# THE EFFECT OF CYSTEINE ON THE CORROSION CHARACTERISTICS OF BIOALLOY

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#### **ABSTRACT**

Many alloys exhibit the shape memory effect, but only the Cu-Al-Zn, Cu-Al-Ni, and Ni-Ti alloys are presentand of commercial importance. The greatest use of these alloys is in biomedicine (filters, orthodontic implants, guide wires for catheters, etc.). Shape-memory alloys may be found in sensors in automobiles, consumer products, and generally in smart materials.

The electrochemical behavior of Cu-Al-Zn alloy in phosphate buffer without and in the presence of cysteine was investigated.

Electrochemical researchwas performed in a traditional three-electrode system by means of the Tafel extrapolation method. The results showed that the presence of cysteine leads to a decrease in the corrosion rate and the density (values) of the corrosion current, which indicates that the tested inhibitor is efficient. In other words, it provides alloys with protection against corrosion.

## 1. INTRODUCTION

Shape memory alloys (SMA) are materials that have the ability to return to a former shape when subjected to an appropriate thermomechanical procedure. Pseudoelastic and shape memory effects are some of the behaviors presented by these alloys [1].

The key transformation responsible for the behavior of shape-memory alloys is martensitic transformation. Martensitic transformation is a reversible crystallographic reorientation process between two stable phases. The transformation takes place when the Gibbs energy of martensite becomes lower than the Gibbs energy of austenite at a temperature below the critical temperature t0, which represents the temperature of thermodynamic equilibrium at which the Gibbs energies of both phases are equal [2,3].

The chemical composition of the alloy, the heat treatment procedure, the cooling rate, the grain size, and the number of transformation cycles are the main factors that influence the transformation temperature of shape-memory alloys [3].

During the phase transformation, several physical properties of the material change.

The transformation of austenite into martensite is associated with the release of heat (exothermic phase transformation). The reverse transformation of martensite into austenite is associated with supplying energy for the reaction (endothermic phase transformation). When a shape memory alloy undergoes a phase transformation, it transforms from a highly ordered phase (austenitic phase) to a low-ordered phase (martensitic phase).

NiTi alloys possess good properties of shape memory effect, pseudoelasticity, and biocompatibility. However, these alloys are significantly more expensive than copper-

based shape memory alloys. Right after NiTi, the second commercially applied shape-memory alloy is the Cu-Al-Zn alloy, which is cheaper and simpler to manufacture, and has better electrical and thermal conductivity. The disadvantage compared to the NiTi alloy is that the Cu-Al-Zn alloy has worse shape memory properties.

However, shape memory Cu-Al-Zn alloys show a significant shape memory effect in a certain range of chemical compositions.

Metal biomaterials have by far been used in biomedicine for the longest time. Numerous biomaterials and medical devices are commonly used today as prosthetic devices in dental, orthopedic, cardiovascular, ophthalmological, and reconstructive surgery. They are also successfully used in interventions, such as angioplasty (stents) and hemodialysis (membranes), for surgical sutures or bioadhesives, but also as devices for the controlled release of drugs. E.g. they are used for fixing and replacing hard tissue (artificial hip, artificial knee, plates for fixing broken bones, various applications in dentistry, etc.) and for making surgical instruments.

The reactions of living organisms to biomaterials are different; under certain circumstances, some material is well accepted by the organism, while, under other circumstances, the same material is not accepted by the organism [4,5]. Corrosion is defined as the destructive and unintentional degradation of a material caused by its environment. Today, corrosion is one of the important causes of the global material and energy crisis and is the cause of significant losses in the economy of every country [6]. The harmful effects of corrosion can cause numerous negative consequences, such as to lead to a decrease in the utility value of materials, more expensive maintenance and shortened durability of structures, production stoppages, accidents, etc.

Corrosion control of metal is of technical, economical, environmental, and aesthetical importance. The use of inhibitors is the best way to prevent metal and alloys from corrosion [7].

A corrosion inhibitor is a substance applied to an environment that significantly reduces the corrosion rate of materials (especially metals) exposed to that environment.

Organic compounds, predominantly, those with O, N, S, P, and/or  $\pi$  electrons in their molecular structure have received considerable attention as metal corrosion inhibitors. It is believed that organic compounds get adsorb onto a metal surface through their active center (heteroatom or  $\pi$ -bond) and form a protective layer that prevents the corrosive agents in the aggressive environment from gaining access to the metal surface.

Compared to the inorganic metal corrosion inhibitors, the organic inhibitors are less toxic to both humans and the environment making discontinuation not to be the best option. [8,9].

Cysteine has the ability to control the corrosion of various metals. Generally, cysteine is a very interesting amino acid that contains the amino group [-NH<sub>2</sub>], carboxyl group [-

COOH], and thiol group [-SH]. It can coordinate with metals through the nitrogen atom, the oxygen atom of the carboxyl group, and Sulphur atom of the thiol group [10].

