



Review

Influence of Parameters and Regimes of the Electrodeposition on Hardness of Copper Coatings

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Abstract: Correlation among morphological, structural and hardness characteristics of electrodeposited copper coatings is presented in this review paper. Cu coatings were produced applying constant galvanostatic (DC) and pulsating current (PC) regimes on hard silicon (Si(111)) and brass substrates. The parameters of the electrochemical deposition were analyzed, which include the kinds of electrolyte and cathode, the coating thickness and the electrolyte stirring, as well as the parameters defining PC regime, such as the average current density and the current density amplitude, were analyzed. Morphology and structure of Cu coatings were examined by scanning electron microscope (SEM), atomic force microscope (AFM) and by X-ray diffraction (XRD), while hardness was examined by Vickers microindentation. The coatings of Cu on both Si(111) and brass cathodes belong to “soft film (coating) on hard substrate” composite hardness system, and the Chicot–Lesage (C–L) composite hardness model was applied to estimate a hardness of the Cu coatings. Analyzing the examined parameters and regimes of electrodeposition, the critical relative indentation depth (RID)_c of 0.14 has been defined by the C–L model. Based on done analyses, it is shown that this RID value, separating a zone where measured hardness corresponds to the coating hardness and a zone where it is necessary to apply the C–L model to determine an absolute hardness of the Cu coatings, has an universal character for the electrolytically produced Cu coatings on Si(111) and brass substrates.

Keywords: copper; electrodeposition; morphology; structure; hardness; the Chicot–Lesage composite hardness model



Citation: Mladenović, I.O.; Nikolić, N.D. Influence of Parameters and Regimes of the Electrodeposition on Hardness of Copper Coatings. *Metals* **2023**, *13*, 683. <https://doi.org/10.3390/met13040683>

Academic Editors: Belén Díaz Fernández and Marcello Cabibbo

Received: 22 February 2023

Revised: 16 March 2023

Accepted: 28 March 2023

Published: 30 March 2023



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1. Introduction

The coatings or films of copper obtained by various production methods occupy an attention of both scientific and technological communities. The cause of this attention lies in specific features of Cu, which include high thermal and electrical conductivity, malleability, corrosion resistance, and good adhesion with a substrate [1]. Thanking to these characteristics, copper coatings are used in various industries such as aerospace, automotive, electronics and telecommunications. All ways of copper coating production on conductive and non-conductive substrates can be divided in two groups: non-electrochemical and electrochemical methods. The main non-electrochemical methods are electroless plating, chemical vapour deposition (CVD), physical vapour deposition (PVD), thermal spray and sputtering techniques [2,3]. Plasma spray is a special kind of thermal spray technique that is used for a formation of various metallic and non-metallic materials, such as nitrides, carbides, and metal oxides [4]. This technique is not suitable for temperature-sensitive materials such as polymers and polymer matrix composites. The cold spray [5,6] and electroless plating [2,3] techniques are suitable for a metallization of temperature-sensitive materials.

Electrochemical deposition processes, or shortly, electrodeposition processes found wide application in obtaining of metal coatings of various morphological, structural and mechanical characteristics on conductive substrates. They belong to environmentally friendly, low cost, time saving and facile methods for a production of metal coatings of desired quality at both micro and nano scale [7]. The desired characteristics of coatings are

achieved by selection of regimes and parameters of the electrodeposition. For an application in industry, current regimes, including the constant galvanostatic (DC), the pulsating current (PC) and the reversing current (RC) regimes are used. Although the potentiostatic regimes, involving the constant potentiostatic and the pulsating overpotential (PO) regime as the type of periodically changing regime, are important in academic investigations in examination of mechanism of metal electrodeposition, they give important information necessary for a transfer of knowledge towards application at the industrial scale. The quality of coatings depends on the following parameters of the electrodeposition: the kind of electrolyte, the kind of cathode, temperature of electrolyte, way of electrolyte stirring, electrodeposition time (coating thickness), current density of the electrodeposition, etc. [8].

The main convenience of electrodeposition processes is a good adhesion of coatings with the electrode surface and an easy control of coating thickness by a choice of parameters and regimes of the electrodeposition; however, an impossibility of deposition on non-conductive substrates is the main shortfall of this technique [8]. On the other hand, non-electrochemical methods are suitable for a formation of coatings on non-conductive substrates, but these methods generally give coatings of worse adhesion than the electrochemical methods and show certain limitations related with the coating thickness [5].

The obtaining of compact and uniform Cu coatings of good mechanical characteristics is a huge challenge. The mechanical features of the materials are those associated with an ability of the material to resist mechanical forces and loads. The thirteen main mechanical properties of metals are as follows: strength (yield, fracture, ultimate), elasticity, plasticity, ductility (elongation, area reduction), malleability, brittleness, stiffness, hardness (macro, micro and nano), creep, fatigue, resilience, toughness, and weldability [9]. Among all these mechanical characteristics, the coating hardness is one of the most important for an application of Cu coatings in above mentioned industries.

In this review paper, comprehensive analysis of influence of various parameters of the electrodeposition on hardness of electrodeposited copper coatings has been presented. The constant galvanostatic (DC) and the pulsating current (PC) regimes are used to produce Cu coatings. The influence of the following electrodeposition parameters will be presented: the kind of electrolyte, the kind of cathode, the thickness of coating, and electrolyte stirring. The following parameters of the PC regime were examined: the average current density (frequency) and the current density amplitude. All electrodepositions were performed on hard Si(111) and brass substrates. Hardness of the Cu coatings was determined applying the Chicot–Lesage (C–L) composite hardness model. This analysis will be presented through the following Sections:

- Influence of parameters of the electrodeposition on morphology and structure of Cu coatings produced by the DC regime—Section 2,
- Influence of parameters of the electrodeposition on hardness of Cu coatings produced by the DC regime—Section 3,
- Influence of parameters of the PC regime on morphology and structure of electrolytically produced Cu coatings—Section 4,
- Influence of parameters of the PC regime on hardness of electrolytically produced Cu coatings—Section 5, and
- Correlation among morphology, structure and hardness of electrolytically produced copper coatings—Section 6.

In addition, the short literature survey of Cu coatings obtained with the reinforcements (Copper matrix composites) is presented in Section 7.

2. Influence of Parameters of the Electrodeposition on Morphology and Structure of Cu Coatings Produced by the DC Regime

2.1. Influence of the Kind of Electrolyte

The kind of electrolyte plays a crucial role in creating morphological and structural characteristics of metal coatings with strong consequences on a hardness of the produced coatings. Depending on it, the coatings with micro and nano characteristics can be obtained.

In the case of Cu, the coatings at the nano level are obtained with the help of specific substances known as leveling/brightening additives added to the electrolyte. In the twenty-first century, a combination of additives including chloride ions, polyethylene glycol (PEG) and 3-Mercapto-1-propanesulfonic acid (MPSA) found wide application in electrodeposition of Cu coatings of nanostructured characteristics with mirror bright appearance [10,11]. In this review paper, the two types of electrolytes are analyzed [12]:

(I) the basic sulfate electrolyte composition of $240 \text{ g L}^{-1} \text{ CuSO}_4 \cdot 5 \text{ H}_2\text{O}$ in $60 \text{ g L}^{-1} \text{ H}_2\text{SO}_4$; this electrolyte is denoted as *electrolyte I*,

(II) the electrolyte with added leveling/brightening additives: the composition of this electrolyte is $240 \text{ g L}^{-1} \text{ CuSO}_4 \cdot 5 \text{ H}_2\text{O}$, $60 \text{ g L}^{-1} \text{ H}_2\text{SO}_4$, $0.124 \text{ g L}^{-1} \text{ NaCl}$, $1 \text{ g L}^{-1} \text{ PEG 6000}$ (polyethylene glycol), and $0.0015 \text{ g L}^{-1} \text{ MPSA}$ (3-Mercapto-1-propanesulfonic acid); this electrolyte is denoted as *electrolyte II*.

Figure 1 shows the morphologies of Cu coatings electrodeposited in the DC mode at a current density (j) of 50 mA cm^{-2} from the basic sulfate electrolyte (*electrolyte I*), (Figure 1a) and the electrolyte including additives for leveling and brightness (*electrolyte II*), (Figure 1b) on brass cathodes. The thickness of coatings is $20 \text{ }\mu\text{m}$, and magnetic stirring (MS) of electrolyte was used during the electrochemical deposition processes. The other electrodeposition conditions for production of these coatings are presented in Ref. [12].

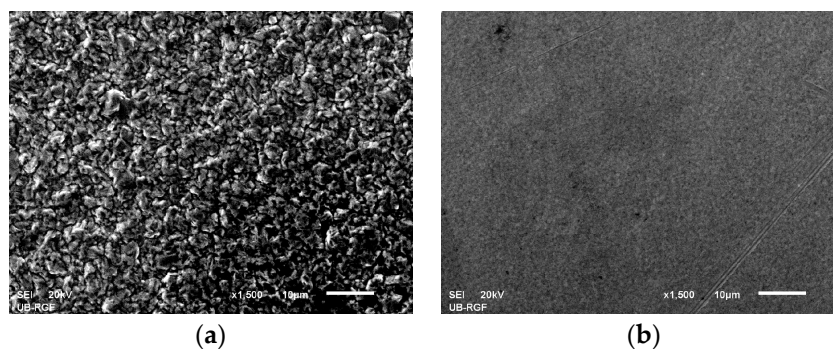


Figure 1. The SEM micrographs of the morphology of Cu coatings electrodeposited at a current density of 50 mA cm^{-2} on the brass cathode from: (a) *electrolyte I*, and (b) *electrolyte II*. The thickness of the coatings: $20 \text{ }\mu\text{m}$; The electrolyte stirring: Magnetic stirring (MS); Temperature: $22.0 \pm 0.50 \text{ }^\circ\text{C}$. Magnification: $\times 1500$ [12].

The fine-grained coating with mat appearance was produced from the basic sulfate electrolyte (Figure 1a), and the very smooth coating with mirror bright appearance was produced from the electrolyte with additives (Figure 1b). In the mirror bright coating, it is not noticed clear boundary among grains.

The topography and the histogram of the Cu coating produced from *electrolyte I* are presented in Figure 2a,b, respectively, while for the coating produced from *electrolyte II* are presented in Figure 2c,d, respectively.

The AFM analysis of the given coatings confirmed a clear difference between them. The roughness analysis of the coatings was performed applying AFM accompanying software, which define the arithmetic average of the absolute (R_a) roughness of coatings [12]. The value of R_a roughness for the coating produced from the basic sulfate solution ($R_a = 231.60 \pm 10.6 \text{ nm}$) was incomparably larger than that produced from the solution with the additives ($R_a = 28.25 \pm 2.1 \text{ nm}$); thus, proving the strong leveling/brightening influence of an added combination of additives. The R_a roughness of the initial surface area of the brass was $31.53 \pm 2.45 \text{ nm}$ [12].

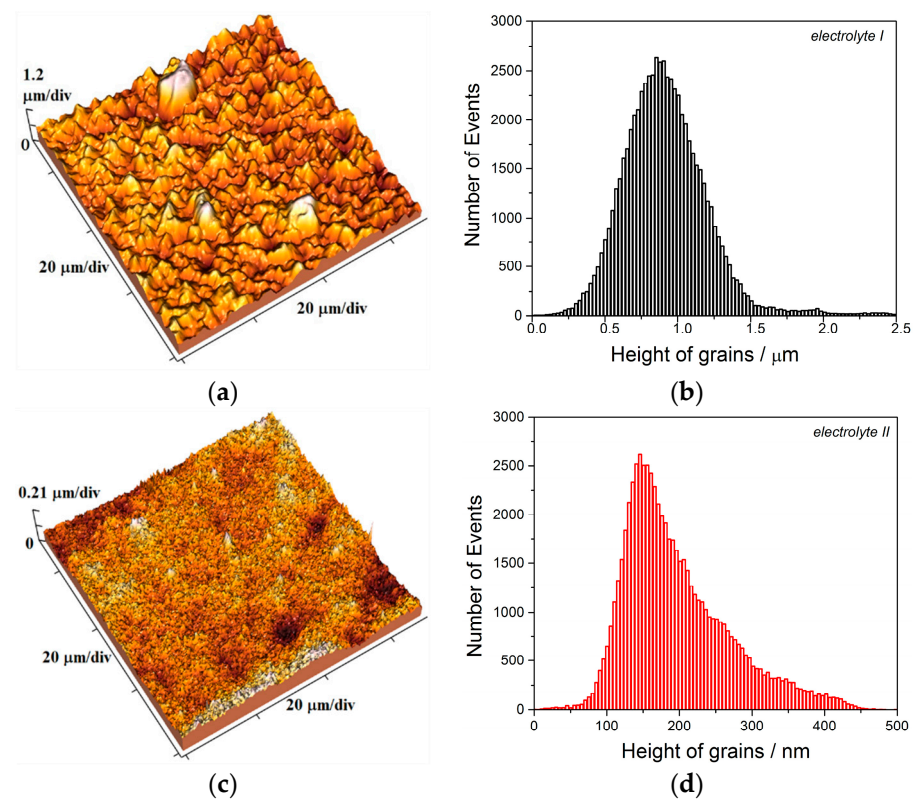


Figure 2. The 3D (three dimensional) AFM images and the corresponding histograms of the Cu coatings produced from: (a,b) *electrolyte I*, and (c,d) *electrolyte II*. The thickness of the coatings: 20 μm . The current density: $j = 50 \text{ mA cm}^{-2}$; The cathode: brass; The electrolyte stirring: Magnetic stirring (MS); Temperature: $22.0 \pm 0.50 \text{ }^{\circ}\text{C}$. Scan size: $(70 \times 70) \mu\text{m}^2$ [12].

2.2. Influence of the Type of Cathode

Morphology as one of the main characteristics of electrodeposited metals is determined by the type of solution, while parameters which include a kind of cathode and an electrolysis time (thickness) had not any significant effect on macro morphology of coatings. The best way to look at an effect of these parameters is roughness analysis of the coatings electrodeposited under the same conditions.

The 3D AFM images and histograms of the Cu coatings electrodeposited under the same conditions as those shown in Figures 1 and 2, but on the Si(111) substrate are presented in Figure 3 (Figure 3a,b—the coating electrodeposited from *electrolyte I* and Figure 3c,d—the coating electrodeposited from *electrolyte II*). The way of a preparation of Si(111) for electrochemical deposition is described elsewhere [13].

Electrodeposition was performed on Si(111) substrate of the initial R_a roughness of $24.18 \pm 2.05 \text{ nm}$ [12]. The obtained R_a values are $216.81 \pm 8.5 \text{ nm}$ for the coating electrodeposited from *electrolyte I* and $20.96 \pm 1.9 \text{ nm}$ for the coating electrodeposited from *electrolyte II*. They are comparable with the values obtained on the brass cathode. The slightly lower values are result of smoother initial surface area of the Si(111) relative to the brass.

