects on protein adsorption and cellular response. *Colloids Surf., B* **2014**, 122, 823–834.
- (4) Anbazhagan, E.; Rajendran, A.; Natarajan, D.; Kiran, M.; Pattanayak, D. K. Divalent ion encapsulated nano titania on Ti metal as a bioactive surface with enhanced protein adsorption. *Colloids Surf., B* **2016**, 143, 213–223.
- (5) Vishwakarma, A.; Bhise, N. S.; Evangelista, M. B.; Rouwkema, J.; Dokmeci, M. R.; Ghaemmaghami, A. M.; Vrana, N. E.; Khademhosseini, A. Engineering Immunomodulatory Biomaterials To Tune the Inflammatory Response. *Trends Biotechnol.* **2016**, 34, 470–482.
- (6) Milošev, I. CoCrMo Alloy for Biomedical Applications. In *Biomedical Applications*; Springer, 2012; pp 1–72.
- (7) Karimi, S.; Nickchi, T.; Alfantazi, A. Effects of bovine serum albumin on the corrosion behaviour of AISI 316L, Co–28Cr–6Mo, and Ti–6Al–4V alloys in phosphate buffered saline solutions. *Corros. Sci.* **2011**, 53, 3262–3272.
- (8) Yu, F.; Addison, O.; Davenport, A. J. A synergistic effect of albumin and H<sub>2</sub>O<sub>2</sub> accelerates corrosion of Ti6Al4V. *Acta Biomater.* **2015**, 26, 355–365.
- (9) Yan, Y.; Yang, H.; Su, Y.; Qiao, L. Albumin adsorption on CoCrMo alloy surfaces. *Sci. Rep.* **2016**, 5, No. 18403.
- (10) Rahimi, E.; Offioach, R.; Hosseinpour, S.; Davoodi, A.; Baert, K.; Lutz, A.; Terryn, H.; Lekka, M.; Fedrizzi, L. Effect of hydrogen peroxide on bovine serum albumin adsorption on Ti6Al4V alloy: A scanning Kelvin probe force microscopy study. *Appl. Surf. Sci.* **2021**, 563, No. 150364.
- (11) Guo, S.; Pranantyo, D.; Kang, E.-T.; Loh, X. J.; Zhu, X.; Janczewski, D.; Neoh, K. G. Dominant Albumin–Surface Interactions under Independent Control of Surface Charge and Wettability. *Langmuir* **2018**, 34, 1953–1966.
- (12) Kiesel, I.; Paulus, M.; Nase, J.; Tiemeyer, S.; Sternemann, C.; Rüster, K.; Wirkert, F. J.; Mende, K.; Büning, T.; Tolan, M. Temperature-driven adsorption and desorption of proteins at solid–liquid interfaces. *Langmuir* **2014**, 30, 2077–2083.
- (13) Rezwan, K.; Meier, L. P.; Rezwan, M.; Vörös, J.; Textor, M.; Gauckler, L. J. Bovine serum albumin adsorption onto colloidal Al<sub>2</sub>O<sub>3</sub> particles: a new model based on zeta potential and UV–Vis measurements. *Langmuir* **2004**, 20, 10055–10061.
- (14) Ascencio, M.; Pekguleryuz, M.; Omanovic, S. An investigation of the corrosion mechanisms of WE43 Mg alloy in a modified simulated body fluid solution: the effect of electrolyte renewal. *Corros. Sci.* **2015**, 91, 297–310.
- (15) Mueller, W. D.; Fernandez Lorenzo de Mele, M.; Nascimento, M. L.; Zeddies, M. Degradation of magnesium and its alloys: dependence on the composition of the synthetic biological media. *J. Biomed. Mater. Res., Part A* **2009**, 90, 487–495.
- (16) Li, Z.; Song, G.-L.; Song, S. Effect of bicarbonate on biodegradation behaviour of pure magnesium in a simulated body fluid. *Electrochim. Acta* **2014**, 115, 56–65.