Phosphate buffers are widely used because they help maintain a constant pH level in a particular environment. Generally speaking, most researchers using phosphate buffers try to maintain a pH of 7.4 because the properties closely match those of the human body.

Compounds that contain polar functional groups that allow them to be efficiently adsorbed have found application in corrosion inhibition because of their effectiveness. Amino acids are also included in such compounds [11].

Amino acids are considered green inhibitors, and they are characterized by non-toxicity, biodegradability, cheap production and a high degree of purity of the produced compound, and solubility in aqueous media [12].

The presence of heteroaromatics in the structure of amino acids (they contain at least one carboxylic acid and an amino group, usually attached to the same carbon) increases their ability to be used as corrosion inhibitors. Previous tests have shown that the effectiveness of inhibition primarily depends on the molecular structure of the amino acid being tested, the size of the molecule, and the type of interaction with the metal surface [13,14]. The highest efficiency of inhibition in the presence of cysteine was attributed to the fact that the amino acid containing S can be adsorbed as a bidentate ligand, whereby coordination takes place both through the amino group (or carboxyl) and through the –SH group [15].

## 2. EXPERIMENTAL

The aim of this study is to focus on the effect of cysteine concentration (2 mM, 4 mM, 8 mM) on the corrosion behavior of Cu-18,05Zn-5,35Al bioalloys in phosphate buffer, pH = 7. This tested bioalloy was prepared using solution annealing at a temperature of  $850 \, ^{0}$ C for 10 min., then quenched into water at  $15 \, ^{0}$ C.

Phosphate buffer was prepared by dissolving 3,405 g of potassium dihydrogen phosphate (KH<sub>2</sub>PO<sub>4</sub>) in distilled water in a 500 mL volumetric flask and 4,450 g disodium hydrogen phosphate (Na<sub>2</sub>HPO<sub>4</sub>) in distilled water in a volumetric flask. A pH value of 7,00 was adjusted by adding 100 mL KH<sub>2</sub>PO<sub>4</sub> and 150 mL Na<sub>2</sub>HPO<sub>4</sub> to a 250 mL volumetric flask. A 0,5mol/L cysteine solution was prepared by dissolving 1,5145 g of cysteine in distilled water in a 25 mL volumetric flask.

The sample was prepared for electrochemical research after polishing with grinding paper, degreasing in ethanol, and washing with distilled water. The work surface of the tested bioalloy was 25 cm<sup>2</sup>.

Electrochemical researchwas performed in a traditional three-electrode system by means of the Tafel extrapolation method. Tafel extrapolation is a mathematical technique used to estimate either the corrosion current ( $I_{corr}$ ) or the corrosion potential ( $E_{corr}$ ) in an electrochemical cell, and by extension, the corrosion rate [16]. The three-electrode system consists of a working electrode, a counter electrode, and a reference electrode. The most common lab reference electrodes are the Saturated Calomel Electrode and the Ag/AgCl electrode.

Potentiostat/galvanostat Vertex one, Ivium Technologiesis being used for electrochemical testing of corrosion rate.

#### 3. RESULTS AND DISCUSSION

The corrosion current density ( $I_{\rm corr}$ ) and corrosion potential ( $E_{\rm corr}$ ) were obtained by employing the Tafel extrapolation method. The Tafel extrapolation line is based on the polarization curve obtained when the overpotential is between -0,300 V do 0,05 V. The scan rate was 20 mV/s.

The efficiency of the inhibitor, Ei (%), was calculated based on following equation:

$$Ei = \frac{\pi - \pi'}{\pi} \tag{1}$$

where:  $\pi$  - corrosion rate in solution without corrosion inhibitor (mm year<sup>-1</sup>)  $\pi$ '- corrosion rate in solution with corrosion inhibitor (mm year<sup>-1</sup>).

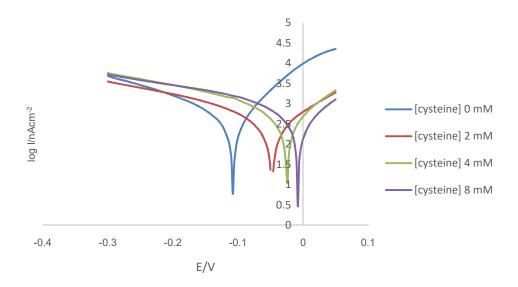


Figure 1. Tafel extrapolation graph of the effect of cysteine concentration on the corrosion behavior of Cu-18,05Zn-5,35Al bioalloy

Table 1. The corrosion parameters determined by Tafel extrapolations method

Inhibitor	c <sub>inh.</sub> (mM)	E <sub>corr.</sub> (V)	I (A)	I (A/cm <sup>2</sup> )	$R_{p}\left(\Omega\right)$	$\beta_a$ (V/dec)	$\beta_k$ (V/dec)	v <sub>corr.</sub> (mm/yr .)
-	0	-0,1034	4,65×10 <sup>-7</sup>	4,65×10 <sup>-6</sup>	$5,09 \times 10^4$	0,078	0,182	0,05405
Cysteine	2	-0,0209	$3,75\times10^{-7}$	$3,75\times10^{-6}$	$8,15\times10^{4}$	0,095	0,272	0,04364
Cysteine	4	-0,0048	5,96×10 <sup>-7</sup>	5,96×10 <sup>-6</sup>	$4,15\times10^{4}$	0,070	0,298	0,06928
Cysteine	8	0,0290	7,07×10 <sup>-7</sup>	7,07×10 <sup>-6</sup>	$3,06\times10^{4}$	0,058	0,374	0,08221