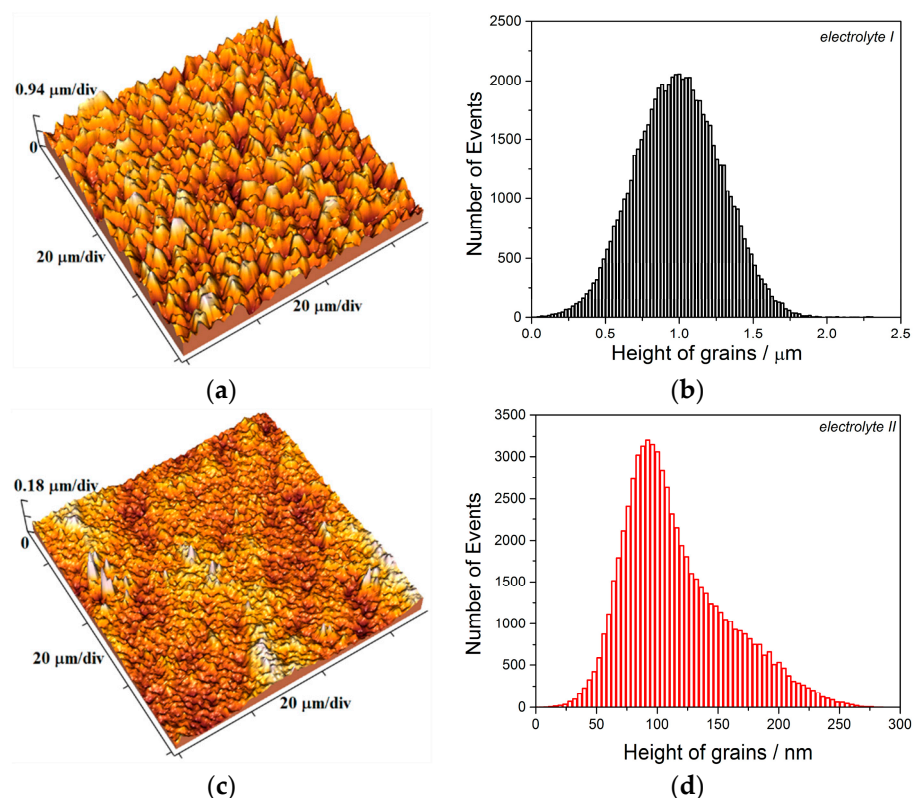


Figure 3. The 3D (three dimensional) AFM images and the corresponding histograms of the Cu coatings electrodeposited from: (a,b) *electrolyte I*, and (c,d) *electrolyte II*. The thickness of the coatings: 20 μm . The current density: $j = 50 \text{ mA cm}^{-2}$; The cathode: Si(111); The electrolyte stirring: Magnetic stirring (MS); Temperature: $22.0 \pm 0.50 \text{ }^{\circ}\text{C}$. Scan size: $(70 \times 70) \mu\text{m}^2$ [12].

2.3. Influence of Coating Thickness

Figure 4 gives AFM images and histograms of the Cu coatings thicknesses of 40 μm electrodeposited from *electrolyte I* (Figure 4a,b) and *electrolyte II* (Figure 4c,d) on Si(111) substrates. The values of R_a roughness are $317.20 \pm 8.9 \text{ nm}$ for the coating produced from *electrolyte I* and $15.36 \pm 1.5 \text{ nm}$ for the coating produced from *electrolyte II*.

The opposite trend in a change of roughness with the increase of coating thickness from 20 to 40 μm can be attributed to various mechanisms of metal electrodeposition without and with additives. The increase of R_a roughness of the coatings electrodeposited from the basic sulfate solution (*electrolyte I*) is a consequence of non-dendritic amplification of the electrode surface area with an electrolysis time [8], while decrease of R_a roughness for the coatings electrodeposited from *electrolyte II* can be ascribed to good leveling/brightening effect of these additives [10,11].

The coatings thicknesses of 40 μm were very convenient for the structural analysis. The diffractograms obtained for the coatings electrodeposited from electrolytes without (*electrolyte I*) and with added additives for leveling and brightening (*electrolyte II*) are presented in Figure 5. The diffraction peaks recorded at 2θ angles of 43.3° , 50.4° , 74.1° and 89.9° corresponding to (111), (200), (220) and (311) crystal planes confirm the belonging of a crystal lattice of Cu to the face-centered cubic (FCC) type [14]. The various strong intensities of the diffraction peaks were observed for these two types of Cu coatings: (220) crystal plane for the Cu coating electrodeposited from *electrolyte I* and (200) crystal plane for the Cu coating electrodeposited from *electrolyte II*.

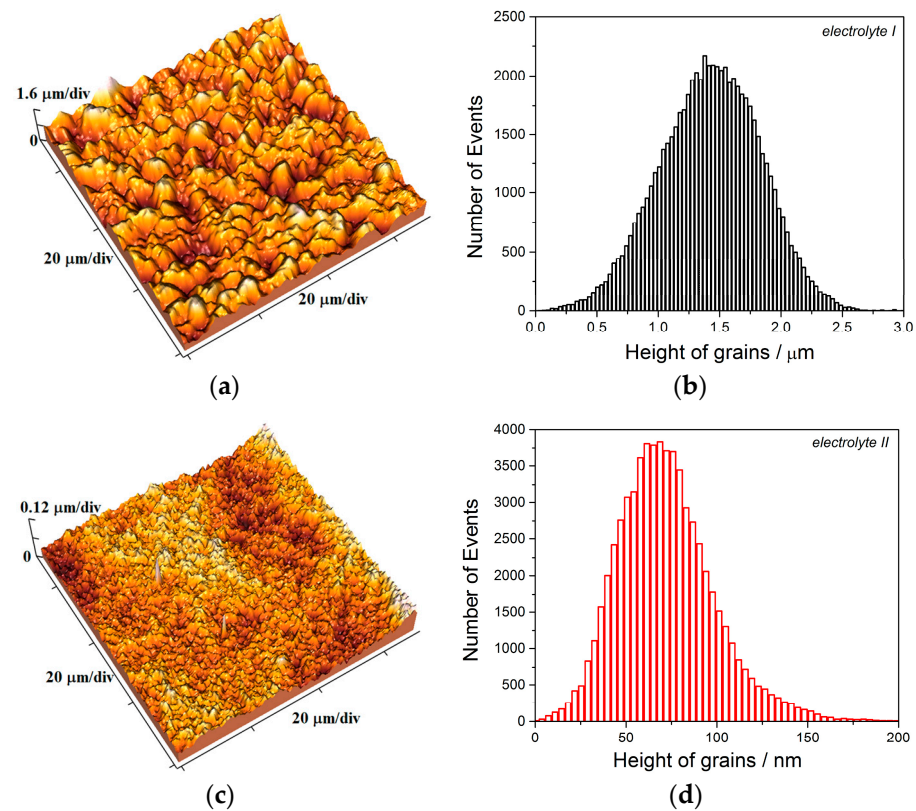


Figure 4. The 3D (three dimensional) AFM images and the corresponding histograms of the Cu coatings electrodeposited from: (a,b) *electrolyte I*, and (c,d) *electrolyte II*. The thickness of the coatings: 40 μm . The current density: $j = 50 \text{ mA cm}^{-2}$; The cathode: Si(111); The electrolyte stirring: Magnetic stirring (MS); Temperature: $22.0 \pm 0.50 \text{ }^{\circ}\text{C}$. Scan size: $(70 \times 70) \mu\text{m}^2$ [12].

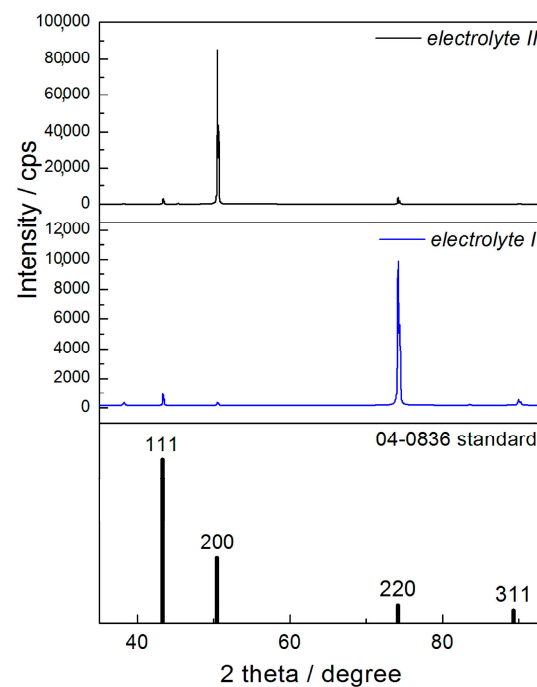


Figure 5. The diffractograms of the Cu coatings thicknesses of 40 μm produced by electrodeposition from *electrolytes I* and *II*, and Cu standard (04-0836) [12].

The preferred orientation of the Cu coatings was precisely estimated by a determination of two types of texture coefficients: the texture coefficients, $TC(hkl)$, and the relative texture coefficients, $RTC(hkl)$ [12–15]. The detailed procedure for a calculation of these coefficients can be found in Refs. [13,16]. The $TC(hkl)$ coefficients larger than 1 are a proof of the preferred orientation in a given crystal plane [12–16]. As regards $RTC(hkl)$ coefficients, the existence of a preferred orientation in any crystal plane is closely related with number of crystal planes under consideration. In this case, since four crystal planes are considered, the values of $RTC(hkl)$ coefficients larger than 25% indicate the existence of the preferred orientation.

Table 1 gives the values of $TC(hkl)$ and $RTC(hkl)$ coefficients for considered Cu coatings. The strong preferred orientation in (220) crystal plane for the coating electrodeposited from *electrolyte I* and the strong preferred orientation in (200) crystal plane for the coating electrodeposited from *electrolyte II* is confirmed by a calculation of these coefficients.

Table 1. The values of TC and RTC coefficients calculated for the Cu coatings electrodeposited from *electrolyte I* and *electrolyte II*; (electrolyte I—I; electrolyte II—II; s—Cu standard). R is an intensity of the diffraction peak [12].

Plane (hkl)	R (in%)		R_s (in%)	TC		RTC (in%)	
	R_I	R_{II}		TC_I	TC_{II}	RTC_I	RTC_{II}
(111)	7.9	3.31	54.6	0.145	0.061	1.71	1.47
(200)	3.2	92.12	25.1	0.127	3.67	1.49	88.31
(220)	84.0	4.05	10.9	7.71	0.37	90.67	8.90
(311)	4.9	0.52	9.4	0.521	0.055	6.13	1.32

2.4. Influence of Electrolyte Stirring

In the DC regime, it is necessary to apply an electrolyte stirring to obtain compact and uniform coatings suitable for application in industrial purposes. The most often used methods of an electrolyte mixing are magnetic stirring (MS), ultrasound assistance (US) and barbotage [12,17–19]. A stirring of electrolyte during electrodeposition process achieves a strong influence on mass transfer, nucleation and growth, and as a result of it, on morphological, structural and mechanical features of electrodeposited coatings [19]. As already mentioned, magnetic stirring (MS) was applied for obtaining of Cu coatings given in previous Sections.

Ultrasound (US) represents a significant and widely used way of an electrolyte stirring [17,19–22]. This way enables an application of different intensities of electrolyte mixing through a regulation of value of power (P_i) relative to maximal power of ultrasonic device [19].

Figure 6 shows the morphologies of the Cu coatings obtained from *electrolyte I* with an intensity of US which corresponds to P_i of 10% (i.e., 10% of maximal power of ultrasonic device of 200 W [19]), Figure 6a and P_i of 50%, Figure 6b.

The both Cu coatings are fine-grained, with a clear difference between grains and similar to each other at the macro level.

Since the AFM analysis includes the Cu coating obtained with P_i of 30%, Figure 7 shows the 3D images and the histograms of topography for the Cu coatings electrodeposited with intensities of US of 10, 30 and 50% ($P_i = 10\%$, Figure 7a,b; $P_i = 30\%$, Figure 7c,d; $P_i = 50\%$, Figure 7e,f).

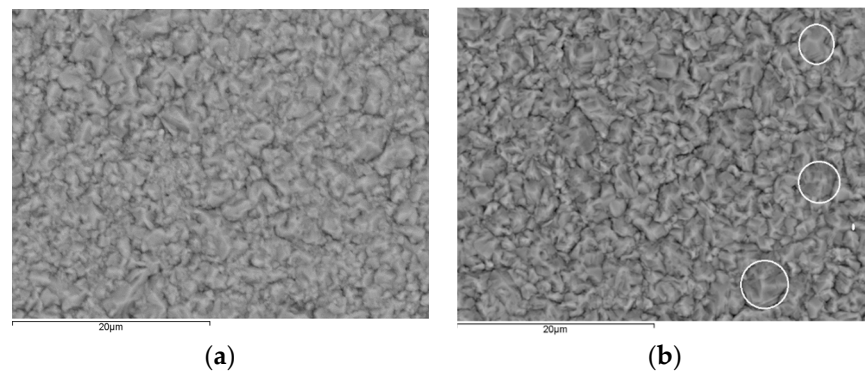


Figure 6. The SEM micrographs of the morphology of Cu coatings obtained by electrodeposition at the low temperature ($T = 7.0 \pm 1.0$ °C) at a current density of 50 mA cm^{-2} on the brass cathodes from *electrolyte I* with an intensity of ultrasound corresponding to: (a) $P_i = 10\%$, and (b) $P_i = 50\%$. The coating thickness: 16.6 μm [19].

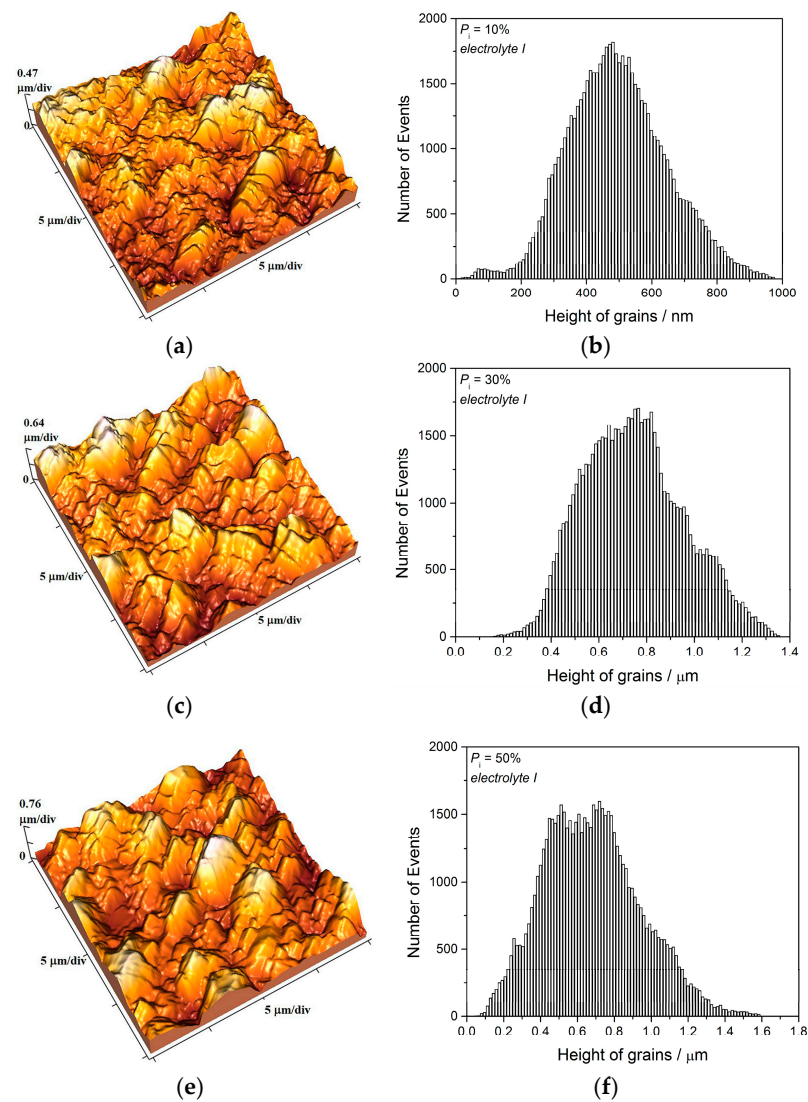


Figure 7. The 3D AFM images and the corresponding histograms of Cu coatings electrodeposited at the low temperature ($T = 7.0 \pm 1.0$ °C) at a current density of 50 mA cm^{-2} on the brass cathodes from *electrolyte I* with an intensity of ultrasound corresponding to: (a,b) $P_i = 10\%$, (c,d) $P_i = 30\%$, and (e,f) $P_i = 50\%$. The coating thickness: 16.6 μm . Scan size: $(20 \times 20) \text{ μm}^2$ [19].

