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Effect of protein adsorption on the corrosion behaviour of 70Cu-30Ni alloy in artificial seawater

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Abstract
Copper alloys often used in cooling circuits of industrial plants can be affected by biocorrosion induced by biofilm formation. The objective of this work was to study the influence of protein adsorption, which is the first step in biofilm formation, on the electrochemical behaviour of 70Cu-30Ni (wt. %) alloy in static artificial seawater and on the chemical composition of oxide layers. For that purpose, electrochemical measurements performed after 1 h of immersion were combined to surface analyses. A model is proposed to analyse impedance data. In the presence of bovine serum albumin (BSA, model protein), the anodic charge transfer resistance deduced from EIS data at $E_{corr}$ is slightly higher, corresponding to lower corrosion current. Without BSA, two oxidized layers are shown by XPS and ToF-SIMS: an outer layer mainly composed of copper oxide (Cu$_2$O redeposited layer) and an inner layer mainly composed of oxidized nickel, with a global thickness of ~ 30 nm. The presence of BSA leads to a mixed oxide layer (CuO, Cu$_2$O, Ni(OH)$_2$) with a lower thickness (~ 10 nm). Thus, the protein induces a decrease of the dissolution rate at $E_{corr}$ and hence a decrease of the amount of redeposited Cu$_2$O and of the oxide layer thickness.

**Keywords:** copper alloy; seawater; protein adsorption; EIS; surface analysis.
1. Introduction

Power plants require cooling circuits with water as the cooling agent; therefore, they are generally located on seacoasts owing to the ready availability of abundant seawater. Tubes are the basic components of heat exchangers; they provide the heat transfer surface between one fluid flowing inside the tubes and other fluid flowing outside the tubes [1]. In cooling circuits, the water is usually circulating but, frequently, plant outages cause provisional stagnant condition that can persist for hours or, at worst, for days. This condition is of particular concern for corrosion risk, especially at the beginning of plant operation [2]. Therefore, hydrodynamics is one of the parameters to be studied.

Copper and copper alloys are commonly used in condensers and heat exchangers due to their high thermal conductivity, good resistance to corrosion and good mechanical workability. Cu-Ni alloys are preferred in marine environments because of their corrosion resistance due to the formation of a thin, adherent, protective surface film which forms naturally and quickly upon exposure to clean seawater. That surface film is predominantly made up of cuprous oxide, often containing nickel and iron oxides, cuprous hydroxychloride and cupric oxide; it becomes more complex when aggressive condition from the water causes corrosion [2].

Cooling circuits of industrial plants are ideal incubators for microorganisms because they offer plenty of water, are maintained at temperatures between 30 and 60°C, at pH of 6 to 9, and provide a continuous source of nutrients, such as inorganic or organic compounds. The microorganisms present in cooling water circuits can be divided into planktonic or sessile cells. Sessile ones attach to surfaces and form what is known as biofilm. The development of a biofilm is considered to be a multistage process involving the following major steps: a) formation of an organic conditioned film on the solid surface by adsorption of biomolecules such as proteins [3], b) transport of microorganisms from the water to the surface, c) adhesion of microorganisms onto the surface, d) replication of the attached cells and production of exopolymers, e) detachment of parts of the biofilm that are swept along by the flowing water to repeat the process of biofilm formation elsewhere [4]. Biofouling is hence a consequence of biofilm formation. The significant negative effects of biofouling are the blockage of water free flow in the cooling circuit and consequent mechanical damage to pumps, clogging of condenser tubes, reduction of the heat transfer efficiency and microbially induced corrosion (MIC) also called biocorrosion [5,6]. Cu-Ni alloys have also shown good resistance to biofouling; the reason for the antifouling behaviour of these alloys is still not fully understood but the protective surface film should play a role [7].

There are two opposite goals for the control of microbial adhesion and biofilm formation: one is the prevention of biofilms, and the other one is their promotion. Controlling the adsorption of biomolecules, which is the first step in biofilm formation, by modifying the surface properties of the material may represent a good strategy for inhibiting microbial growth [8]. The global objective of this work was to study the influence of biomolecules adsorption on the electrochemical behaviour of 70Cu-30Ni (wt. %) alloy and the chemical composition of oxide layers in seawater environment, in different hydrodynamics conditions (static conditions, and using a rotating electrode). For that purpose, artificial seawater (ASW) was chosen, either without any biomolecule or added with bovine serum albumin (BSA), a model protein often used to study the protein-surface interactions due to its low cost and to a good knowledge of its properties [9-11]. Electrochemical measurements (corrosion potential ($E_{corr}$) vs time, polarization curves and electrochemical impedance spectroscopy (EIS)) performed, in this work, in static conditions during the very first steps of oxide layers formation (1 h
immersion time) were combined to surface analysis by X-ray photoelectron spectroscopy (XPS) and time-of-flight secondary ions mass spectrometry (ToF-SIMS). Electrochemical measurements in well-controlled hydrodynamics conditions, using a rotating electrode, have been performed, and will be the purpose of a forthcoming paper. Although 1 h immersion time is a short time for the formation of oxides, it is not a short time for the adsorption of BSA since an adsorption plateau (steady-state) is reached within 10-20 min [11-13].