Based on the values of corrosion parameters determined by Tafel extrapolation (Table 1) it was confirmed that cysteine can be applied in order to reduce the corrosion rate. In the investigated concentration range of the cysteine in phosphate buffer, the lowest rate of corrosion is at the concentration of inhibitor(cysteine) of 2 mmol/L and which amounts to 0.04364 mm/year and its effectiveness amounts to 19,10%.

By further increasing of the concentration of cysteine, the inhibitory effect decrease, because the corrosion rate is greater than the rate of corrosion of the examined bioalloy without the presence of the inhibitor. In other words, at a concentration of 4 mmol/L and 8 mmol/L inhibitor functions as an activator because the corrosion rate is higher than the rate of corrosion without an inhibitor.

## 4. CONCLUSION

As the obtained results have shown, the presence of cysteine in concentration 2 mmol/L leads to a decrease in the corrosion rate and the density (values) of the corrosion current, as well as to an increase in polarization resistance values, which indicates that the corrosion inhibitor is efficient in appropriate concentration.

#### 5. REFERENCES

- [1] Machado LG, Savi MA.: Medical applications of shape memory alloys. Braz J Med Biol Res. 2003 Jun;36(6):683-91. doi: 10.1590/s0100-879x2003000600001. Epub 2003 Jun 3. PMID: 12792695., 2003
- [2] I. Ivanić, M. Gojić, S. Kožuh. Slitine s prisjetljivosti oblika (I. dio): najznačajnija svojstva, *Kemija u industriji* 63. 2014. 9-10; 323-329

- [3] K. Otsuka, C. M. Wayman. Shape memory materials. University of Cambridge, Cambridge. 1998
- [4] Tanikić, D. I., Manić, M. T., Ranđelović, S. S., Brodić, D. T. Legure koje pamte oblik i njihova upotreba u medicini. *Vojno tehničk iglasnik*. 2014 62(4), 59-71
- [5] L. G. Machado, M. A. Savi. Medical applications of shape memory alloys. *Braz. J. Med. Biol. Res.* 2003 36, str. 683–691
- [6] Juraga, I., Alar, V., Šimunović, V., Stojanović, I. *Korozija i metode zaštite od korozije*. Zagreb: Fakultet strojarstva i brodogradnje. 2012
- [7] Manoj Acharya, Jinendra Singh Chouhan, Anita Dixit, D. K. Gupta: Green Inhibitors for Prevention of Metal and Alloys Corrosion: An Overview, 2013
- [8] S. A. Umoren, M. M.Solomon: Synergistic corrosion inhibition effect of metal cations and mixtures of organic compounds: A Review, Journal of Environmental Chemical Engineering, 5(1), 246-273., 2017
- [9] Malinović B., Djuričić T., Zorić D.: Corrosion behaviour of stainless steel EN 1.4301 in acid media in presence of PBTCA inhibitor, University of Banja Luka, Faculty of Technology, Banja Luka, 2020
- [10] A. Sahaya Raja, R. Venkatesan, R. Sonisheeba, J. Thomas Paul raj S. Sivakumar, P. Angel, J. Sathiyabama: Corrosion Inhibition by Cysteine An Over View, Corrosion Research Centre, PG and Research Department of Chemistry G.T.N Arts College, Dindigul, TN, India, 2014
- [11] Antonijević M. M., Petrović M. B. Inhibitori korozije bakra. Zaštita materijala. 2007.,48
- [12] B. El Ibrahimi, Jmiai A., Bazzi L., El Issami S. Amino acids and their derivatives as corrosion inhibitors for metals and alloys. *Arabian Journal of Chemistry*. 2017, str. 1878-5352
- [13] Radovanović, B. M. *Uticaj organskih inhibitora na koroziono ponašanje mesinga u rastvoru natrijum sulfata*. Doktorska disertacija. Univerzitet u Beogradu. Tehničkifakultet u Boru. 2012
- [14] D. Q. Zhang, Q. R. Cai, X. M. He, L. X. Gao and G. D. Zhou. *Mater. Chem. Phys.* 2008 *112*, str. 353
- [15] Ivanov, S., Rajčić Vujasinović M., Stević, Z. Uticaj hloridnih jona na koroziono ponašanje hladno deformisane bakarne žice u alkalnoj sredini, *Zaštita materijala*, 2006., *Vol.47*, str. 33-38
- [16] Neeraj K Namboodiri, Tafel extrapolation method, 2020