The R_a roughness values of electrodeposited Cu coatings were 138.1 ± 4.31 nm for the coating obtained with P_i of 10%, 181.2 ± 7.34 nm for that obtained with P_i of 30% and 212.4 ± 8.2 nm for the coating obtained with P_i of 50% [19]. The increasing trend of the roughness value was noted with the increase of the intensity of US.

The morphologies of Cu coatings electrodeposited under the same electrolysis conditions as those shown in Figure 6, but from *electrolyte II* are presented in Figure 8.

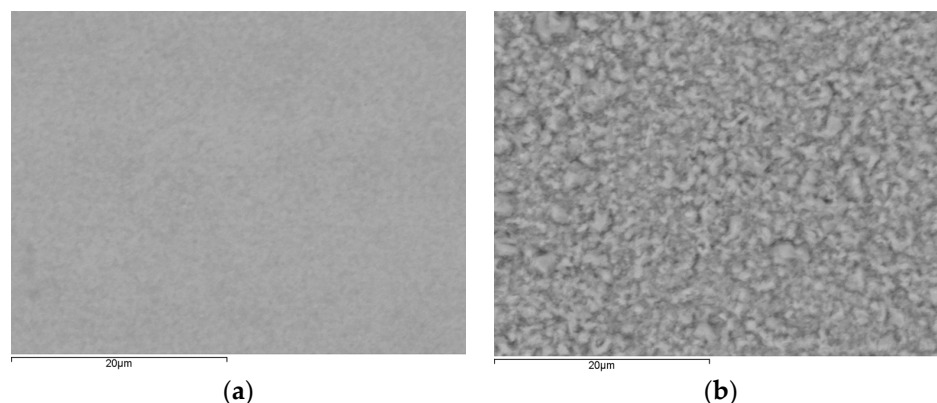


Figure 8. The SEM micrographs of the morphology of Cu coatings electrodeposited at the low temperature ($T = 7.0 \pm 1.0$ °C) at a current density of 50 mA cm^{-2} on the brass cathodes from *electrolyte II* with an intensity of ultrasound corresponding to: (a) $P_i = 10\%$, and (b) $P_i = 50\%$. The coating thickness: $16.6 \mu\text{m}$ [19].

The intensity of applied ultrasound had a strong influence on morphology of the electrodeposited Cu coatings. The Cu coating electrodeposited with P_i of 10% was smooth, without visible boundary among grains. The coating of Cu produced with P_i of 50% was fine-grained, and boundary among grains can be seen.

The 3D AFM images and the histograms of topography of the coatings with included those obtained with P_i of 30% are shown in Figure 9 ($P_i = 10\%$, Figure 9a,b; $P_i = 30\%$, Figure 9c,d; $P_i = 50\%$, Figure 9e,f).

The R_a roughness values of the coatings electrodeposited from *electrolyte II* were 26.87 ± 2.1 nm for the coating obtained with P_i of 10%, 53.07 ± 3.25 nm for that obtained with P_i of 30% and 90.5 ± 3.4 nm for the coating obtained with P_i of 50% [19]. Similar to the coatings obtained from *electrolyte I*, the coating roughness increased with the increase of the intensity of ultrasound.

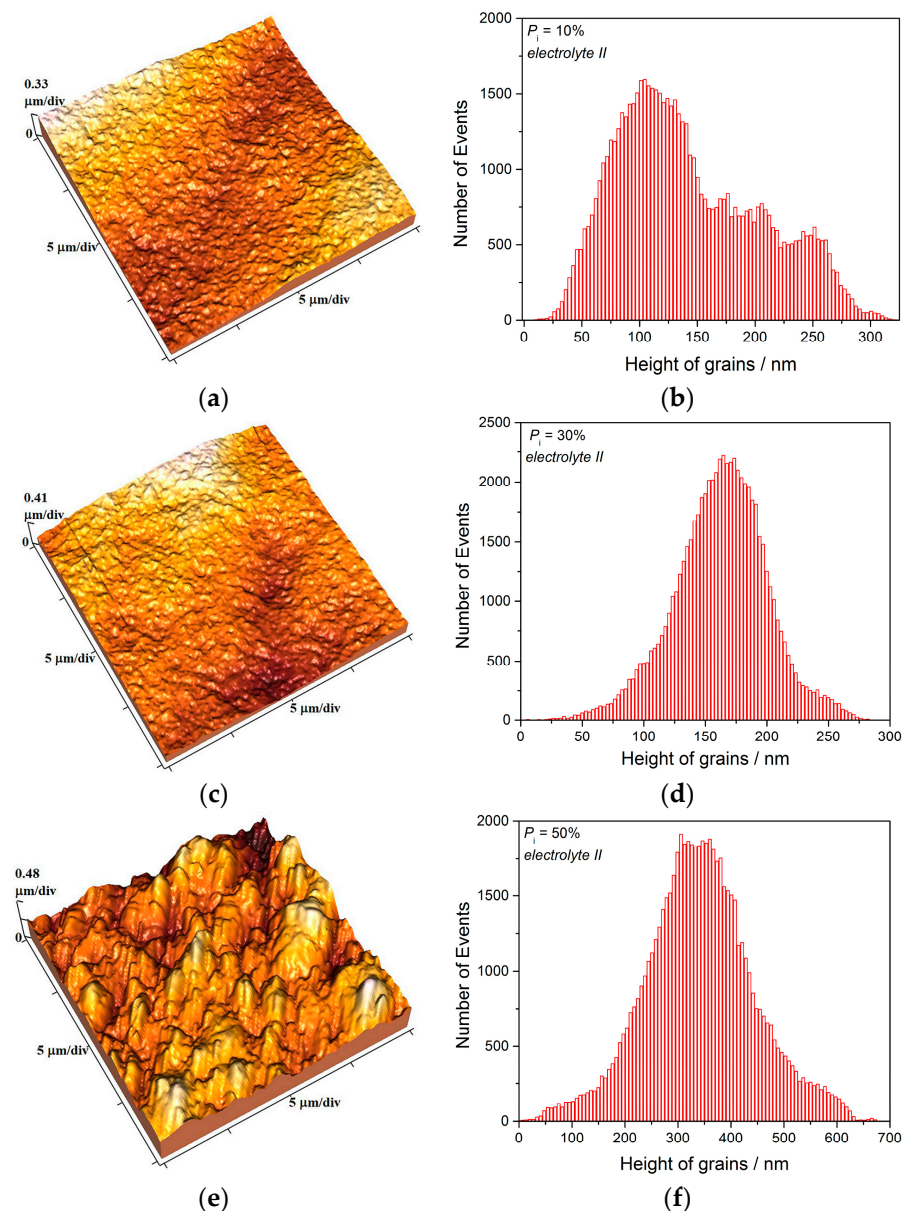


Figure 9. The 3D AFM images and the corresponding histograms of Cu coatings electrodeposited at the low temperature ($T = 7.0 \pm 1.0$ °C) at a current density of 50 mA cm^{-2} on the brass cathodes from *electrolyte II* with an intensity of ultrasound corresponding to: (a,b) $P_i = 10\%$, (c,d) $P_i = 30\%$, and (e,f) $P_i = 50\%$. The coating thickness: $16.6 \text{ } \mu\text{m}$. Scan size: $(20 \times 20) \text{ } \mu\text{m}^2$ [19].

3. Influence of Parameters of the Electrodeposition on Hardness of Cu Coatings Produced by the DC Regime

Hardness is the most important mechanical characteristic of metal coatings, and microindentation technique represents one of often used ways to determine it [13]. The measured hardness usually consists of a contribution both substrate (cathode) and coating, and for that reason, it is called composite hardness [12,13,23,24]. The largest challenge in a determination of an absolute (a true, an intrinsic) coating hardness is to eliminate a contribution of substrate to measured composite hardness. It can be achieved on two ways: (a) by use of low indentation loads, and (b) applying the composite hardness models. The main lack of the first way is a huge error of measurement of diagonal size for thin coatings caused by use of low indentation loads, and for that reason, this way is only suitable for thick coatings [12]. On the other hand, composite hardness models are mathematic models including large number of approximations, and their application requires a knowledge of

many phenomena, such as applied load, duration of the applied load at an indentation (dwell time) and indentation depth [25], elastic properties of the coatings [26], indentation size effect (ISE) [27–30], interfacial adhesion [31,32], etc.

A lot of mathematical models are developed which can be used for a determination of the absolute hardness of metal coatings. Some of them are Chicot–Lesage (C–L) [23,24,33,34], Korsunsky (K) [35–37], Chen–Gao (C–G) [38–41], Burnett–Rickerby (B–R) [31,32], Jonson–Hogmark (J–H) [42], Puchi–Cabrera (P–C) [43,44] models, etc. There is no an universal model applicable for all substrate/coating systems, and some of them are adapted to “soft film (coating) on hard substrate” system, such as C–L and C–G models, while the other models, such as K, J–H and P–C models, are adapted to “hard film (coating) on soft substrate” system.

In this Section, Vickers microindentation technique was used to determine an absolute hardness of the Cu coatings electrodeposited on Si(111) and brass substrates under various electrodeposition conditions. The indentation loads between 0.049 and 2.942 N, and the dwell time of 25 s were used [12,13,45]. The Cu coatings electrodeposited on Si(111) and brass substrates belong to “soft film (coating) on hard substrate” system, and the Chicot–Lesage (C–L) composite hardness model was applied for a determination of absolute hardness of the coatings. This model was selected because the real values of coating hardness can be determined by a simple measurement of diagonal size of Vickers imprint in a surface area of the coating. This model does not require use complex software packages and is based on solution of a quadratic equation for every applied load, i.e., point by point, whereby the positive value of a quadratic equation solution is taken as a value for the coating hardness. The basis of the C–L model is given in Ref. [13].

The composite hardness, H_c usually includes a contribution of both substrate (cathode) and coating, and it is presented by Equation (1) [12,13,45]:

$$H_c = \frac{1.8544 \cdot P}{d^2} \quad (1)$$

In Equation (1), P is applied load, in N, and d , in m is size of diagonal made in a coating during an indentation process.

The typical imprint in the Cu coating obtained by Vickers microindentation is shown in Figure 10.

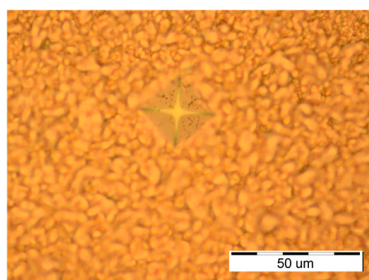


Figure 10. The imprint obtained by Vickers microindentation in the electrolytically produced Cu coating. The applied load: 0.49 N. The dwell time: 15 s [46].

Vickers microindentation belongs to the static method for a determination of hardness of coatings (films). Aside from this static method, there are also the dynamic methods which involve both micro- and nanoindentation [47,48]. Unlike the static method where a determination of coating hardness is based on a measurement of diagonal size of imprint in a coating made by applied load, the dynamic methods are based on continuous processes where the applied load and the displacement of indenter are simultaneously recorded. The nanoindentation method is especially suitable to determine a hardness of thin coatings owing to an avoidance of an influence of substrate with the small applied loads [49].

In order to determine an absolute hardness of any coating, knowledge of hardness of cathode is of crucial importance. The PSR (Proportional Specimen Resistance) model [50]

is often used for that purpose, and the values of substrate hardness, H_s were 7.42 GPa for Si(111) [13] and 1.41 GPa for the brass [45].

The following Sub-sections give a survey of the effects of types of both electrolyte and cathode, the thickness of the coating and the electrolyte stirring on hardness of the Cu coatings.

3.1. The Influence of Type of Electrolyte on Hardness of Electrolytically Produced Cu Coatings

The dependencies of the composite hardness, H_c for 20 μm thick Cu coatings produced from *electrolyte I* and *electrolyte II* on the brass cathodes on relative indentation depth (RID) are presented in Figure 11a. The RID is given by Equation (2) [12,13,45,51]:

$$\text{RID} = \frac{h}{\delta} = \frac{d}{7\delta} \quad (2)$$

where h is an indentation depth, δ is a coating thickness, and d is a diagonal size, since $h = d/7$. The effect of cathode hardness in the composite hardness grows with an increase of RID value, i.e., with an increase of a depth of indentation. Although there is no sharp boundary where a substrate commences to effect on a coating hardness, as well as where a composite hardness is being equaled with hardness of a substrate, RID values between 0.1 and 1 are usually considered as boundaries for these influences. The RID zone between 0.1 and 1 is so-called composite zone, and it is a zone where there is strong contribution of cathode hardness to composite hardness. From this reason, for H_c values dominantly situated in this zone, it is necessary to apply the corresponding composite hardness model to determine an absolute hardness of any coating.

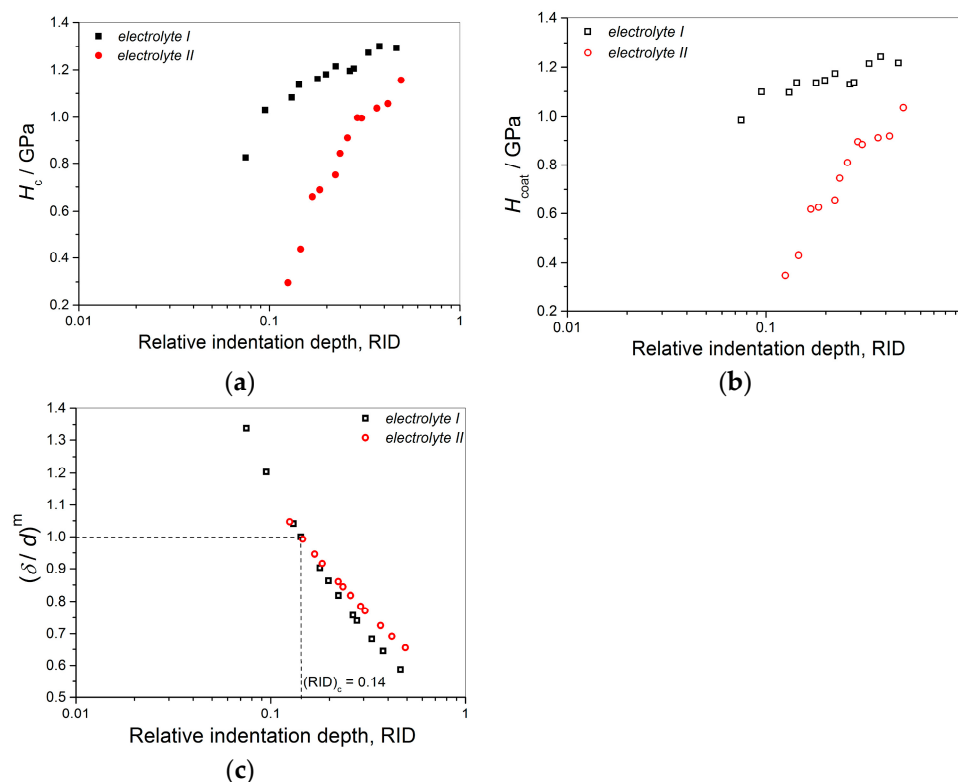


Figure 11. The dependencies of: (a) H_c , (b) H_{coat} , and (c) parameter $(\delta/d)^m$ on RID for the 20 μm thick Cu coatings produced at a current density of 50 mA cm^{-2} on the brass from *electrolyte I* and *electrolyte II* [12].