2. Materials and methods

2.1 Samples and solution

The material under study was 70Cu-30Ni alloy (Cu 68.5, Ni 30, Mn 0.8, Fe 0.7 wt. %). The samples provided by RSE S.p.A. were disks cut from real new condenser tubes, and then flattened at Chimie ParisTech. The geometrical surface area exposed to the solution was either 0.45 cm$^2$, and the tightness between the disk and the sample holder was ensured by a Viton® O-ring. Before electrochemical measurements, samples were mechanically polished with SiC papers down to grade 1200, then degreased in an ultrasonic bath three times in acetone for 10 min, once in ethanol for 10 min, and once in ultra-pure water for 10 min, dried under an argon flow, and finally exposed to UV for 20 min. Before surface analysis, samples were mechanically polished first with SiC papers down to grade 1200, then with 6 μm, 3 μm, and 1 μm diamond paste. Their subsequent treatment was the same as before electrochemical measurements.

The solution used was aerated artificial seawater (ASW; composition (g/L): NaCl (24.615), KCl (0.783), Na$_2$SO$_4$ (4.105), MgCl$_2$(H$_2$O)$_6$ (11.060), CaCl$_2$ (1.160), NaHCO$_3$ (0.201); pH = 8.0; ionic strength = 0.7155 M), without and with 20 mg.L$^{-1}$ of BSA (about 99% purity (Fraction V); Sigma-Aldrich).

All the experiments were performed at room temperature.

2.2 Electrochemical measurements

The electrochemical measurements were performed with a three-electrode cell, designed and manufactured at Chimie ParisTech, with a volume of solution of about 0.1 L. The working electrode was the 70Cu-30Ni alloy (disk samples), the counter-electrode was a platinum wire, and the reference electrode was a saturated calomel electrode (SCE; 0.245 V vs SHE). Experiments were carried out in stagnant conditions (static working electrode and solution).

The corrosion potential ($E_{corr}$) was measured during the first hour of immersion, then cathodic or anodic polarization curve was plotted separately starting from +20 or -20 mV vs $E_{corr}$, respectively, using a scan rate of 0.5 mV.s$^{-1}$. Electrochemical impedance diagrams were plotted at $E_{corr}$ after 1 h of immersion, with a frequency domain ranging from $10^2$ Hz to $10^3$ Hz, 7 points per decade, and an amplitude of 10 mV peak-to-peak. Electrochemical measurements were performed with an EC-Lab SP-200 system from Bio-Logic.

2.3 Surface analysis

Three 70Cu-30Ni samples were analyzed by XPS and ToF-SIMS: 1) after polishing, 2) after 1 h of immersion at $E_{corr}$ in ASW without BSA, and 3) after 1 h of immersion at $E_{corr}$ in ASW with BSA. After the electrochemical measurements, the samples were gently dipped in ultra-
pure water three times to remove the BSA molecules loosely bound to the surface, then dried with argon before introduction in the fast-entry lock chamber of the XPS spectrometer.

XPS analyses were performed with a Thermo Electron Escalab 250 spectrometer, using a monochromatised Al Kα X-ray source (1486.6 eV). The analyser pass energy was 100 eV for survey spectra and 20 eV for high resolution spectra. The spectrometer was calibrated using Au 4f/2 at 84.1 eV. The following core levels were recorded: Cu 2p (and Auger lines), Ni 2p, O 1s, C 1s, and N 1s. All spectra were referred to the C 1s peak for the carbon involved in C-C and C-H bonds, located at 285 eV. The fitting of the complex C 1s signal was based on published data [11]. Curve fitting of the spectra was performed with the Thermo Electron software “Avantage”. The inelastic mean free path values were calculated by the TPP2M formula [14], and the photoemission cross-sections were taken from Scofield [15].

ToF-SIMS analyses were done using a ToF-SIMS V spectrometer (ION-TOF GmbH). The analysis chamber was maintained at less than 10⁻⁹ Pa in operation conditions. The depth profiles were performed using the instrument in dual beam mode. A pulsed 25 keV Bi⁺ primary ion source (LMIG) at a current of 1.2 pA (high mass resolution mode), rastered over a scan area of 100×100 μm², was used as the analysis beam. The sputtering was performed using a 1 keV Cs⁺ ion beam at a current of 50 nA, and rastered over an area of 300×300 μm². The depth profiles were obtained in negative polarity meaning only negative ions were analysed. The distribution of the ionized fragments, all measured quasi-simultaneously, were plotted versus Cs⁺ ion sputtering time. The intensity was reported using a logarithmic scale, which gave equal emphasis to signals of all intensities. The variation of the ion intensity with sputtering time reflects the variation of the in-depth concentration but is also dependent on the matrix from which the ions are emitted. Data acquisition and processing were performed using the IonSpec software.

3. Results

3.1 Electrochemical measurements

During the first hour of immersion, the corrosion potential \( E_{corr} \) decreases till reaching a steady-state value of -0.230 ± 0.011 V vs SCE without BSA. The presence of 20 mg.L⁻¹ of BSA induces a more anodic \( E_{corr} \) value (-0.203 ± 0.008 V vs SCE i.e. difference of ~ 30 mV with the value found without protein).