- (17) Zeller-Plumhoff, B.; Gile, M.; Priebe, M.; Slominska, H.; Boll, B.; Wiese, B.; Würger, T.; Willumeit-Römer, R.; Meißner, R. H. Exploring key ionic interactions for magnesium degradation in simulated body fluid—a data-driven approach. *Corros. Sci.* **2021**, 182, No. 109272.
- (18) Gonzalez, J.; Lamaka, S. V.; Mei, D.; Scharnagl, N.; Feyerabend, F.; Zheludkevich, M. L.; Willumeit-Römer, R. Mg Biodegradation Mechanism Deduced from the Local Surface Environment under Simulated Physiological Conditions. *Adv. Healthcare Mater.* **2021**, 10, No. 2100053.
- (19) Fu, J.; Su, Y.; Qin, Y.-X.; Zheng, Y.; Wang, Y.; Zhu, D. Evolution of metallic cardiovascular stent materials: a comparative study among stainless steel, magnesium and zinc. *Biomaterials* **2020**, 230, No. 119641.
- (20) Soto, F.; Wang, J.; Ahmed, R.; Demirci, U. Medical micro/nanorobots in precision medicine. *Adv. Sci.* **2020**, 7, No. 2002203.
- (21) Zeng, R.-C.; Li, X.-T.; Li, S.-Q.; Zhang, F.; Han, E.-H. In vitro degradation of pure Mg in response to glucose. *Sci. Rep.* **2015**, 5, No. 13026.
- (22) Johnson, I.; Jiang, W.; Liu, H. The effects of serum proteins on magnesium alloy degradation in vitro. *Sci. Rep.* **2017**, 7, No. 14335.
- (23) Pang, T. Y.; Kwok, J. S.; Nguyen, C. T.; Fox, K. Evaluating magnesium alloy WE43 for bioresorbable coronary stent applications. *MRS Adv.* **2021**, 6, 54–60.
- (24) Hou, P.; Han, P.; Zhao, C.; Wu, H.; Ni, J.; Zhang, S.; Liu, J.; Zhang, Y.; Xu, H.; Cheng, P.; et al. Accelerating corrosion of pure magnesium co-implanted with titanium in vivo. *Sci. Rep.* **2017**, 7, No. 41924.
- (25) Li, T.; He, Y.; Zhou, J.; Tang, S.; Yang, Y.; Wang, X. Influence of albumin on in vitro degradation behavior of biodegradable Mg–1.5 Zn–0.6 Zr–0.2 Sc alloy. *Mater. Lett.* **2018**, 217, 227–230.
- (26) Höhn, S.; Virtanen, S.; Boccaccini, A. R. Protein adsorption on magnesium and its alloys: A review. *Appl. Surf. Sci.* **2019**, 464, 212–219.

- (27) Rettig, R.; Virtanen, S. Time-dependent electrochemical characterization of the corrosion of a magnesium rare-earth alloy in simulated body fluids. *J. Biomed. Mater. Res., Part A* **2008**, *85*, 167–175.
- (28) Liu, C.; Xin, Y.; Tian, X.; Chu, P. K. Degradation susceptibility of surgical magnesium alloy in artificial biological fluid containing albumin. *J. Mater. Res.* **2007**, *22*, 1806–1814.
- (29) Wang, Y.; Lim, C. S.; Lim, C. V.; Yong, M. S.; Teo, E. K.; Moh, L. N. In vitro degradation behavior of M1A magnesium alloy in protein-containing simulated body fluid. *Mater. Sci. Eng., C* **2011**, *31*, 579–587.
- (30) Yan, W.; Lian, Y.-J.; Zhang, Z.-Y.; Zeng, M.-Q.; Zhang, Z.-Q.; Yin, Z.-Z.; Cui, L.-Y.; Zeng, R.-C. In vitro degradation of pure magnesium—the synergetic influences of glucose and albumin. *Bioact. Mater.* **2020**, *5*, 318–333.