The dependencies of H_{coat} on RID calculated by C–L model are presented in Figure 11b. It follows that mat coating electrodeposited from *electrolyte I* had larger both H_c and H_{coat}

values than mirror bright coating electrodeposited from *electrolyte II*. The values both measured H_c and calculated H_{coat} showed same trend of the dependencies on the RID.

However, it is noticed that for RID values close to 0.1 calculated values of H_{coat} were larger than H_c values, whereby the difference between them increased with a decrease of RID value. This illogicity demands an additional analysis in an implementation of the C–L model to obtain real values of hardness of the Cu coatings. It is shown in Figure 11c the dependencies of the $(\delta/d)^m$ on the RID for the coatings produced from *electrolyte I* and *electrolyte II*, where m is a composite Meyer's index calculated by a linear regression on the way described in Refs. [24,33]. The values of this coefficient are 0.4525 for mat coating electrodeposited from *electrolyte I* and 0.3423 for mirror bright coating electrodeposited from *electrolyte II* [12].

The C–L model is valid up to $(\delta/d)^m = 1$ [23,24,33,34], and taking this value, it is obtained RID value of 0.14 (Figure 11c). For $RID < 0.14$, H_c values are smaller than calculated H_{coat} values, and the coating hardness is equal to the measured composite hardness. For $RID \geq 0.14$, H_c values are larger than H_{coat} values, and the coating hardness corresponds to the calculated value. Hence, the RID value of 0.14 is the limiting (or critical) value $[(RID)_c = 0.14]$ separating a zone where the measured (composite) hardness is equal to the coating hardness and a zone in which calculated values correspond to the coating hardness.

3.2. The Effect of Type of Cathode on Hardness of Electrolytically Produced Cu Coatings

The dependencies of H_c , H_{coat} and $(\delta/d)^m$ on RID obtained for 20 μm thick Cu coatings produced from *electrolyte I* and *electrolyte II* on Si(111) substrate are shown in Figure 12. The values of composite Meyer's index for these coatings are $m = 0.4153$ for the coating electrodeposited from *electrolyte I* and $m = 0.3447$ for the coating electrodeposited from *electrolyte II* [12].

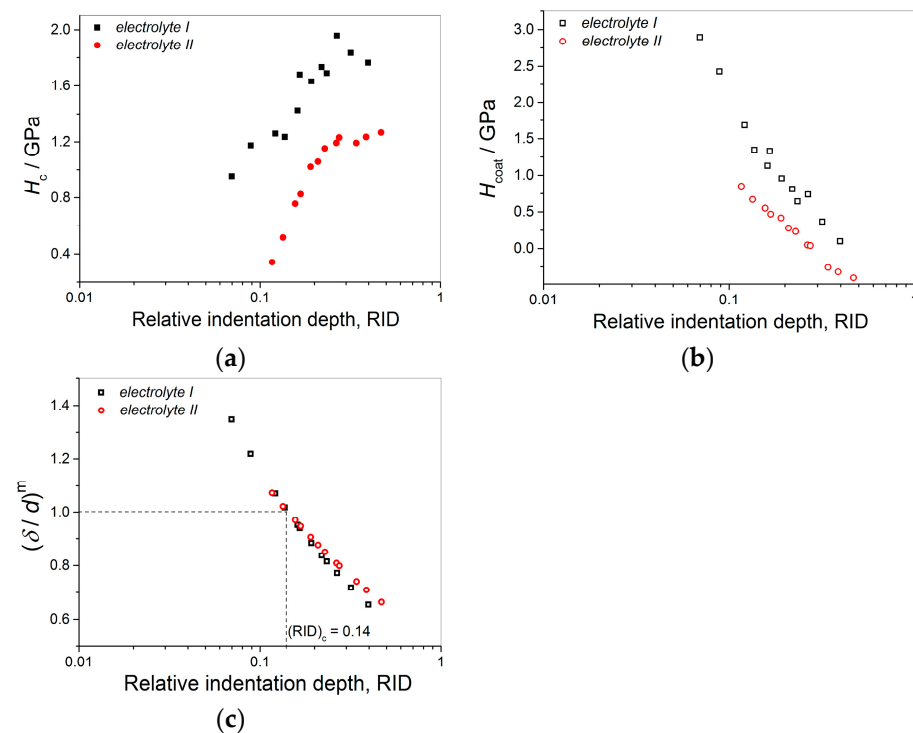


Figure 12. The dependencies of: (a) H_c , (b) H_{coat} , and (c) parameter $(\delta/d)^m$ on RID for the 20 μm thick Cu coatings produced at a current density of 50 mA cm^{−2} on the Si(111) substrate from *electrolyte I* and *electrolyte II* [12].

It is clear from Figure 12b that the dependencies of the calculated H_{coat} on the RID values have opposite trend than those obtained on brass substrate. As seen from Figure 12b,

the dependencies of H_{coat} –RID have a descending trend. For this system, H_{coat} values are considerably larger than H_c values up to RID of 0.14, with a tendency of a decrease of this difference with an increase of RID value. After RID value larger than 0.14, the H_{coat} values become smaller than H_c values, confirming that this value is the limiting or critical one after which an application of the C–L model was necessary for a determination of an absolute hardness of the Cu coatings.

3.3. The Effect of the Coating Thickness on Hardness of Electrolytically Produced Cu Coatings

Figure 13 shows the dependencies of H_c , H_{coat} and $(\delta/d)^m$ on RID for 40 μm thick Cu coatings obtained by the electrodeposition from *electrolyte I* and *electrolyte II* on Si(111) substrate. The values of Meyer's parameters are $m = 0.4290$ for the Cu coating electrodeposited from *electrolyte I* and $m = 0.3367$ for the Cu coating electrodeposited from *electrolyte II* [12].

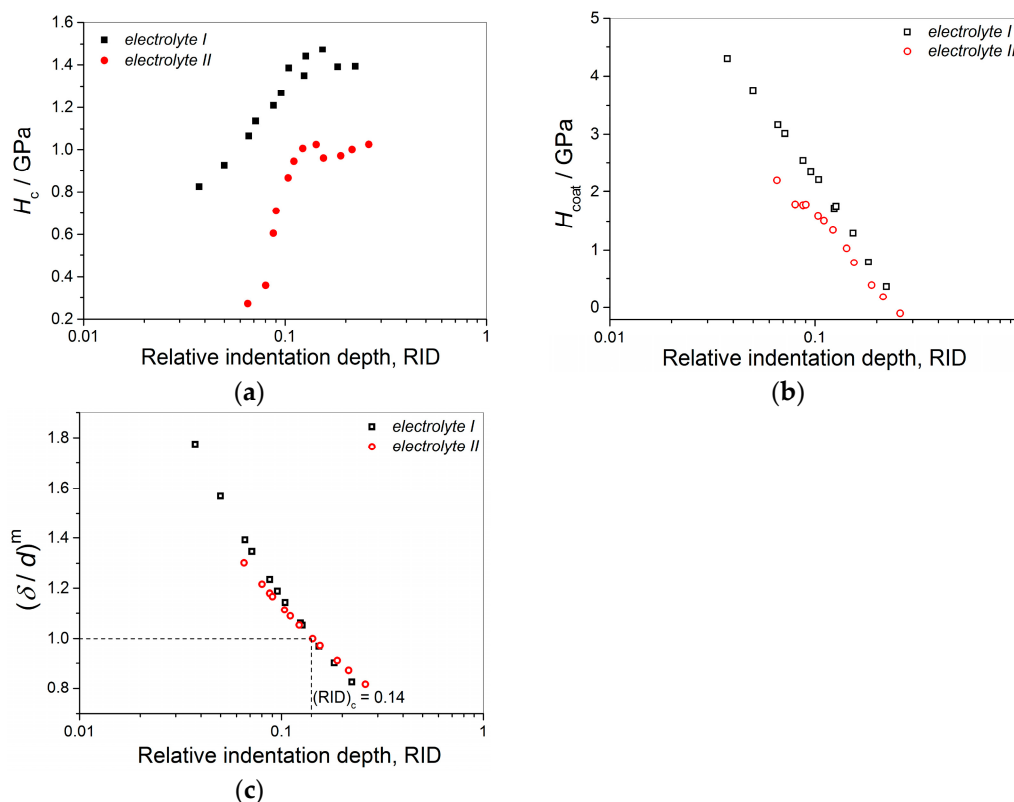


Figure 13. The dependencies of: (a) H_c , (b) H_{coat} , and (c) parameter $(\delta/d)^m$ on RID for the 40 μm thick Cu coatings electrodeposited at a current density of 50 mA cm^{-2} on the Si(111) substrate from *electrolyte I* and *electrolyte II* [12].

The dependencies of H_c and H_{coat} on RID values have the same trend as those obtained for 20 μm thick Cu coatings on the same substrate (Figure 12). The only difference is a shift of hardness values towards lower RID values, i.e., towards the zone of equalized composite (measured) and coating hardness. For this case, the RID value of 0.14 was confirmed as the limiting value separating a zone where the coating hardness is equal to the composite (measured) hardness than a zone where it is necessary to apply the C–L model to determine an absolute hardness of the Cu coatings.

3.4. Influence of Electrolyte Stirring on Hardness of Electrolytically Produced Cu Coatings

Figure 14 shows the dependencies of H_c , H_{coat} and $(\delta/d)^m$ on RID obtained by the electrodeposition from *electrolyte I* on the brass substrate with various intensities of imposed ultrasound. The values of coefficient m for the considered coatings are given in Table 2.

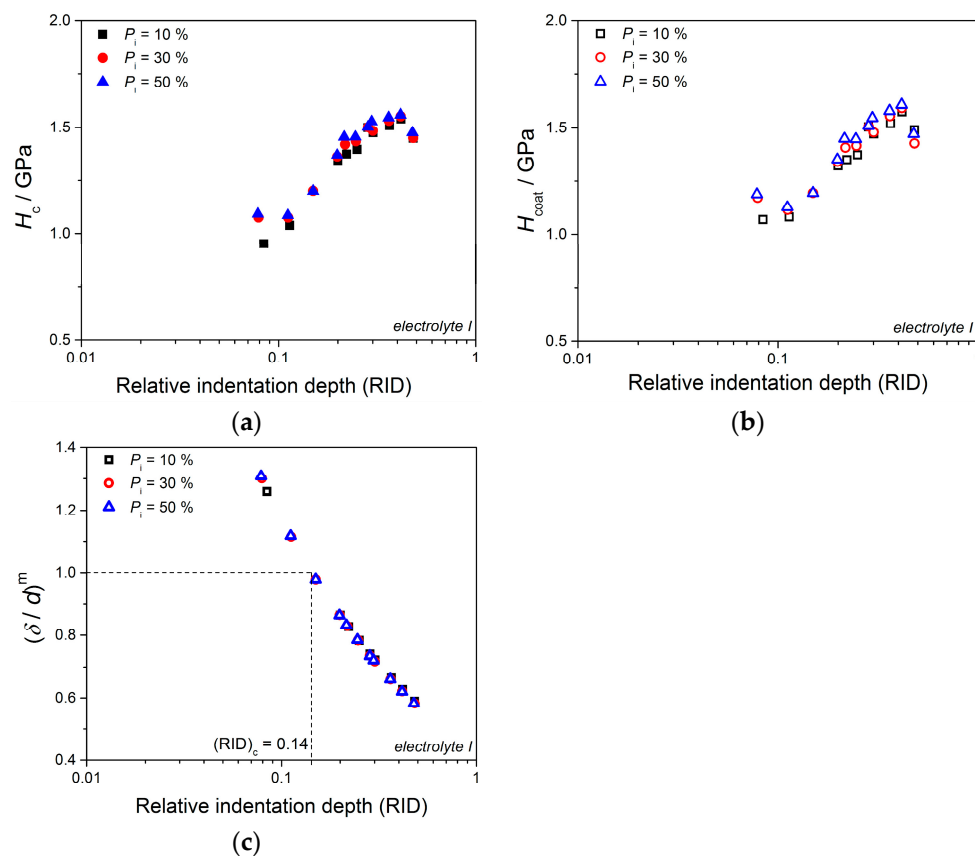


Figure 14. The dependencies of: (a) H_c , (b) H_{coat} , and (c) parameter $(\delta/d)^m$ on RID for the Cu coatings produced from *electrolyte I* on the brass with various intensities of imposed ultrasound [19].

Table 2. The values of Meyer's index (m) for the Cu coatings electrodeposited with different intensities of ultrasound [19].

$P_i/\%$	10	30	50
m (<i>electrolyte I</i>)	0.4503	0.4594	0.4750
m (<i>electrolyte II</i>)	0.4407	0.4565	0.4725

The dependencies of values of H_c and H_{coat} on RID have „S“ shape, as seen from Figure 14a,b. Both of the hardness values grow with an increase of RID values up to about 0.40. Then, their sharp decrease commences up to the value corresponding approximately to a hardness of the brass cathode, H_s , of 1.41 GPa [45]. For these coatings, the limiting or critical RID value of 0.14 (Figure 14c) separating the two zones is also confirmed.

The dependencies of H_c , H_{coat} and $(\delta/d)^m$ on RID obtained for the Cu coatings produced from *electrolyte II* on the brass with various intensities of imposed ultrasound are shown in Figure 15. The values of m coefficients for the considered coatings are given in Table 2.

The values of both H_c and H_{coat} increased with the increase of RID value, approaching to value corresponding to hardness of the brass cathode (Figure 15a,b). Certainly, the „S“ shape of these dependencies is not obtained, probably due to lower values of both H_c and H_{coat} relative to the coatings electrodeposited from *electrolyte I*. The limiting RID value of 0.14 was also observed for the coatings electrodeposited in the presence of leveling/brightening addition agents under imposed ultrasound of various intensities.

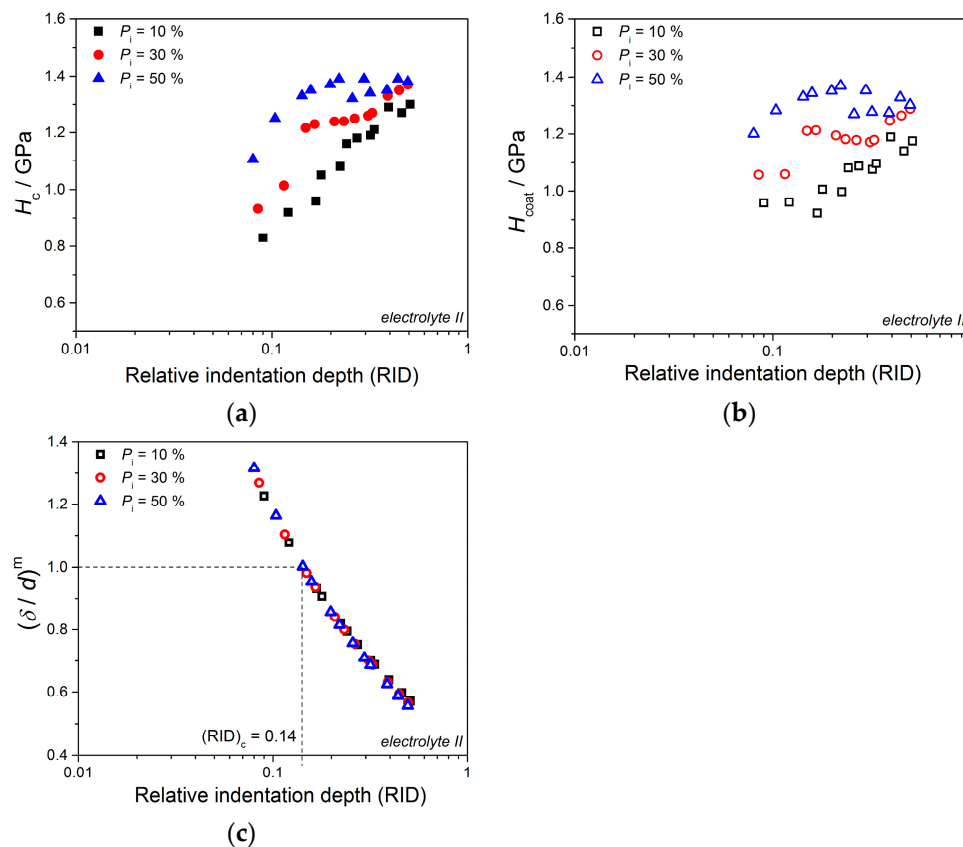


Figure 15. The dependencies of: (a) H_c , (b) H_{coat} , and (c) parameter $(\delta/d)^m$ on RID for the Cu coatings produced from *electrolyte II* on the brass with various intensities of imposed ultrasound [19].