Figure 1(a) shows the cathodic polarization curves obtained without and with 20 mg.L⁻¹ of BSA. Two current plateaus illustrating mass transport limited reactions can be observed. The short plateau at around -0.30 V vs SCE (\( j_\text{lim} \sim 10 \mu A.cm^{-2} \)) illustrates the first step of dissolved oxygen reduction with production of \( H_2O_2 \), that can be written at \( pH = 8.0 \) as follows [16,17]:

\[
O_2 + 2H_2O + 2e^- \rightarrow H_2O_2 + 2OH^- \quad (1)
\]

The wide plateau observed for potentials ranging from -0.40 to -1.00 V vs SCE corresponds to the second step of dissolved oxygen reduction with transfer of 4 electrons, that can be written at \( pH = 8.0 \) as follows:

\[
O_2 + 2H_2O + 4e^- \rightarrow 4OH^- \quad (2)
\]
The ratio between the two plateau current densities is comprised between 2 and 3 (theoretical value: 2). This observation is consistent with the presence on the alloy surface of a layer composed of reducible products (oxides) in the oxygen reduction domain. The current increase (in absolute value) observed below -1.00 V vs SCE illustrates water reduction (hydrogen evolution reaction).

The second plateau current density is divided by 2 in the presence of the protein ($|j| \sim 40 \mu A.cm^{-2}$ without BSA to be compared to $|j| \sim 20 \mu A.cm^{-2}$ with BSA). This difference in plateau current is not necessarily due to the BSA but may also be induced by a difference in natural convection from one experiment to another.

The anodic polarization curves obtained without and with 20 mg.L$^{-1}$ of BSA are presented in Figure 1(b). In the absence of protein, the anodic polarization curves show very high dissolution currents. In particular, a pseudo-plateau, corresponding to a current density of $\sim 2.5 mA.cm^{-2}$, can be observed at around 0.4 V vs SCE; therefore, this plateau current is not a passive current, and hence the oxide film formed on the surface is not a passive layer. Moreover, visual observation of the electrode surface after an anodic scan shows a green layer, the thickness of which increases with the end anodic potential. This green layer corresponding to high anodic currents may be formed by redeposition due to the saturation of the solution in copper ions. In the presence of BSA, lower current densities can be observed near $E_{\text{corr}}$, for potentials lower than 0.0 V vs SCE; this difference in current densities is partly due to the difference in $E_{\text{corr}}$ values without and with BSA. Above 0.0 V vs SCE, the two anodic polarization curves overlap. Thus, only a slight influence of the protein on the anodic electrochemical behaviour can be seen. Moreover, after the anodic polarization curve, the green color is not observed in the solution with BSA.

Figure 2 shows the impedance diagrams in the complex plane (opposite of the imaginary part of the impedance $-Z_j$ vs real part of the impedance $Z_r$; the electrochemical impedance $Z$ is a complex number that can be written as: $Z(f) = Z_r(f) + jZ_j(f)$, with $f$ the ac-frequency (in Hz)) plotted at $E_{\text{corr}}$ after 1 h of immersion in aerated artificial seawater, without protein and with 20 mg.L$^{-1}$ of BSA. The diagrams exhibit two capacitive loops: one high frequency (HF) depressed semi-circle, and a low frequency (LF) loop. The size of the HF loop is slightly increased in the presence of BSA. After impedance measurement at $E_{\text{corr}}$ without or with BSA, there is no sign of pitting corrosion on the electrode surface.

Surface analysis

For the sample after polishing, it is observed from XPS and ToF-SIMS data (not shown here) a mixed copper oxides (Cu$_2$O and CuO) and nickel hydroxide (Ni(OH)$_2$) layer, with a thickness of $\sim$1.3 nm and the following atomic composition: 43 at. % Cu$_2$O + 25 at. % CuO + 32 at. % Ni(OH)$_2$. An enrichment in copper of the alloy beneath the oxide layer is also detected (77 at. % Cu + 23 at. % Ni to be compared to 68 at. % Cu + 32 at. % Ni for the bulk alloy).
Differences in chemical composition and thickness of the oxide layers are observed without and with BSA.

After immersion in the BSA-free ASW, the XPS Cu 2p_{3/2} core level peak with a binding energy located at 932.3 eV (Fig. 3(a)) and the Cu Auger line (L_3M_{45}M_{45}) at a binding energy of 570.2 eV (Fig. 3(b)) demonstrate the presence of Cu\(^+\) instead of Cu\(^{2+}\) as for the sample after polishing [18,19].

From the XPS Ni 2p_{3/2} core level spectrum with a peak located at 856.1 eV and the corresponding satellite at 861.7 eV, the presence of Ni(OH)\(_2\) in the surface layer is shown (Fig. 4) [20]. The atomic composition of this oxide layer calculated from XPS data (93 at. % Cu\(_2\)O + 7 at. % Ni(OH)\(_2\)) indicates an enrichment in Cu\(_2\)O. It is important to mention that the depth analyzed by XPS is about 10 nm.