- (31) Narth, C.; Gillet, N.; Cailliez, F.; Lévy, B.; de la Lande, A. Electron transfer, decoherence, and protein dynamics: insights from atomistic simulations. *Acc. Chem. Res.* **2015**, *48*, 1090–1097.
- (32) Rahimi, E.; Offoiach, R.; Lekka, M.; Fedrizzi, L. Electronic properties and surface potential evaluations at the protein nanobiofilm/oxide interface: Impact on corrosion and biodegradation. *Colloids Surf., B* **2022**, *212*, No. 112346.
- (33) Ron, I.; Pecht, I.; Sheves, M.; Cahen, D. Proteins as solid-state electronic conductors. *Acc. Chem. Res.* **2010**, *43*, 945–953.
- (34) Nitzan, A.; Ratner, M. A. Electron transport in molecular wire junctions. *Science* **2003**, *300*, 1384–1389.
- (35) Morita, T.; Kimura, S. Long-range electron transfer over 4 nm governed by an inelastic hopping mechanism in self-assembled monolayers of helical peptides. *J. Am. Chem. Soc.* **2003**, *125*, 8732–8733.
- (36) Stankovich, M. T.; Bard, A. J. The electrochemistry of proteins and related substances part III. Bovine serum albumin. *J. Electroanal. Chem. Interfacial Electrochem.* **1978**, *86*, 189–199.
- (37) Bogomolova, A.; Komarova, E.; Reber, K.; Gerasimov, T.; Yavuz, O.; Bhatt, S.; Aldissi, M. Challenges of Electrochemical Impedance Spectroscopy in Protein Biosensing. *Anal. Chem.* **2009**, *81*, 3944–3949.
- (38) Lacina, K.; Sopoušek, J.; Čunderlová, V.; Hlaváček, A.; Václavěk, T.; Lacinová, V. Biosensing based on electrochemical impedance spectroscopy: Influence of the often-ignored molecular charge. *Electrochem. Commun.* **2018**, *93*, 183–186.
- (39) Shevchuk, A. I.; Frolenkov, G. I.; Sánchez, D.; James, P. S.; Freedman, N.; Lab, M. J.; Jones, R.; Klenerman, D.; Korchev, Y. E. Imaging proteins in membranes of living cells by high-resolution scanning ion conductance microscopy. *Angew. Chem.* **2006**, *118*, 2270–2274.
- (40) Zhang, J.; Kuznetsov, A. M.; Medvedev, I. G.; Chi, Q.; Albrecht, T.; Jensen, P. S.; Ulstrup, J. Single-molecule electron transfer in electrochemical environments. *Chem. Rev.* **2008**, *108*, 2737–2791.
- (41) Chi, Q.; Farver, O.; Ulstrup, J. Long-range protein electron transfer observed at the single-molecule level: In situ mapping of redox-gated tunneling resonance. *Proc. Natl. Acad. Sci.* **2005**, *102*, 16203–16208.
- (42) Xu, D.; Watt, G. D.; Harb, J. N.; Davis, R. C. Electrical conductivity of ferritin proteins by conductive AFM. *Nano Lett.* **2005**, *5*, 571–577.
- (43) Zhao, J.; Davis, J. J.; Sansom, M. S.; Hung, A. Exploring the electronic and mechanical properties of protein using conducting atomic force microscopy. *J. Am. Chem. Soc.* **2004**, *126*, 5601–5609.
- (44) Sinensky, A. K.; Belcher, A. M. Label-free and high-resolution protein/DNA nanoarray analysis using Kelvin probe force microscopy. *Nat. Nanotechnol.* **2007**, *2*, 653–659.
- (45) Fuchs, F.; Caffy, F.; Demadrille, R.; Melin, T.; Grevin, B. High-resolution kelvin probe force microscopy imaging of interface dipoles and photogenerated charges in organic donor–acceptor photovoltaic blends. *ACS Nano* **2016**, *10*, 739–746.
- (46) Lee, H.; Lee, W.; Lee, J. H.; Yoon, D. S. Surface potential analysis of nanoscale biomaterials and devices using kelvin probe force microscopy. *J. Nanomater.* **2016**, *2016*, No. 4209130.