4. Influence of Parameters of the PC Regime on Morphology and Structure of Electrolytically Produced Cu Coatings

Electrodeposition by pulse reverse current (PRC) regimes offers many benefits in obtaining metal deposits of desired characteristics. The deposits obtained by these regimes usually possess both lower porosity and more compact structure than the deposits produced by the constant regimes of electrodeposition. The development of electronic industry lowers a price of pulse rectifiers, that is still incomparable higher than the DC units [52], and this lowering of price enables larger application of pulse reverse regimes in commercial purposes. The only pulsating (PC) and reversing current (RC) regimes have potential commercial application from all PRC regimes of the electrodeposition.

The PC regime is defined by the average current density (j_{av}), the current density amplitude (j_A), a deposition pulse (t_c) and a period of pause (t_p) on way given by Equation (3) [8]:

$$j_{\text{av}} = \frac{j_A \cdot t_c}{t_c + t_p} \quad (3)$$

Since the pause-to-pulse ratio, p is defined by Equation (4) [8]:

$$p = \frac{t_p}{t_c} \quad (4)$$

The average current density (j_{av}) is also defined by Equation (5):

$$j_{\text{av}} = \frac{j_A}{1 + p} \quad (5)$$

The frequency, ν is defined by Equation (6):

$$\nu = \frac{1}{t_c + t_p} \quad (6)$$

The pulsating current regime gives optimal results in the millisecond range, in the (10–100) Hz frequency range [8]. In this range of frequency, the surface concentration of depositing ions is constant and corresponds to a concentration in the constant DC regime at the current density equal to the average current density in regime of the pulsating current [8,53]. It means that electrochemical deposition in the PC regime in this frequency range takes place at the average current density.

The regime of reversing current (RC) includes anodic component of the current density, and instead of pause duration, the anodic time is included [8].

In this review paper, influence of the following parameters of electrodeposition and the pulsating current regime on morphology, structure and hardness of the Cu coatings is analyzed [13,45,46]:

(1) the average current density, j_{av} or frequency, ν —these parameters are analyzed keeping constant the current density amplitude and the deposition pulse, while pause duration is varied,

(2) the current density amplitude, j_A —these parameters are analyzed keeping constant both deposition pulse and pause duration,

(3) coating thickness, δ —coating thickness is varied keeping the parameters of pulsating current regime constant (the current density amplitude, the deposition pulse, and the pause duration), and

(4) type of cathode.

The parameters of the pulsating current regime analyzed in this review paper are summarized in Table 3.

Table 3. The parameters of the pulsating current (PC) regime used in the electrodeposition processes. j_A —the current density amplitude, t_c —deposition time (on period), t_p —pause time (off period), p —the pause-to-pulse ratio, j_{av} —the average current density and ν —frequency [13].

Number of Samples	$j_A/\text{mA cm}^{-2}$	t_c/ms	t_p/ms	$\delta/\mu\text{m}$	$j_{av}/\text{mA cm}^{-2}$	ν/Hz
1	100	5	5	40	50	100
2	100	5	7.5	40	40	80
3	100	5	15	40	25	50
4	100	5	28.3	40	15	30
5	120	5	5	40	60	100
6	140	5	5	40	70	100
7	100	5	5	20	50	100

4.1. Influence of the Average Current Density (or Frequency) on Morphology and Structure of the Cu Coatings

Figure 16 shows the Cu coatings produced at j_{av} of 15 (Figure 16a,b), 25 (Figure 16c,d), 40 (Figure 16e,f) and 50 mA cm^{-2} (Figure 16g,h) on the Si(111) electrode. All Cu coatings presented in Section 4 were electrodeposited from *electrolyte I* at a temperature of 22.0 ± 0.50 °C without electrolyte stirring.

The size of grains decreased with an increase of the average current density (or frequency) from 15 to 50 mA cm^{-2} (i.e., from 30 to 100 Hz). The well-defined grains size about 10 μm are mostly formed at 15 mA cm^{-2} (30 Hz). The number of grains of this size decreased with increasing the average current density, in order to size of grains in the coatings produced at 40 and 50 mA cm^{-2} (80 and 100 Hz) was predominant about or below 5 μm [13].

The topography and the histograms of the shown Cu coatings are presented in Figure 17a,c,e,g—AFM images; Figure 17b,d,f,h—histograms.

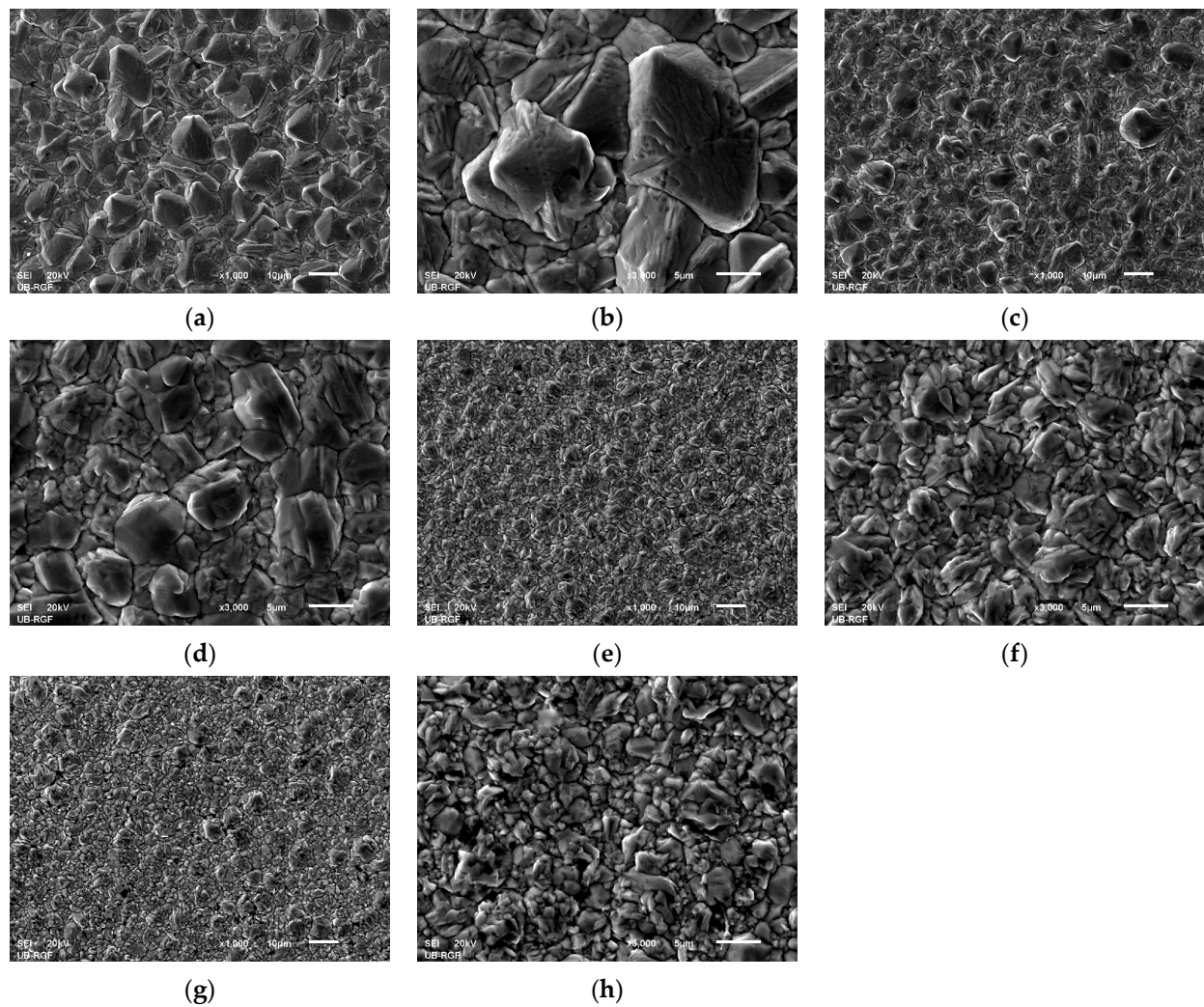


Figure 16. The SEM micrographs of morphology of Cu coatings electrodeposited by the PC regime at j_{av} of: (a,b) 15, (c,d) 25, (e,f) 40 and (g,h) 50 mA cm^{-2} . The current density amplitude: $j_A = 100 \text{ mA cm}^{-2}$; The cathode: Si(111); the coating thickness, $\delta = 40 \mu\text{m}$; Magnification: (a,c,e,g) $\times 1000$, and (b,d,f,h) $\times 3000$ [13].

The R_a roughness values for the coatings are given in Table 4, from which can be seen that the coating roughness decreases with the increase of the average current density.

Table 4. The R_a values of roughness obtained for the Cu coatings produced at different average current densities (or frequencies) [13].

$j_{av}/\text{mA cm}^{-2}$	15	25	40	50
R_a / nm	507.3 ± 3.2	470.5 ± 2.9	385 ± 2.4	169.9 ± 2.1

The XRD analysis of 40 μm thick Cu coatings produced by various PC regimes is shown in Figure 18. The shown diffractograms represent typical diffractograms of Cu with 2θ angles of 43.3° , 50.4° , 74.1° and 89.9° corresponding to the (111), (200), (220) and (311) crystal planes.

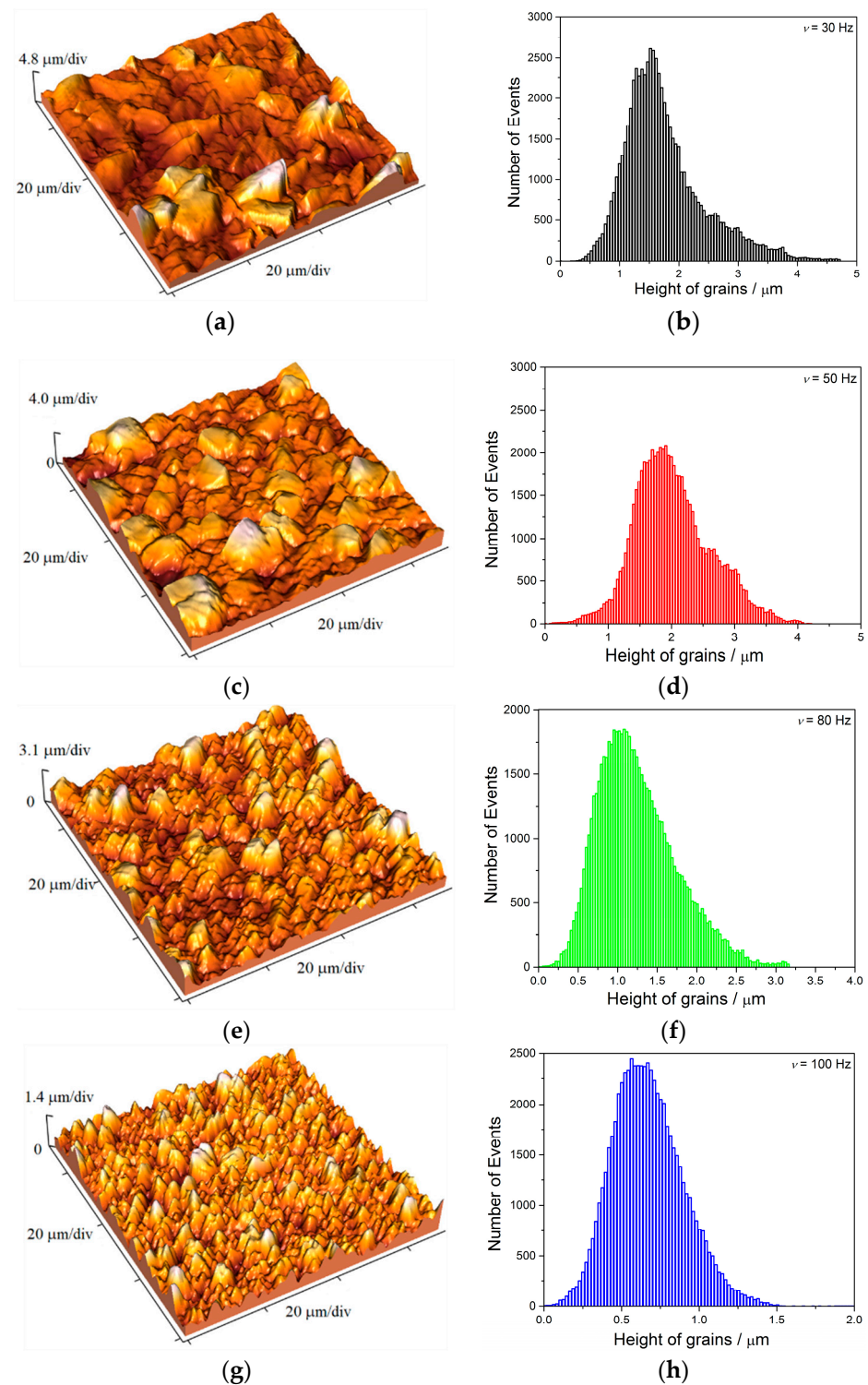


Figure 17. The 3D AFM images and the histograms of Cu coatings electrodeposited by the PC regime: (a,b) $j_{av} = 15 \text{ mA cm}^{-2}$; $\nu = 30 \text{ Hz}$, (c,d) $j_{av} = 25 \text{ mA cm}^{-2}$; $\nu = 50 \text{ Hz}$, (e,f) $j_{av} = 40 \text{ mA cm}^{-2}$; $\nu = 80 \text{ Hz}$, and (g,h) $j_{av} = 50 \text{ mA cm}^{-2}$; $\nu = 100 \text{ Hz}$. The cathode: Si(111); the coating thickness, $\delta = 40 \text{ }\mu\text{m}$. Scan size: $(70 \times 70) \text{ }\mu\text{m}^2$ [13].

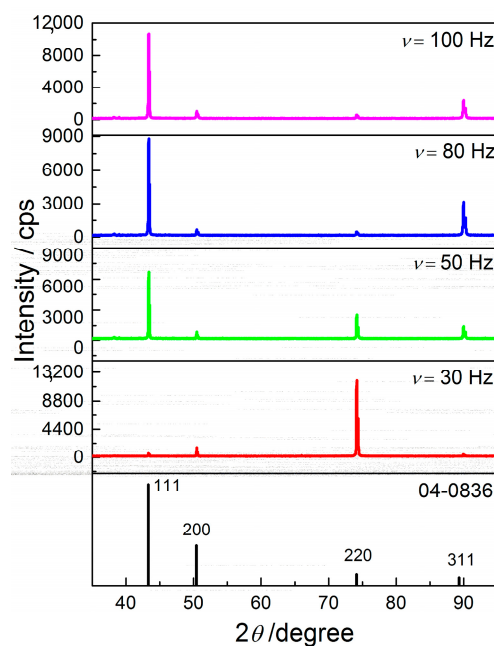


Figure 18. The XRD patterns recorded for the Cu coatings produced by the PC regime at various frequencies (or the average current densities), and Cu standard (04-0836) [13].