A characteristic ToF-SIMS negative depth profile obtained with 70Cu-30Ni alloy immersed during 1 h at \(E_{corr}\) in ASW without BSA is presented in Figure 5. This profile allows evidencing a possible stratification of the different compounds on the alloy surface. It shows four regions. A first one that extends from 0 s to 10 s of sputtering characterized by an increase of all signals and corresponding to the time necessary to reach a steady state. After 10 s, one enters the second region that extends up to 200 s of sputtering. This region is characterized by an intense CuO\(_2^-\) signal. As soon as one probes deeper into this region, a progressive increase of the NiO\(_2^-\) signal is observed. This indicates that the outer surface layer is mainly composed of copper oxide, with the presence of some Ni oxide and/or hydroxide, in agreement with XPS data. In the third region that extends from 200 s to 290 s, a sharp decrease of the CuO\(_2^-\) signal and a very intense NiO\(_2^-\) signal are observed. This region is assigned to the formation of a nickel oxide and/or hydroxide inner layer in which the presence of oxidized copper cannot be excluded. Finally, after 290 s of sputtering, one enters the fourth region characterized by a sharp decrease of all oxidized signals (18O\(^-\), CuO\(_2^-\) and NiO\(_2^-\)) and a constant and intense plateau for the Ni\(_2^+\) signal that is characteristic of the metallic substrate. From the ToF-SIMS depth profiles, it can be concluded that in the absence of BSA two oxidized layers can be observed: an 18nm-thick outer layer mainly composed of copper oxide and a 12nm-thick inner layer mainly composed of oxidized nickel, with a global thickness of \(\sim\)30 nm.

After immersion in the BSA-containing solution, the XPS Cu 2p_{3/2} core level spectrum exhibits three peaks (Fig. 3(a)): one located at 932.3 eV attributed to Cu\(^0\) and/or Cu\(^+\), another one with a binding energy of 934.5 eV and the corresponding satellite at higher binding energy attributed to Cu\(^{2+}\). Moreover, the Cu Auger line at a binding energy of 570.0 eV (Fig. 3(b)) shows the presence of Cu\(^+\). The XPS Ni 2p_{3/2} core level spectrum recorded in the presence of BSA exhibits the same features as without protein: a peak at a binding energy of 856.0 eV and the corresponding satellite at 861.6 eV showing the presence of Ni(OH)\(_2\) in the surface layer (Fig. 4). From these XPS data, it can be concluded that the presence of BSA leads to a mixed copper oxides (Cu\(_2\)O and CuO) and nickel hydroxide layer, with the following atomic composition: 11 at. % Cu\(_2\)O + 44 at. % CuO + 45 at. % Ni(OH)\(_2\). A lower
amount of Cu$_2$O and higher amounts of CuO and Ni(OH)$_2$ are detected compared to the results obtained in the absence of BSA.

By ToF-SIMS, it is not possible to identify a stratification of the different compounds on the alloy surface in the presence of the protein. The depth profiles show one mixed oxide layer (oxidized copper and nickel) with a lower thickness of ~10 nm.

The N 1s spectrum recorded after immersion in ASW with BSA (Fig. 6(a)) exhibits a major symmetric peak, centered at 400.2 eV, as expected for the amine or amide groups of BSA [21]. The C 1s signal obtained in the same conditions is shown in Figure 6(b). It can be fitted with three contributions corresponding to well identified carbon bonds present in the BSA molecule: C1, at a binding energy of 285.0 eV, assigned to C-C and C-H; C2, at a binding energy of 286.4 eV, attributed to C-N and C-O single bonds; and C3, at a binding energy of 288.3 eV, assigned to O=C-O and O=C-N (peptide bonds) bonds [11].

From the N 1s and C 1s signals, it is possible to calculate different “nitrogen/carbon” or “carbon/carbon” atomic ratios. The values of these ratios obtained for the 70Cu-30Ni alloy immersed in ASW without and with BSA, as well as those estimated in previous studies for the BSA powder [12,22] are presented in Table 1. The good agreement between the values for the sample put into contact with the BSA and the BSA powder provides a fingerprint for the protein, and allows us to conclude that the protein is present on the surface. The thickness estimated from XPS data for this adsorbed layer is ~3 nm which, according to the size of the BSA molecule [10], corresponds to one monolayer.

4. Discussion

4.1 Impedance model (EIS)

It is accepted by many authors that the corrosion of copper is controlled by mass transport processes to and from corroding surfaces, involving O$_2$, Cl$^{-}$, OH$^{-}$, Cu$^{+}$ and CuCl$_2^{-}$ species [23]. In aerated solution, the cathodic reaction is the reduction of dissolved oxygen, and the anodic reactions involve:

$$\text{Cu} + \text{Cl}^{-} \rightleftharpoons \text{CuCl} + e^{-} \quad (3)$$

followed by the formation of the soluble cuprous complex:

$$\text{CuCl} + \text{Cl}^{-} \rightleftharpoons \text{CuCl}_2^{-} \quad (4)$$

Then, cuprous oxide can be formed from CuCl$_2^{-}$ as follows [16]:

$$2\text{CuCl}_2^{-} + 2\text{OH}^{-} \rightarrow \text{Cu}_2\text{O} + \text{H}_2\text{O} + 4\text{Cl}^{-} \quad (5)$$