- (47) Park, J.; Lee, S.; Jang, K.; Na, S. Ultra-sensitive direct detection of silver ions via Kelvin probe force microscopy. *Biosens. Bioelectron.* **2014**, *60*, 299–304.
- (48) Chen, R. J.; Choi, H. C.; Bangsaruntip, S.; Yenilmez, E.; Tang, X.; Wang, Q.; Chang, Y.-L.; Dai, H. An investigation of the mechanisms of electronic sensing of protein adsorption on carbon nanotube devices. *J. Am. Chem. Soc.* **2004**, *126*, 1563–1568.
- (49) Park, J.; Lee, G.; Lee, G.; Lee, C. Y.; Na, S.; Lee, S. W.; Haam, S.; Huh, Y.-M.; Yoon, D. S.; Eom, K.; Kwon, T. Single-molecule recognition of biomolecular interaction via Kelvin probe force microscopy. *ACS Nano* **2011**, *5*, 6981–6990.
- (50) Lee, H.; Lee, S. W.; Lee, G.; Lee, W.; Nam, K.; Lee, J. H.; Hwang, K. S.; Yang, J.; Lee, H.; Kim, S.; et al. Identifying DNA mismatches at single-nucleotide resolution by probing individual surface potentials of DNA-capped nanoparticles. *Nanoscale* **2018**, *10*, 538–547.
- (51) Spajić, I.; Rahimi, E.; Lekka, M.; Offoiach, R.; Fedrizzi, L.; Milošev, I. Al<sub>2</sub>O<sub>3</sub> and HfO<sub>2</sub> Atomic Layers Deposited in Single and Multilayer Configurations on Titanium and on Stainless Steel for Biomedical Applications. *J. Electrochem. Soc.* **2021**, *168*, No. 071510.
- (52) Rahimi, E.; Offoiach, R.; Deng, S.; Chen, X.; Pané, S.; Fedrizzi, L.; Lekka, M. Corrosion mechanisms of magnetic microrobotic platforms in protein media. *Appl. Mater. Today* **2021**, *24*, No. 101135.
- (53) Hou, R.; Willumeit-Römer, R.; Garamus, V. M.; Frant, M.; Koll, J.; Feyerabend, F. Adsorption of Proteins on Degradable Magnesium—Which Factors are Relevant? *ACS Appl. Mater. Interfaces* **2018**, *10*, 42175–42185.
- (54) Blind, O.; Klein, L. H.; Dailey, B.; Jordan, L. Characterization of hydroxyapatite films obtained by pulsed-laser deposition on Ti and Ti-6Al-4v substrates. *Dent. Mater.* **2005**, *21*, 1017–1024.
- (55) F-06, A. *Standard Test Method for Conducting Cyclic Potentiodynamic Polarization Measurements to Determine the Corrosion Susceptibility of Small Implant Devices*; ASTM International: West Conshohocken, PA, 2001.
- (56) Hedberg, Y. S.; Žnidaršič, M.; Herting, G.; Milošev, I.; Odnevall Wallinder, I. Mechanistic insight on the combined effect of albumin and hydrogen peroxide on surface oxide composition and extent of metal release from Ti6Al4V. *J. Biomed. Mater. Res., Part B* **2019**, *107*, 858–867.
- (57) Milošev, I. The effect of biomolecules on the behaviour of CoCrMo alloy in various simulated physiological solutions. *Electrochim. Acta* **2012**, *78*, 259–273.
- (58) Wood, M. H.; Payagalage, C. G.; Geue, T. Bovine Serum Albumin and Fibrinogen Adsorption at the 316L Stainless Steel/Aqueous Interface. *J. Phys. Chem. B* **2018**, *122*, 5057–5065.