The values of $TC(hkl)$ and $RTC(hkl)$ coefficients for given the Cu coatings are summarized in Tables 5 and 6.

Table 5. TC and RTC coefficients calculated for Cu coatings produced by the PC regimes at frequencies of 30 and 50 Hz; s—Cu standard; R is an intensity of diffraction peak [13].

Plane (hkl)	R (in%)		R_s (in%)	TC		RTC (in%)	
	$R_{30\text{ Hz}}$	$R_{50\text{ Hz}}$		$TC_{30\text{ Hz}}$	$TC_{50\text{ Hz}}$	$RTC_{30\text{ Hz}}$	$RTC_{50\text{ Hz}}$
(111)	4.0	58.6	54.6	0.073	1.07	0.88	22.8
(200)	11.2	6.9	25.1	0.45	0.27	5.4	5.8
(220)	81.9	22.2	10.9	7.5	2.04	90	43.5
(311)	2.9	12.3	9.4	0.31	1.31	3.72	27.9

Table 6. TC and RTC coefficients calculated for Cu coatings produced by the PC regimes at frequencies of 80 and 100 Hz; s—Cu standard; R is an intensity of diffraction peak [13].

Plane (hkl)	R (in%)		R_s (in%)	TC		RTC (in%)	
	$R_{80\text{ Hz}}$	$R_{100\text{ Hz}}$		$TC_{80\text{ Hz}}$	$TC_{100\text{ Hz}}$	$RTC_{80\text{ Hz}}$	$RTC_{100\text{ Hz}}$
(111)	67.3	72.7	54.6	1.23	1.33	28.2	35.9
(200)	5.0	7.1	25.1	0.20	0.28	4.6	7.6
(220)	3.6	4.0	10.9	0.33	0.37	7.6	10
(311)	24.1	16.2	9.4	2.6	1.72	59.6	46.5

The Cu coating produced at j_{av} of 15 mA cm^{-2} (or ν of 30 Hz) exhibited the strong (220) preferred orientation. With an increase of j_{av} (or ν), the share of crystallites oriented in this plane decreased, and crystallites of Cu become predominately oriented in (311), (111) crystal planes [13].

4.2. Influence of the Current Density Amplitude on Morphology and Structure of the Cu Coatings

Figure 19 shows the Cu coatings produced with the current density amplitudes, j_A of 120 (Figure 19a,b) and 140 mA cm^{-2} (Figure 19c,d). These current density ampli-

tudes corresponded to j_{av} of 60 and 70 mA cm^{-2} , since values t_c and t_p are kept constant ($t_c = t_p = 5$ ms).

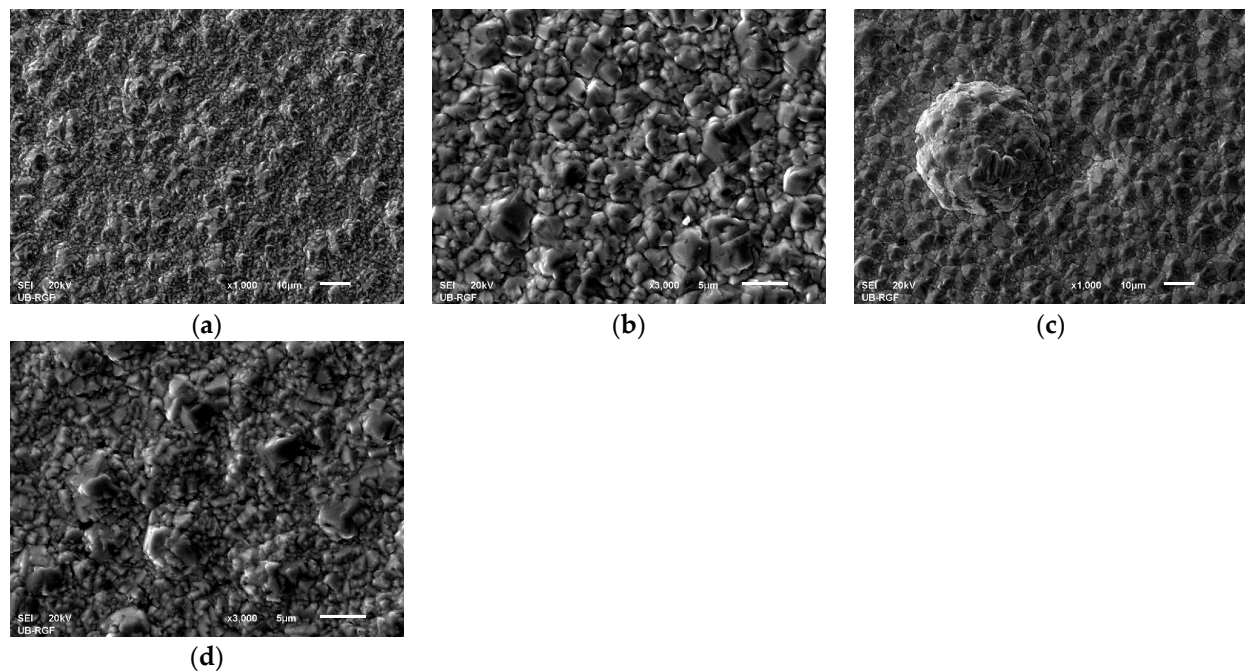


Figure 19. The SEM micrographs of morphology of Cu coatings electrodeposited by the PC regime with j_A of: (a,b) 120, and (c,d) 140 mA cm^{-2} . The cathode: Si(111); the coating thickness, $\delta = 40$ μm ; Magnification: (a,c) $\times 1000$, and (b,d) $\times 3000$ [13].

The size of grains in the Cu coating electrodeposited with j_A of 120 mA cm^{-2} ($j_{av} = 60$ mA cm^{-2}) was about 5 μm or less, and this coating was very similar to those produced with j_{av} of 40 and 50 mA cm^{-2} (Figure 16e–h). The further increase in j_A (or j_{av}) led to formation of globules size of about 30 μm (Figure 19c).

The 3D AFM images and the histograms of the topography of the Cu coatings obtained with the current density amplitudes of 120 and 140 mA cm^{-2} are shown in Figure 20 (Figure 20a,b for the coating obtained with j_A of 120 mA cm^{-2} and Figure 20c,d with j_A of 140 mA cm^{-2}).

The R_a roughness values of these coatings were 237 ± 2.2 nm for the Cu coating produced with j_A of 120 mA cm^{-2} , and 229.1 ± 2.2 nm for that produced with j_A of 140 mA cm^{-2} . The obtained values were slightly higher than that obtained for coating produced with j_A of 100 mA cm^{-2} under the same all other parameters of the PC regime (Figure 16g,h). It is necessary to note that the deposit among globules was analyzed in the coating electrodeposited with j_A of 140 mA cm^{-2} .

The XRD analysis (Figure 21) revealed a strong influence of the current density amplitude on the preferred orientation of the produced Cu coatings. Please note that the diffractogram shown in Figure 18 for ν of 100 Hz ($j_A = 100$ mA cm^{-2}) is repeated in Figure 21.

Analysis of $TC(hkl)$ and $RTC(hkl)$ coefficients given in Table 7 showed that Cu crystallites were mostly oriented in (111) crystal plane, and that a share of crystallites oriented in this plane increased with an increase of the current density amplitude. Generally, the increase of j_{av} from 15 to 70 mA cm^{-2} (Figures 16 and 19, Tables 5–7) causes a change of the preferred orientation of the Cu coatings from the strong (220) to the strong (111) preferred orientation.

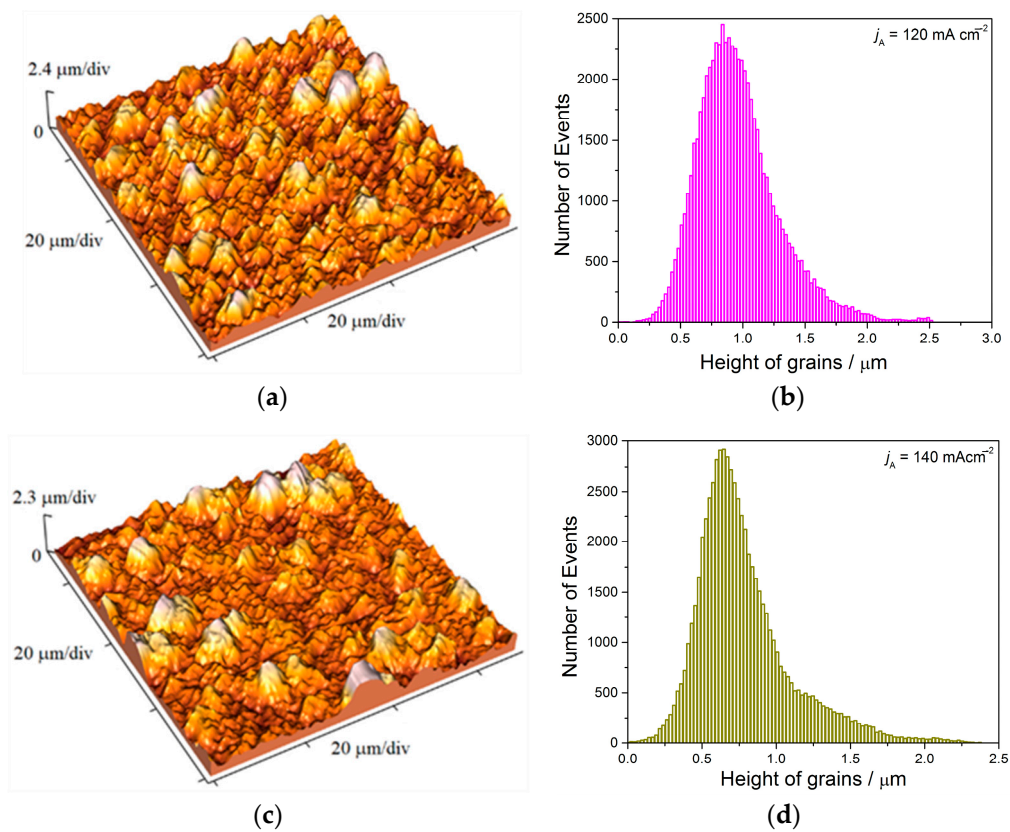


Figure 20. The 3D AFM images and the histograms of Cu coatings electrodeposited by the PC regime: (a,b) $j_A = 120 \text{ mA cm}^{-2}$, and (c,d) $j_A = 140 \text{ mA cm}^{-2}$. The cathode: Si(111); the coating thickness, $\delta = 40 \text{ }\mu\text{m}$. Scan size: $(70 \times 70) \text{ }\mu\text{m}^2$ [13].

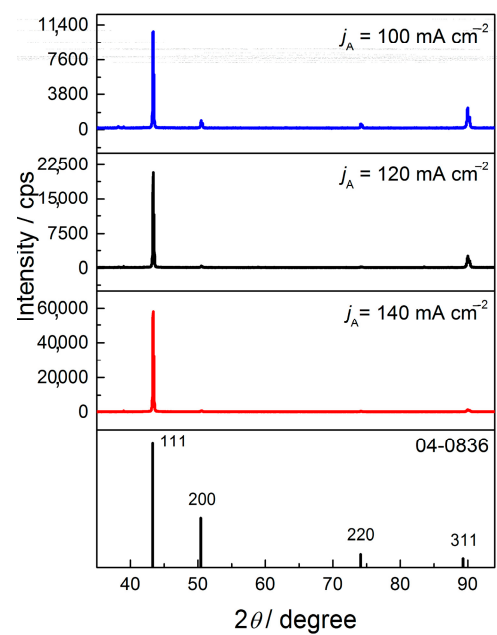


Figure 21. The XRD patterns recorded for the Cu coatings produced by the pulsating current regime with different current density amplitudes, and Cu standard (04-0836) [13].

Table 7. Texture calculations for the Cu coatings obtained by the PC regimes with j_A of 120 and 140 mA cm^{-2} ; s—Cu standard; R is an intensity of diffraction peak [13].

Plane (<i>hkl</i>)	R (in%)		R_s (in%)	TC		RTC (in%)	
	R_{120}	R_{140}		TC_{120}	TC_{140}	RTC_{120}	RTC_{140}
(111)	86.1	95.8	54.6	1.58	1.75	54.3	83.3
(200)	2.0	1.26	25.1	0.080	0.050	2.7	2.4
(220)	1.3	0.76	10.9	0.12	0.070	4.1	3.3
(311)	10.6	2.18	9.4	1.13	0.23	38.9	11.0

4.3. Influence of the Coating Thickness on Morphology of the Cu Coatings

Figure 22 shows the 3D AFM image and the histogram of topography of 20 μm thick Cu coating produced at j_{av} of 50 mA cm^{-2} . The other parameters of PC regime are $j_A = 100 \text{ mA cm}^{-2}$, $t_c = t_p = 5 \text{ ms}$.

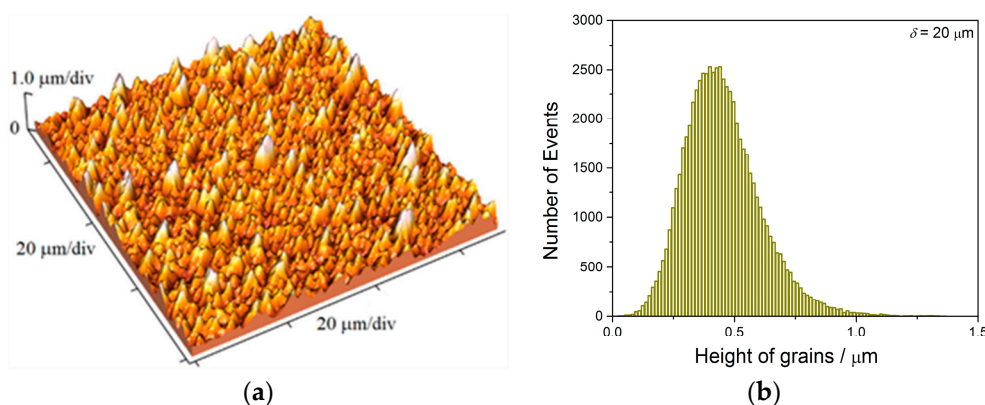


Figure 22. The 3D AFM image and the histogram of 20 μm thick Cu coating electrodeposited by the pulsating current regime at j_{av} of 50 mA cm^{-2} ; $j_A = 100 \text{ mA cm}^{-2}$; $t_c = t_p = 5 \text{ ms}$: (a) topography, and (b) histogram. The cathode: Si(111). Scan size: (70 \times 70) μm^2 [13].

The value of R_a roughness for this coating was $101.5 \pm 1.9 \text{ nm}$. By insight in Table 4, and comparing the values obtained for the same PC regime, it follows that similar to the DC regime, an increase of the electrodeposition time, i.e., the coating thickness causes an increase of the coating roughness.

4.4. Influence of the Type of Cathode on Morphology of the Cu Coatings

Figure 23 gives morphology of 40 μm thick Cu coating produced by the pulsating current regime at j_{av} of 50 mA cm^{-2} ($j_A = 100 \text{ mA cm}^{-2}$; $t_c = t_p = 5 \text{ ms}$) on the brass cathode. At the macro level, this coating was very similar to that obtained on the Si(111) electrode with the same PC parameters (Figure 16g,h). The grains of size of about 5 μm are mostly electrodeposited on this cathode.