At $E_{\text{corr}}$, the anodic and cathodic currents have the same magnitude and the net current is equal to zero. By principles of summation of currents, the anodic and cathodic impedances must be in parallel. Both the anodic and the cathodic reactions are mass transport limited. The anodic impedance can be depicted by a charge transfer resistance ($R_t^a$) in series with an impedance that illustrates mass transport and partial blocking effect by CuCl ($Z_{\theta,D}^a$); whereas,
the cathodic impedance can be depicted by a charge transfer resistance \( R_t^c \) in series with an impedance that illustrates \( \text{O}_2 \) mass transport \((Z_D^c)\). A double layer capacitance \( C_{dl} \) is added in parallel with the anodic and the cathodic impedances. As the impedance response for electrochemical systems often reflects a distribution of reactivity that is commonly represented in equivalent electrical circuits as a constant-phase element (CPE), \( C_{dl} \) is replaced here by \( CPE_{dl} \) which is a constant phase element related to the double layer. The CPE impedance is expressed in terms of model parameters \( \alpha \) and \( Q \) as:

\[
Z_{CPE}(\omega) = \frac{1}{Q(j\omega)^\alpha} \tag{6}
\]

where \( \omega \) is the angular frequency (in \( s^{-1} \)) given by:

\[
\omega = 2\pi f \tag{7}
\]

When \( \alpha = 1 \), the parameter \( Q \) has units of capacitance; otherwise, \( Q \) has units of \( \Omega^{-1}\text{.cm}^{-2}\cdot\text{s}^{(\alpha-1)} \).

Thus, in a first approach, the 70Cu-30Ni/ASW system can be modeled by the general equivalent electrical circuit illustrated in Figure 7(a), where \( R_e \) is the electrolyte resistance.

As the first current plateau for the reduction of dissolved oxygen is observed close to \( E_{corr} \), one hypothesis is that this plateau can be extrapolated down to \( E_{corr} \); thus, \( R_t^c \) can be neglected and the cathodic mass transport impedance is a Warburg impedance \((W_c)\) given by:

\[
Z_{W_c} = \frac{1}{k_c \sqrt{j\omega}} \tag{8}
\]

with \( k_c \) expressed in \( s^{0.5}\cdot\Omega^{-1}\cdot\text{cm}^{-2} \). Taking into account experimental cathodic voltammograms, the 70Cu-30Ni/ASW system can be modeled by the simplified circuit presented in Figure 7(b).

For the same experimental impedance data as those presented in Figure 2 in the complex plane, the absolute value of the imaginary part of the impedance \(|Z_j|\) was plotted as a function of the frequency in logarithmic coordinates (Fig. 8(a)) [24].

In the HF range, a pseudo-straight line with a slope lower than 1 in absolute value but varying slightly with the frequency can be observed. This slope value lower than 1 suggests a CPE-like behavior (it should be equal to 1 in case of pure capacitive behavior). The CPE parameters \( \alpha \) and \( Q \) can be graphically obtained in the case of a \( R//\text{CPE} \) circuit, following the method presented by Orazem et al. [24]. The parameter \( \alpha \) is calculated from the slope of the \( \log|Z_j| \) vs \( \log f \) curve in the HF range:

\[
\alpha = \left. \frac{d \log|Z_j(f)|}{d \log f} \right|_{\text{at HF}} \tag{9}
\]
and $Q$ is obtained from $\alpha$ as follows:

$$Q = -\frac{1}{Z_f(f)(2\pi f)^{\alpha}} \times \sin\left(\frac{\alpha \pi}{2}\right)$$

(10)

However, as the slope varies with the frequency, $\alpha$ cannot be determined from Figure 8(a). To better visualize a possible constant value of the slope in a narrow frequency range, the $d \log Z_f / d \log f$ vs $\log f$ curves were calculated from those presented in Figure 8(a) (derivative curves; Fig. 8(b)). In the case of a R//CPE circuit, a plateau would be observed at HF corresponding to a value of $-\alpha$. In Figure 8(b), no plateau is visible, in particular, at very HF, and again the value of $\alpha$ cannot be graphically obtained. It can be concluded from Figure 8 that the HF loop of the experimental impedance diagrams cannot be modeled by a R//CPE circuit, and therefore that the CPE parameters $\alpha$ and $Q$ cannot be graphically estimated.

In conclusion, the HF loop of the experimental impedance diagrams corresponds to the $CPE_{dl}||R_t^{\alpha}||W_c$ equivalent circuit (Fig. 7(c)); thus, it illustrates mainly the anodic charge transfer (diameter equal to $R_t^{\alpha}$), and its depressed shape is partly due to the CPE and partly due to the cathodic Warburg impedance in parallel. Since no plateau is visible at very HF in Figure 8(b), the effect of $W_c$ is not negligible compared to that of $CPE_{dl}$ even at $10^4$ Hz, and there is no clear frequency domain specifically assigned to each process. Whereas, the LF loop is related to the anodic mass transport and partial blocking effect by CuCl ($Z_{dl}^{\alpha}$).

The circuits of Figures 7(a) and (b) do not take into account the presence of an oxide layer, as shown by surface analysis. In fact, those circuits are in series with the impedance of the oxide film, depicted by a film resistance in parallel with a film capacitance ($R_f//C_f$). If $R_f$ is very low, then the $R_f//C_f$ circuit can be neglected and the oxide film cannot be seen by impedance. Thus, EIS data are in agreement with an oxide layer with conductive or semi-conductive properties.

On the other hand, the presence of Cu$_2$O and CuO on the 70Cu-30Ni alloy was evidenced by surface analyses. These copper oxides are well-known p-type semi-conductors [25-28], which validates the impedance models proposed in Figure 7.