- (59) Ron, I.; Sepunaru, L.; Itzhakov, S.; Belenkova, T.; Friedman, N.; Pecht, I.; Sheves, M.; Cahen, D. Proteins as electronic materials: Electron transport through solid-state protein monolayer junctions. *J. Am. Chem. Soc.* **2010**, *132*, 4131–4140.
- (60) Melitz, W.; Shen, J.; Kummel, A. C.; Lee, S. Kelvin probe force microscopy and its application. *Surf. Sci. Rep.* **2011**, *66*, 1–27.
- (61) Glatzel, T.; Sadewasser, S.; Shikler, R.; Rosenwaks, Y.; Lux-Steiner, M. C. Kelvin probe force microscopy on III–V semiconductors: the effect of surface defects on the local work function. *Mater. Sci. Eng., B* **2003**, *102*, 138–142.
- (62) Cherniavskaya, O.; Chen, L.; Weng, V.; Yuditsky, L.; Brus, L. E. Quantitative Noncontact Electrostatic Force Imaging of Nanocrystal Polarizability. *J. Phys. Chem. B* **2003**, *107*, 1525–1531.
- (63) Greiner, M. T.; Lu, Z.-H. Thin-film metal oxides in organic semiconductor devices: their electronic structures, work functions and interfaces. *NPG Asia Mater.* **2013**, *5*, No. e55.
- (64) Zhu, H.; Ramprasad, R. Effective work function of metals interfaced with dielectrics: a first-principles study of the Pt-HfO<sub>2</sub> interface. *Phys. Rev. B* **2011**, *83*, No. 081416.

- (65) Olsson, F.; Persson, M. A density functional study of adsorption of sodium-chloride overlayers on a stepped and a flat copper surface. *Surf. Sci.* **2003**, *540*, 172–184.
- (66) Sadeghi, A.; Barattoff, A.; Ghasemi, S. A.; Goedecker, S.; Glatzel, T.; Kawai, S.; Meyer, E. Multiscale approach for simulations of Kelvin probe force microscopy with atomic resolution. *Phys. Rev. B* **2012**, *86*, No. 075407.
- (67) Liscio, A.; Palermo, V.; Samorì, P. Nanoscale quantitative measurement of the potential of charged nanostructures by electrostatic and Kelvin probe force microscopy: unraveling electronic processes in complex materials. *Acc. Chem. Res.* **2010**, *43*, 541–550.
- (68) Palermo, V.; Liscio, A.; Palma, M.; Surin, M.; Lazzaroni, R.; Samorì, P. Exploring nanoscale electrical and electronic properties of organic and polymeric functional materials by atomic force microscopy based approaches. *Chem. Commun.* **2007**, 3326–3337.
- (69) Liu, J.; Song, Y.; Chen, J.; Chen, P.; Shan, D.; Han, E.-H. The special role of anodic second phases in the micro-galvanic corrosion of EW75 Mg alloy. *Electrochim. Acta* **2016**, *189*, 190–195.
- (70) Michaelson, H. B. The work function of the elements and its periodicity. *J. Appl. Phys.* **1977**, *48*, 4729–4733.
- (71) Rahimi, E.; Rafsanjani-Abbasi, A.; Imani, A.; Davoodi, A. TiO<sub>2</sub>/Cu<sub>2</sub>O coupled oxide films in Cl<sup>−</sup> ion containing solution: Volta potential and electronic properties characterization by scanning probe microscopy. *Mater. Chem. Phys.* **2018**, *212*, 403–407.
- (72) Rohwerder, M.; Turcu, F. High-resolution Kelvin probe microscopy in corrosion science: scanning Kelvin probe force microscopy (SKPFM) versus classical scanning Kelvin probe (SKP). *Electrochim. Acta* **2007**, *53*, 290–299.
- (73) Dong, H.; Lin, F.; Boccacini, A. R.; Virtanen, S. Corrosion behavior of biodegradable metals in two different simulated physiological solutions: Comparison of Mg, Zn and Fe. *Corros. Sci.* **2021**, *182*, No. 109278.