The 3D AFM images and the histograms of topography of 20 and 40 μm thick coatings of Cu produced on brass cathode at j_{av} of 50 mA cm^{-2} are shown in Figure 24 (20 μm , Figure 24a,b and 40 μm , Figure 24c,d).

The R_a roughness values were $146.0 \pm 1.8 \text{ nm}$ for 20 μm , and $215.6 \pm 2.2 \text{ nm}$ for 40 μm thick coatings of Cu [45]. These values were higher than those obtained on the Si(111) substrate, that can be ascribed to, similar to the Cu coatings electrodeposited in the DC regime, higher initial roughness of the brass than the Si(111) cathodes.

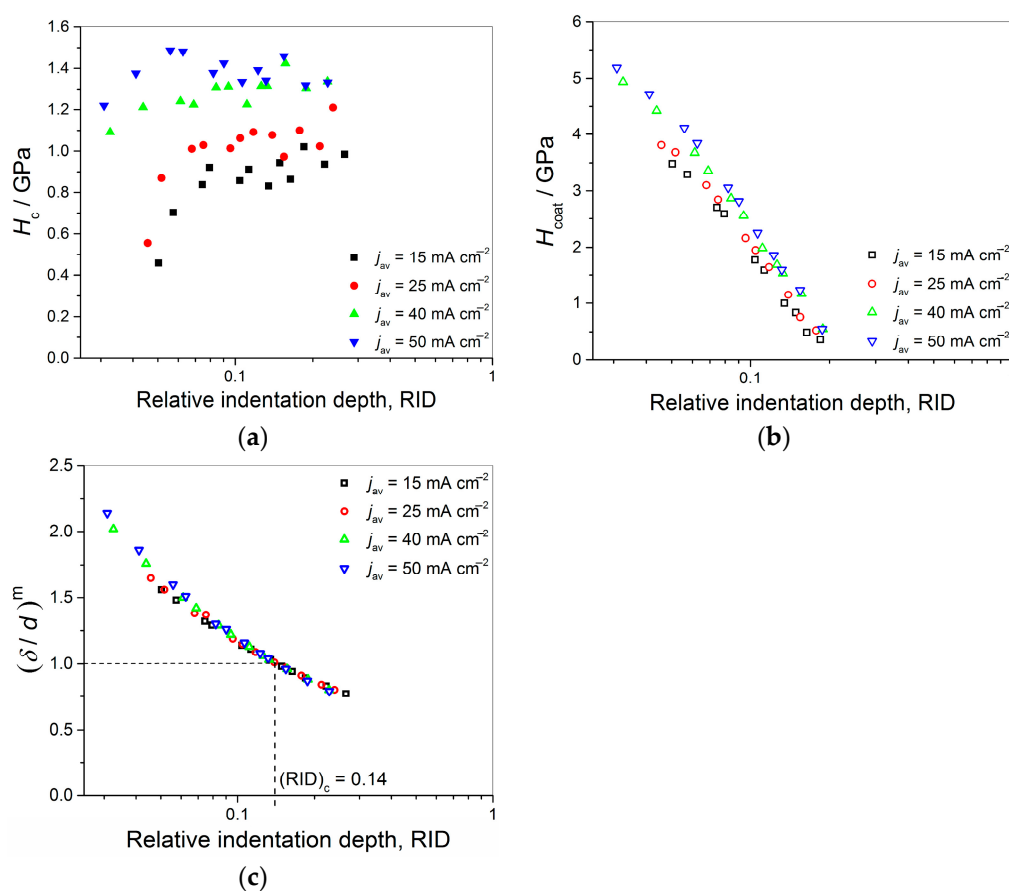


Figure 25. The dependencies of: (a) H_c , (b) H_{coat} , and (c) parameter $(\delta/d)^m$ on RID for the Cu coatings produced by the PC regime from electrolyte I on Si(111) substrate at frequencies of 30, 50, 80 and 100 Hz (i.e., at average current densities of 15, 25, 40 and 50 mA cm^{-2}) [13,54].

The m coefficient values for these coatings are given in Table 8 [13,54].

Table 8. The Meyer's index, m values for the Cu coatings electrodeposited by different PC regimes [13,54].

$j_{av}/\text{mA cm}^{-2}$	15	25	40	50	60	70
m	0.4288	0.4372	0.4770	0.4979	0.4346	0.3447

It follows from Figure 25b that the calculated H_{coat} values have typical descending trend of a change with the increase of RID values characteristic for Si(111) substrate. At the small RID values, the H_{coat} values were notably larger than the measured H_c values, and this difference decreased with an increase of RID value. For $\text{RID} \geq 0.14$, the measured hardness became larger than the calculated coating hardness. Taking into consideration the validity of the C–L model up to $(\delta/d)^m = 1$ [23,24,33,34], it is confirmed the limiting (or critical) RID value of 0.14 separating the zone where the measured hardness is equal to the coating hardness from the zone where an application of the C–L model is necessary to obtain the absolute coating hardness.

This limiting RID value of 0.14 is also confirmed for the Cu coatings obtained with different the current density amplitudes, and it is also valid for the Cu coatings of different thicknesses produced by the PC regime [54]. Figure 26 shows the dependencies of the H_c , H_{coat} , $(\delta/d)^m$ on the RID for 40 μm thick Cu coatings obtained by the pulsating current regime with the current density amplitudes, j_A of 120 and 140 mA cm^{-2} (or j_{av} of 60 and 70 mA cm^{-2} , respectively), and for 20 μm thick Cu coating electrodeposited at j_{av} of

50 mA cm^{−2} (see Table 3). The m coefficient values for the Cu coatings electrodeposited at j_{av} of 60 and 70 mA cm^{−2} are presented in Table 8, while m value for 20 µm thick Cu coating was 0.3591 [13,54].

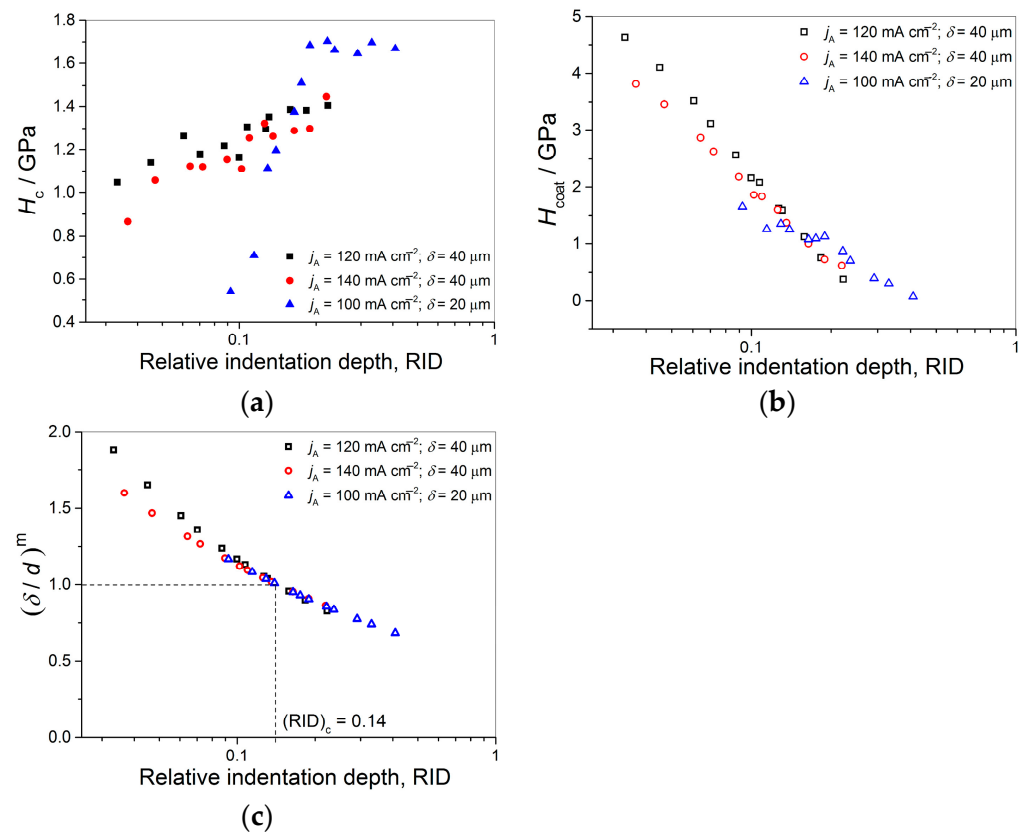


Figure 26. The dependencies of: (a) H_c , (b) H_{coat} , and (c) parameter $(\delta/d)^m$ on RID for the 40 µm thick Cu coatings produced by the PC regime from *electrolyte I* on Si(111) substrate at j_{av} (j_A) of 60 and 70 mA cm^{−2} (120 and 140 mA cm^{−2}, respectively), and 20 µm thick Cu coating electrodeposited under the same conditions at j_{av} (j_A) of 50 mA cm^{−2} (100 mA cm^{−2}) [13,54].

The effect of the kind of cathode on hardness of electrolytically produced Cu coatings is examined through analysis of the Cu coatings produced by the pulsating current regime on the brass substrate. Figure 27 shows the dependencies of the H_c , H_{coat} , $(\delta/d)^m$ on RID for 20 and 40 µm thick Cu coatings electrodeposited on the brass substrate at j_{av} (j_A) of 50 mA cm^{−2} (100 mA cm^{−2}). The m coefficient value for 20 µm thick coating is 0.4141, and 0.3744 for 40 µm thick coating [45].

According to expectation, the dependencies of H_c and H_{coat} on RID were very similar to each others (Figure 27a,b). The H_c and calculated H_{coat} values approached to a hardness of brass substrate at the high RID. At the low applied loads, H_{coat} were larger than H_c whereby the differences between H_c and H_{coat} were smaller than on Si(111) cathode. The validity of the C–L model $((RID)_c = 0.14$ for $(\delta/d)^m = 1$; Figure 27c) is confirmed for the Cu coatings electrodeposited by the PC regime on the brass cathode.

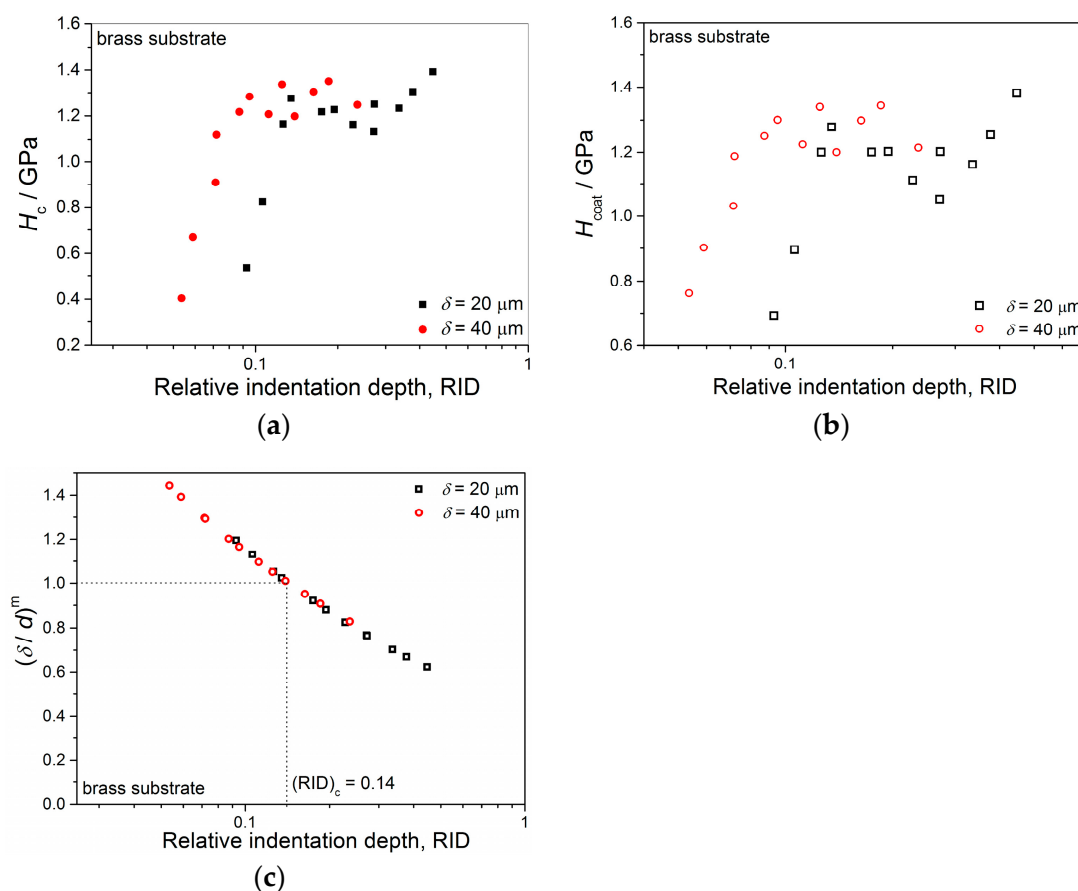


Figure 27. The dependencies of: (a) H_c , (b) H_{coat} , and (c) parameter $(\delta/d)^m$ on RID for 20 and 40 μm thick Cu coatings produced by the pulsating current regime from *electrolyte I* on the brass cathode at j_{av} (j_A) of 50 mA cm^{-2} (100 mA cm^{-2}) [45].

6. Correlation among Morphology, Structure and Hardness of Electrolytically Produced Copper Coatings

The two kinds of compact and uniform Cu coatings were obtained by electrodeposition in the DC regime. The fine-grained mat coatings are produced from the basic sulfate electrolyte (*electrolyte I*), and the very smooth mirror bright coatings from the electrolyte in the presence of additives for leveling and brightness (*electrolyte II*). The analysis of histograms confirmed the strong difference between them. In the mat Cu coatings, the grain heights are at the micro level, while in mirror bright Cu coatings the grain heights were at the nano level. The addition of the additives affected not only morphology of the coatings, but to crystal structure of the produced coatings. The mirror bright Cu coatings exhibited the strong (200) preferred orientation, while those produced from the basic sulfate solution exhibited the strong (220) preferred orientation.

The change in morphology and structure of the Cu coatings can be attributed to a synergistic effect of the used leveling/brightening addition agents on electrodeposition of copper. The adsorbed chloride layer is formed on electrode surface by an addition of small concentrations of chloride ions (1–10 mM). This adsorbed layer mediates copper reduction and introduces an additional reaction pathway, resulting in an overall depolarization of the reduction process. Both depolarization and the increase in peak current observed in the cyclic voltammograms (CVs) revealed a catalytic effect of chloride on Cu reduction [55]. PEG is copolymer and nonionic surfactant, acting as a suppressor of the electrodeposition process only in the presence of chloride ions [56]. There is no unique mechanism explaining adsorption/suppression effect of PEG in the presence of chloride ions on Cu electrodeposition process. The various mechanisms are proposed and they

are summarized in Ref. [56]. The MPSA is top brightening addition agent added in parts per million (ppm) concentrations to acid sulfate Cu electrolytes [56,57]. The addition of this additive to the basic sulfate electrolyte with already added chloride/PEG additives activates electrodeposition process enabling a formation of mirror bright coatings [10,11,57]. The typical model explaining synergistic effect of this combination of additives causing a formation of mirror bright coatings is known as the model of “local perforation” [58,59].