As the LF loop of impedance diagrams is not well defined (described only by a few points), the single HF loop was analysed by regression of the equivalent circuit presented in Figure 7(c), in which $CPE_{dl}$, $R_t^{\alpha}$ and $W_c$ are in parallel, using Simad® software developed at Laboratoire Interfaces et Systèmes Electrochimiques. The regression results are presented in Figure 9 and in Table 2. The experimental frequency range taken into account for the regression is $0.416 \text{ Hz} - 100 \text{ kHz}$ without BSA and $0.582 \text{ Hz} - 100 \text{ kHz}$ with BSA, but the fitted curves in Figure 9 are shown in the whole frequency range ($0.002 \text{ Hz} - 100 \text{ kHz}$), with parameters values corresponding to those given in Table 2.

Here Fig. 9

Here Table 2

If the CPE behavior is assumed to be associated with surface distributed time constants for charge-transfer reactions (time-constant distribution along the electrode surface), then it is possible to apply the equation derived by Brug et al. to calculate the effective capacitance associated with the CPE [29,30]:
The capacitance values calculated from the impedance diagrams shown in Figure 2, taking for \( R_e \) and \( R_t^a \) the values extracted from the regression procedure (Table 2), are given in Table 2. These capacitance values, of the order of several tens of \( \mu F \cdot cm^{-2} \), are typical of those for a double layer capacitance, which validates the equivalent electrical circuits proposed in Figure 7. Hence, the HF loop illustrates mainly the anodic charge transfer and its diameter is equal to \( R_t^a \).

In the presence of BSA, \( R_t^a \) is 1.2 times higher than without protein. This indicates that albumin leads to a slight decrease of the corrosion current (inversely proportional to \( R_t^a \)), and hence of the corrosion rate.

4.2 Surface layers models (combined XPS and ToF-SIMS)

After short-term immersion (1 h of exposure) at \( E_{corr} \) in ASW without BSA, combined XPS and ToF-SIMS data show two oxidized layers: an outer layer mainly composed of cuprous oxide (\( Cu_2O \)) and an inner layer mainly composed of oxidized nickel. Thus, the surface layers can be depicted by the model presented in Figure 10(a). This duplex structure, with a cuprous oxide-rich outer layer and a nickel oxide-rich inner layer, has already been observed by Souchet et al. for the early stages (1-2 h) of low temperature (100-200°C) air oxidation of CuNi alloys [31-32] but this result is not well-known for the oxidation of such alloys in aqueous solution.

The presence of BSA leads to a mixed oxide layer composed of \( CuO \), \( Cu_2O \), and \( Ni(OH)_2 \) (Fig. 10(b)). Compared to the results obtained without BSA, there is a marked decrease of the \( Cu_2O \) content in the oxide layer, and its thickness is lower (~ 10 nm).

Here Fig. 10

There are basically two mechanisms of de-alloying for binary alloys proposed in the literature [16]:

1) simultaneous dissolution of both components of the alloy followed by redeposition of one component (usually the more noble one),

2) selective dissolution of one element from the alloy.

Beccaria and Crousier studied the de-alloying of Cu-Ni alloys exposed to natural seawater for 660 h and found simultaneous dissolution of both components with possible redeposition of copper for nickel contents lower than 50 %, whereas for nickel concentrations higher than 50% selective dissolution of copper took place [33]. Results obtained by Mansfeld et al. for long-term exposure (1-3 months) to natural seawater suggest that de-alloying of 70Cu-30Ni is initially due to simultaneous dissolution of copper and nickel and subsequent redeposition of copper in agreement with the results of Beccaria and Crousier [16].

Our results obtained for short-term exposure to ASW are in agreement with the conclusions drawn by Beccaria and Crousier, and Mansfeld et al.; the cuprous oxide \( Cu_2O \) detected on the surface by XPS and ToF-SIMS is formed by redeposition of dissolved copper.

\[
C_{eff} = Q^{1/\alpha} \left( R_e^{-1} + R_t^a^{-1} \right)^{(\alpha-1)/\alpha}
\]
5. Conclusions

The influence of BSA adsorption on the electrochemical behavior of 70Cu-30Ni alloy and the chemical composition of oxide layers were studied in static artificial seawater by combined electrochemical measurements and surface analyses.

From polarization curves, high anodic dissolution currents are shown (no passive current). A model is proposed to analyse electrochemical impedance data obtained at $E_{corr}$. The HF loop of the experimental diagrams illustrates mainly the anodic charge transfer, and its depressed shape is partly due to the cathodic Warburg impedance in parallel and partly due to the CPE; whereas, the LF loop is related to the anodic mass transport and partial blocking effect by CuCl. EIS measurements show uniform reactivity of the surface and are in agreement with an oxide layer with conductive or semi-conductive properties (the oxide layer cannot be seen by impedance). The BSA has a slight effect on the electrochemical behavior of the copper alloy. Thus, $E_{corr}$ value after 1 h of immersion is $\sim 30$ mV more anodic with BSA, and EIS results indicate a decrease of the corrosion current induced by the protein.