- (74) Tie, D.; Feyerabend, F.; Hort, N.; Willumeit, R.; Hoeche, D. XPS studies of magnesium surfaces after exposure to Dulbecco's modified eagle medium, Hank's buffered salt solution, and simulated body fluid. *Adv. Eng. Mater.* **2010**, *12*, B699–B704.
- (75) Mena-Morcillo, E.; Veleza, L. Degradation of AZ31 and AZ91 magnesium alloys in different physiological media: Effect of surface layer stability on electrochemical behaviour. *J. Magnesium Alloys* **2020**, *8*, 667–675.
- (76) Ishizaki, T.; Kudo, R.; Omi, T.; Teshima, K.; Sonoda, T.; Shigematsu, I.; Sakamoto, M. Corrosion resistance of multilayered magnesium phosphate/magnesium hydroxide film formed on magnesium alloy using steam-curing assisted chemical conversion method. *Electrochim. Acta* **2012**, *62*, 19–29.
- (77) Takemoto, S.; Hattori, M.; Yoshinari, M.; Kawada, E.; Oda, Y. Corrosion behavior and surface characterization of titanium in solution containing fluoride and albumin. *Biomaterials* **2005**, *26*, 829–837.
- (78) Mei, D.; Lamaka, S. V.; Gonzalez, J.; Feyerabend, F.; Willumeit-Römer, R.; Zheludkevich, M. L. The role of individual components of simulated body fluid on the corrosion behavior of commercially pure Mg. *Corros. Sci.* **2019**, *147*, 81–93.
- (79) Leung, C.; Kinns, H.; Hoogenboom, B. W.; Howorka, S.; Mesquida, P. Imaging surface charges of individual biomolecules. *Nano Lett.* **2009**, *9*, 2769–2773.
- (80) Kopac, T.; Bozgeyik, K.; Yener, J. Effect of pH and temperature on the adsorption of bovine serum albumin onto titanium dioxide. *Colloids Surf., A* **2008**, *322*, 19–28.
- (81) Bathawab, F.; Bennett, M.; Cantini, M.; Reboud, J.; Dalby, M. J.; Salmerón-Sánchez, M. Lateral chain length in polyalkyl acrylates determines the mobility of fibronectin at the cell/material interface. *Langmuir* **2016**, *32*, 800–809.
- (82) Combes, C.; Rey, C. Adsorption of proteins and calcium phosphate materials bioactivity. *Biomaterials* **2002**, *23*, 2817–2823.
- (83) Rabe, M.; Verdes, D.; Seeger, S. Understanding protein adsorption phenomena at solid surfaces. *Adv. Colloid Interface Sci.* **2011**, *162*, 87–106.
- (84) Mulheran, P. A.; Pellenc, D.; Bennett, R.; Green, R.; Sperrin, M. Mechanisms and dynamics of protein clustering on a solid surface. *Phys. Rev. Lett.* **2008**, *100*, No. 068102.
- (85) Rabe, M.; Verdes, D.; Seeger, S. Understanding cooperative protein adsorption events at the microscopic scale: a comparison between experimental data and Monte Carlo simulations. *J. Phys. Chem. B* **2010**, *114*, 5862–5869.
- (86) Jachimska, B.; Tokarczyk, K.; Łapczyńska, M.; Puciul-Malinowska, A.; Zapotoczny, S. Structure of bovine serum albumin adsorbed on silica investigated by quartz crystal microbalance. *Colloids Surf., A* **2016**, *489*, 163–172.
- (87) Barinov, N. A.; Prokhorov, V. V.; Dubrovina, E. V.; Klinov, D. V. AFM visualization at a single-molecule level of denaturated states of proteins on graphite. *Colloids Surf., B* **2016**, *146*, 777–784.
- (88) Rahimi, E.; Offoia, R.; Baert, K.; Terry, H.; Fedrizzi, L.; Lekka, M. Albumin Protein Adsorption on CoCrMo Implant Alloy: Impact on the Corrosion Behaviour at Localized Scale. *J. Electrochem. Soc.* **2022**, *169*, No. 031507.

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