The electrolyte stirring is another important parameter of the electrodeposition affecting a quality of metal coatings, whereby a quality of the coating is strongly influenced by intensity of stirring of the electrolyte. The various intensity of electrolyte stirring can be imposed by application of ultrasound of various powers, and it can be considered as follows: the compact and uniform Cu coatings are produced at the current density, j of 50 mA cm^{-2} which belongs to the mixed activation-diffusion control of the electrodeposition process.

For this type of electrodeposition control, the overpotential of electrodeposition can be presented by Equation (7) [8,60]:

$$\eta = \frac{b_c}{2.3} \ln \frac{j}{j_0} + \frac{b_c}{2.3} \ln \frac{1}{1 - (j/j_L)} \quad (7)$$

where: b_c is the cathodic Tafel slope, j_0 is the exchange current density and j_L is the limiting diffusion current density.

The first term, η_{act}

$$\eta_{\text{act}} = \frac{b_c}{2.3} \ln \frac{j}{j_0} \quad (8)$$

is an activation part of overpotential required for the charge transfer, and the second term, η_{diff}

$$\eta_{\text{diff}} = \frac{b_c}{2.3} \ln \frac{1}{1 - (j/j_L)} \quad (9)$$

is a diffusion part of overpotential and it is due to mass transfer limitations (diffusion limitation primarily).

Ultrasound imposed to the electrodeposition process only effects on the diffusion part of overpotential. The imposed ultrasound causes a mixing of the solution in the near-electrode layer, causing the decrease of the diffusion layer thickness, and the increase of the limiting diffusion current density. The degree of the diffusion control in the mixed activation-diffusion control decreases (i.e., it decreases the diffusion part in Equation (7)) by electrolyte stirring, and this decrease enhances with an intensification of electrolyte stirring.

The effect of imposed ultrasound is equivalent to the other ways of electrolyte stirring, such as imposed magnetic field and magnetic stirring [60,61], but also to vigorous hydrogen evolution occurring parallel to metal electrochemical deposition [62]. The concept of „effective overpotential“ originally proposed to explain a change of the hydrodynamic conditions in the near-electrode layer caused by vigorous hydrogen evolution [62] is applicable for all other ways of electrolyte stirring. The production of different morphologies of the Cu coatings under different intensities of applied ultrasound (Figures 6–9) can be considered as follows: the solution stirring in the near-electrode layer intensifies with an increase of the intensity of imposed ultrasound. As a result of this intensification, the degree of the diffusion control in the mixed activation-diffusion control decreases. Since the activation part is independent of the electrolyte stirring, it means that the contribution of the activation control in the mixed activation-diffusion control increases with an increase of the intensity of ultrasound. This causes a formation of large and regular crystal grains in the coatings produced under strong imposed ultrasound (see parts in circles in Figure 6b). Namely, formation of regular and well defined grains is result of dominant activation control in the overall control of the electrochemical deposition process [8].

The equivalent effect on morphology and structure of electrodeposited Cu coatings is achieved by the pulsating current regime. The range of the average current densities between 15 and 70 mA cm^{-2} belongs to the mixed activation-diffusion control of the

electrochemical deposition process [13]. With the increase of the average current density, the share of the activation control decreased, while the share of diffusion control increased in the mixed activation-diffusion control of the electrochemical deposition. The regular and well defined grains formed with j_{av} of 15 mA cm^{-2} (Figure 16a,b) are characteristic of predominant activation control, while globules are formed with j_{av} of 70 mA cm^{-2} (Figure 19c) when diffusion becomes predominant process in the mixed activation-diffusion control of electrochemical deposition.

The change in coating morphology is accompanied by change of the preferred orientation from the strong (220) to the strong (111) preferred orientation with the increase of j_{av} value. This change of the preferred orientation can be explained by different growth rate on different crystal planes [63]. The rates of growth are associated with the surface energy values of crystal planes γ and for the FCC type of crystal lattice following the trend $\gamma_{111} < \gamma_{100} < \gamma_{311} < \gamma_{110}$ [64,65]. Following the values of surface energy, the electrodeposition rate is the smallest on the (111) crystal plane, and for that reason, the (111) plane represents “slow-growing crystal plane” surviving the electrochemical deposition process [66]. All other crystal planes, i.e., (100), (311) and (110) planes, are the “fast-growing crystal planes” and they disappear firstly during electrochemical deposition process. In this case, with an intensification of the electrodeposition process, i.e., with increasing the average current density from 15 to 70 mA cm^{-2} , it increases a share of Cu crystallites oriented in the (111) crystal plane, while it decreases a share of crystallites in all other crystal planes, especially in (220) crystal plane (Figures 18 and 21, and Tables 5–7).

The mat Cu coatings produced from the basic sulfate electrolyte (*electrolyte I*) showed larger coating hardness than mirror bright Cu coatings produced in the presence of leveling/brightening additives (*electrolyte II*). The processes occurring on boundary among grains are responsible for this difference [12]. Namely, mat coatings have fine-grained structure with numerous boundaries among grains. The boundaries among grains are a disruption sites for a dislocation motion, and they prevent a movement of the dislocations [46,67]. It can be said that the boundaries among grains behave as mini resistors which resist during indentation in the metal coating. On the other hand, mirror bright Cu coatings were very smooth, without clear boundary among grains, and dislocation motion is a dominant mechanism affecting the coating hardness.

Analysis of histograms for the coatings electrodeposited from *electrolyte I* by both DC and PC regimes clearly revealed their microcrystalline (mc) character, and for them, Hall-Patch equation [68] predicting an increase of coating hardness with a decrease of grain size is valid. On the other hand, the histogram analysis of smooth mirror bright coatings revealed their nanocrystalline (nc) character with size of grains below critical value for a validity of Hall-Patch equation, and for these coatings, the inverse Hall-Petch equation begins to hold [69].

The values of both composite and calculated coating hardness on Si(111) were larger than the values obtained on the brass cathode, what clearly indicate that a determination of coating hardness is complex phenomena depending on many factors, such as the indenter geometry, plastic pile-up effect, film/substrate adhesion, elastic properties of the film and substrate and the friction between the indenter and film [70–76]. The pile-up effect affecting the coating hardness value can be the easiest looked at through analysis of the Cu coatings electrodeposited from *electrolyte I* and *electrolyte II*, as seen from Figure 28 showing Vickers indents.

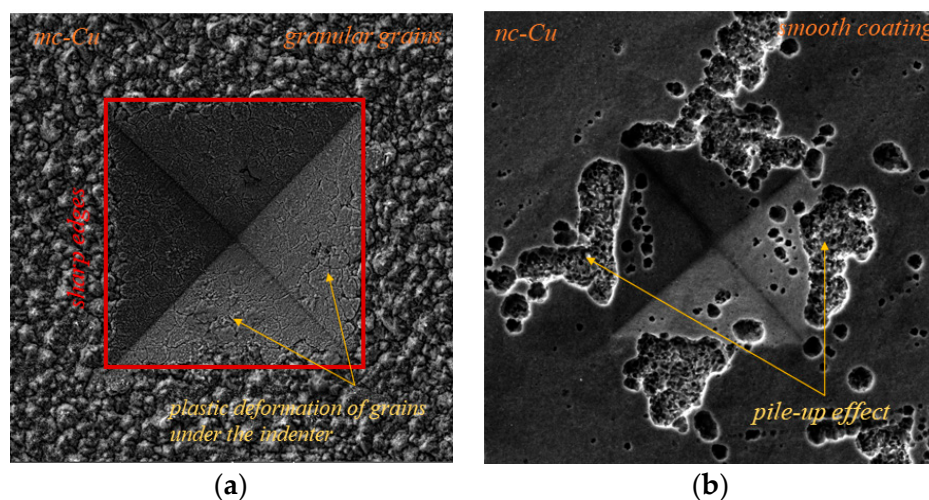


Figure 28. The deformation of the Cu coatings around an indenter as a function of the applied load: (a) the Cu coating produced from *electrolyte I* in the DC mode at $j = 33 \text{ mA cm}^{-2}$, ($P = 4.903 \text{ N}$, $d = 84.4 \text{ }\mu\text{m}$), and (b) the Cu coating produced from *electrolyte II* in the DC mode at $j = 67 \text{ mA cm}^{-2}$ with ultrasound assistance ($P = 0.981 \text{ N}$, $d = 39.6 \text{ }\mu\text{m}$) [18].

In the Cu coating electrodeposited from *electrolyte I*, the main mechanism is a plastic deformation of the coating under the indenter (Figure 28a) [18]. The plastic deformation of fine-grained coatings produced from *electrolyte I* occurs throughout the grain with the slip of dislocations mostly in the grain interior (intragranular slip) [77]. In the nanocrystalline coatings electrodeposited from *electrolyte II*, a pile-up effect occurs at edge of the indent (Figure 28b), and this effect is not observed in the Cu coating electrodeposited from *electrolyte I*.

Anyway, the limiting (or critical) RID value of 0.14 [$(\text{RID})_c = 0.14$] showed an universal character for electrolytically produced Cu coatings. This value was observed for both the Cu coatings electrodeposited in the DC regime under various electrodeposition conditions (the types of electrolyte and cathode, coating thickness and electrolyte stirring) and for various PC regimes. This value is related with the coating thickness as $(h_c/\delta) = 0.14$, i.e., as $h_c = 0.14\delta$. For values of an indentation depth smaller than this critical value, h_c , (i.e., for $h_c < 0.14\delta$) a substrate (cathode) had not any significant influence on the coating hardness and the measured hardness can be equaled with the coating hardness. For $h_c \geq 0.14\delta$, the cathode hardness strongly contributes to the measured composite hardness, and it is necessary to apply the C–L model to obtain an absolute hardness of the Cu coatings. This criteria ($h_c = 0.14\delta$) is situated between the value $h_c < 0.20\delta$ [70] established for polycrystalline soft coatings on hard substrates. Such as, the Cu coatings produced on Si(111) and brass cathodes, for which a substrate had no influence on the coating hardness, and Buckle's one-tenth rule predicting that a film thickness of 10 times the indentation depth is sufficient to neglect the influence of the substrate, i.e., a substrate commences to effect on the coating hardness after $h_c > 0.10\delta$ [51].

The coating hardness values calculated applying the C–L model were in a line with those obtained in other investigations for the Cu coatings obtained by different electrochemical deposition processes. Depending on parameters of the electrodeposition, the values of hardness of Cu coatings electrodeposited by the DC regimes were between 0.70 and 1.65 GPa [61,78,79]. For example, the Cu coating hardness value determined by Korsunsky model was 0.80 GPa [41]. The values of coating hardness for the Cu coatings produced by different periodically changing regimes of the electrochemical deposition were in the 1.10–2.0 GPa range, and they were slightly higher than the values obtained by the DC mode [80].

7. Copper Matrix Composites

In the last decades, an effort in a production of Cu coatings suitable for application in various industries is moved towards an improvement of their mechanical characteristics. It is attained by an incorporation of various particles, known as reinforcements, into the Cu coatings. The particles are suspended into an electrolyte and incorporated (co-deposited) into a Cu coating during electrodeposition process, producing copper matrix composites (CMCs) [81]. All reinforcements can be situated into one of two groups [82]:

- I. Metal ceramics, such as zirconia (ZrO_2) [83], Al_2O_3 [84,85], nitrides (Si_3N_4) [81,86,87], silicon carbide (SiC) [88,89], silica (SiO_2) [90], TiO_2 [91,92], etc., and
- II. Carbon nanomaterials, such as carbon nanotube (CNT) [82,93], multi walled carbon nanotube (MWCNT) [94,95], and graphene (Gr) and its derivatives such as (graphene oxides (GO) and reduced graphene oxides (RGO)) [96–102].

The addition of the first group of the reinforcements to the electrolyte improves a hardness of electrodeposited Cu coatings, but always causes a decrease of the electrical conductivity due to inferior electrical conductivity of metal ceramic particles [82]. On the other hand, carbon nanomaterials possess excellent both electrical and thermal conductivities, making them superior in a production of copper matrix composites relative to those obtained with ceramic reinforcements. The combination of two types of particles, such as Gr and CNT, is often used for a synthesis of Cu matrix composites of enhanced mechanical characteristics [82].

The processes of electrodeposition from electrolytes with suspended particles by which copper matrix composites are produced are known as electrolytic co-deposition or composite plating processes [81]. All parameters and regimes of the electrodeposition reported in Sections 2 and 4 are also applied for obtaining of Cu matrix composites [81–102].

Anyway, various research groups reported about an enhanced hardness of Cu coatings electrodeposited with the reinforcements relative to those obtained from reinforcements-free electrolytes [81,83,97]. Also, the existence of optimal concentration of particles such as ZrO_2 giving the coatings of maximum hardness is reported [83]. The decrease of the coating hardness after an attained maximum value can be attributed to a process of agglomeration with the high, applied concentration of the particles. The increase in hardness of coatings can be attributed to [97]: (a) particle strengthening—related with the incorporation of hard particles in a coating and a volume fraction above 20%, (b) dispersion strengthening—related with the incorporation of fine particles ($<1\ \mu\text{m}$) and a volume fraction lower than 15%, and (c) grain refining—related with the nucleation of small grains on surface of the incorporated particles, resulting in a general structural refinement. The presence of smaller grains in coating impedes a dislocation motion, resulting in an increase of microhardness.

8. Conclusions

The effect of various parameters and regimes of the electrochemical deposition on morphology, structure and hardness of copper coatings is summarized. The coatings of copper were obtained by galvanostatically varying the following parameters: the kind of solution (the basic sulfate solution and a solution with an added leveling/brightening additives), the kind of cathode (Si(111) and brass), the electrolysis time (or coating thickness) and stirring of the electrolyte (magnetic stirring and stirring by ultrasound). The Cu coatings of different thicknesses were produced from basic sulfate electrolyte on both Si(111) and brass cathodes by pulsating current (PC) regime varying the average current density (or frequency) and the current density amplitude. Morphology and structure of the Cu coatings were characterized by scanning electron microscopy (SEM) and by atomic force microscopy (AFM) techniques and X-ray diffraction (XRD), respectively, and hardness of the Cu coatings was examined by Vickers microindentation. The coatings of Cu on the Si(111) and the brass substrates belong to “soft film on hard substrate” composite hardness system, and the Chicot–Lesage (C–L) composite hardness model was applied to estimate a hardness of the Cu coatings.

On the basis of presented analyses, it follows that application of the C–L model represents valuable way for an estimation of hardness of the copper coatings. Applying this model, it is established the critical (or limiting) relative indentation depth (RID) of 0.14 separating the zone where measured (composite) hardness is equal to hardness of the coating from the zone where it is necessary to apply the C–L model to determine absolute hardness of the Cu coatings. Considering representative parameters in the DC regime and applying the PC regime as type of PRC regimes, it can be concluded that this critical RID value has universal character for the Cu coatings obtained by different electrodeposition regimes.

Author Contributions: Conceptualization, N.D.N.; investigation, I.O.M.; writing—original draft preparation N.D.N.; methodology, N.D.N. and I.O.M.; formal analysis, I.O.M.; writing—review and editing, N.D.N. and I.O.M.; software, I.O.M.; validation, N.D.N. and I.O.M.; visualization, N.D.N. and I.O.M.; supervision, N.D.N.; funding acquisition, N.D.N. All authors have read and agreed to the published version of the manuscript.

Funding: This work was financially supported by Ministry of Science, Technological Development and Innovation of Republic of Serbia (Grant No. 451-03-47/2023-01/200026).

Institutional Review Board Statement: Not applicable.

Informed Consent Statement: Not applicable.

Data Availability Statement: The data presented in this study are available on request from the corresponding author or co-authors. The data are not publicly available.

Acknowledgments: This work was funded by Ministry of Science, Technological Development and Innovation of Republic of Serbia.

Conflicts of Interest: The authors declare no conflict of interest.

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