From XPS and ToF-SIMS analyses, different surface chemical compositions of 70Cu-30Ni are shown without and with BSA. In the absence of BSA, two oxidized layers can be observed: an outer layer mainly composed of copper oxide ($Cu_2O$ redeposited layer) and an inner layer mainly composed of oxidized nickel, with a global thickness of $\sim 30$ nm. In the presence of BSA, the protein is detected on the surface and the thickness of the adsorbed layer is $\sim 3$ nm, corresponding to one monolayer. The presence of BSA leads to a mixed oxide layer ($CuO$, $Cu_2O$, $Ni(OH)_2$) with a lower thickness ($\sim 10$ nm).

The combination of electrochemical measurements and surface analyses allows us to conclude that in static conditions the BSA induces a decrease of the dissolution rate at $E_{corr}$ and hence a decrease of the amount of redeposited $Cu_2O$ and of the oxide layer thickness.

As the water of cooling circuits is typically under flow and as the corrosion of copper is controlled by mass transport processes to and from corroding surfaces, electrochemical measurements in well-controlled hydrodynamics conditions, using a rotating electrode, have been performed. These experiments, which clarify the role of mass transport on the electrochemical behavior and strengthen the proposed impedance model, will be the purpose of a forthcoming paper.

Acknowledgements

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References


Tables

**Table 1: XPS atomic ratios.** Atomic ratios calculated from the XPS N 1s and C 1s core level spectra recorded for the BSA powder, and for 70Cu-30Ni after 1 h of immersion at $E_{\text{corr}}$ in aerated artificial seawater without and with 20 mg.L$^{-1}$ of BSA.

<table>
<thead>
<tr>
<th></th>
<th>N/Ctotal$^{(a)}$</th>
<th>N/(C2+C3)$^{(a)}$</th>
<th>C1/Ctotal$^{(a)}$</th>
<th>C2/Ctotal$^{(a)}$</th>
<th>C3/Ctotal$^{(a)}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>BSA powder</td>
<td>0.22</td>
<td>0.48</td>
<td>0.54</td>
<td>0.26</td>
<td>0.20</td>
</tr>
<tr>
<td>ASW</td>
<td>0.03</td>
<td>0.14</td>
<td>0.78</td>
<td>0.13</td>
<td>0.09</td>
</tr>
<tr>
<td>ASW+BSA</td>
<td>0.20</td>
<td>0.45</td>
<td>0.56</td>
<td>0.22</td>
<td>0.22</td>
</tr>
</tbody>
</table>

(a)The atomic ratio $X/Y$ is given by:

$$X / Y = I_X \sigma_Y \lambda_{X,Y}^{BSA} T_Y / I_Y \sigma_X \lambda_{X,Y}^{BSA} T_X$$

where $I_{X,Y}$ is the intensity of the peak (peak surface area) associated to element X or Y ($X$ and $Y$ equal to N or C), $\lambda_{X,Y}^{BSA}$ the attenuation length of photoelectrons emitted by the X or Y 1s core level in the BSA layer, $\sigma_{X,Y}$ the photoionisation cross-section of X or Y 1s, and $T_{X,Y}$ the transmission factor of X or Y 1s. The C 1s signal is fitted with three contributions C1, C2 and C3, corresponding to well identified carbon bonds present in the BSA molecule.
Table 2: Regression results and effective capacitance. Parameters values (electrolyte resistance $R_e$, anodic charge transfer resistance $R_t^a$, constant of the cathodic Warburg impedance $k_c$, and CPE parameters $\alpha$ and $Q$) obtained from the regression of the equivalent circuit presented in Figure 7(c) to experimental impedance data shown in Figure 2, and effective capacitance $C_{eff}$ associated with the CPE calculated from Eq. (11).

<table>
<thead>
<tr>
<th></th>
<th>$R_e$ / $\Omega$.cm$^2$</th>
<th>$R_t^a$ / $\Omega$.cm$^2$</th>
<th>$k_c$ / s$^{0.5}$.$\Omega$-1.cm$^{-2}$</th>
<th>$\alpha$</th>
<th>$Q$ / F.cm$^{-2}$.s$^{(\alpha-1)}$</th>
<th>$C_{eff}$ / $\mu$F.cm$^{-2}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Without BSA</td>
<td>12</td>
<td>1410</td>
<td>2.5$\times$10$^{-4}$</td>
<td>0.78</td>
<td>1.97$\times$10$^{-04}$</td>
<td>37</td>
</tr>
<tr>
<td>With BSA</td>
<td>12</td>
<td>1700</td>
<td>1.2$\times$10$^{-4}$</td>
<td>0.76</td>
<td>2.01$\times$10$^{-04}$</td>
<td>30</td>
</tr>
</tbody>
</table>
Figure captions

Figure 1: Voltammograms. (a) Cathodic and (b) anodic voltammograms of 70Cu-30Ni after 1 h of immersion at $E_{corr}$ in aerated artificial seawater, (—) without BSA, and (— — —) with 20 mg.L$^{-1}$ of BSA. Scan rate: 0.5 mV.s$^{-1}$.

Figure 2: Experimental Nyquist diagrams. Experimental impedance diagrams in the complex plane of 70Cu-30Ni plotted at $E_{corr}$ after 1 h of immersion in aerated artificial seawater, (○) without and (●) with 20 mg.L$^{-1}$ of BSA.

Figure 3: Cu 2p XPS spectra and Cu Auger lines. (a) X-ray photoelectron spectroscopy (XPS) Cu 2p core level spectra and (b) Cu L$_3$M$_{45}$M$_{45}$ Auger lines of 70Cu-30Ni after 1 h of immersion at $E_{corr}$ in aerated artificial seawater without and with 20 mg.L$^{-1}$ of BSA. The intensity is expressed in arbitrary unit (a.u.).

Figure 4: Ni 2p XPS spectra. X-ray photoelectron spectroscopy (XPS) Ni 2p core level spectra of 70Cu-30Ni after 1 h of immersion at $E_{corr}$ in aerated artificial seawater without and with 20 mg.L$^{-1}$ of BSA. The intensity is expressed in arbitrary unit (a.u.).

Figure 5: ToF-SIMS depth profile. Characteristic time-of-flight secondary ions mass spectrometry (ToF-SIMS) negative depth profile of 70Cu-30Ni after 1 h of immersion at $E_{corr}$ in aerated artificial seawater without BSA.

Figure 6: N 1s and C 1s XPS spectra. X-ray photoelectron spectroscopy (XPS) (a) N 1s and (b) C 1s core level spectra of 70Cu-30Ni after 1 h of immersion at $E_{corr}$ in aerated artificial seawater with 20 mg.L$^{-1}$ of BSA. Solid line: experimental spectra; dashed line: peak decomposition. The intensity is expressed in counts per second (CPS).

Figure 7: Impedance models. Equivalent electrical circuits to model the 70Cu-30Ni/ASW system: (a) general circuit, (b) simplified circuit taking into account experimental cathodic voltammograms, and (c) circuit used to analyse the HF loop of experimental impedance diagrams. $R_e$ is the electrolyte resistance, $CPE_{dl}$ a constant phase element related to the double layer, $R^a_t$ the anodic charge transfer resistance, $Z_{D^c}$ an impedance that illustrates anodic mass transport and partial blocking effect by CuCl, $R^c_t$ the cathodic charge transfer resistance, $Z_{D^c}$ a cathodic impedance that illustrates O$_2$ mass transport, and $W_c$ the cathodic Warburg impedance.

Figure 8: Experimental impedance data. Experimental impedance data of 70Cu-30Ni obtained at $E_{corr}$ after 1 h of immersion in aerated artificial seawater, (□) without and (●) with 20 mg.L$^{-1}$ of BSA. (a) Imaginary part of the impedance as a function of frequency, and (b) derivative curves calculated from Figure 8(a) ($d \log |Z_j| / d \log f$ vs $\log f$). Same data as in Figure 2.

Figure 9: Fit of the impedance model to the data. High frequency loops of Nyquist diagrams obtained for 70Cu-30Ni at $E_{corr}$ after 1 h of immersion in aerated artificial seawater, (□) without and (●) with 20 mg.L$^{-1}$ of BSA. (□, ●) Experimental curves and (■, ○) fit of the impedance model presented in Figure 7(c) to the data. Same data as in Figures 2 and 8.
**Figure 10: Surface layers models.** Models of the surface layers deduced from combined X-ray photoelectron spectroscopy (XPS) and time-of-flight secondary ions mass spectrometry (ToF-SIMS) results for 70Cu-30Ni after 1 h of immersion at $E_{corr}$ in aerated artificial seawater (a) without and (b) with 20 mg.L$^{-1}$ of BSA.
Figure 1

(a) 

(b)
Figure 2

\[ Z_j / \Omega \cdot \text{cm}^2 \]
\[ Z_r / \text{k} \Omega \cdot \text{cm}^2 \]

-0.416 Hz
-0.002 Hz

Figure 2
Figure 3

(a) Copper 2p spectroscopy showing binding energies with and without BSA.

(b) Protein binding energy with BSA and without BSA at different intensities.
Figure 4

The figure shows a comparison of Ni 2p intensity plots with and without BSA. The plots are labeled as "With BSA" and "Without BSA" respectively. The binding energy is given in eV, and the intensity is in arbitrary units (a.u.). Peaks are labeled as Ni 2p\(_{1/2}\) and Ni 2p\(_{3/2}\).
Figure 5

![Graph showing intensity counts vs. sputtering time for different oxidation states: Cu oxide, oxidized Ni interface, Cu oxide / oxidized Ni interface, oxide / alloy interface.](image)

Key:
- $95^{65}\text{CuO}_2$
- $90^{59}\text{NiO}_2$
- $18^{18}\text{O}$
- $116^{58}\text{Ni}_2$

Intensity / counts

Sputtering time / s
Figure 6

(a) and (b) show the binding energy distribution and intensity/counts per second (CPS) plots, respectively. The peaks at specific binding energies (eV) correspond to different carbon species (C1, C2, C3).
Figure 7
Figure 8
Figure 10

(a) 93% Cu$_2$O, 7% Ni(OH)$_2$
Cu$_2$O

~ 18 nm

~ 12 nm

NiO and/or Ni(OH)$_2$

ALLOY

68% Cu, 32% Ni

(b) BSA

~ 3 nm

~ 10 nm

44% CuO, 11% Cu$_2$O, 45% Ni(OH)$_2$

ALLOY

68% Cu, 32% Ni
Vitae

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