# EFFECT OF YTTRIUM OXIDE MODIFICATION OF AI-SI ALLOY ON MICROHARDNESS AND MICROSTRUCTURE OF SURFACE LAYERS

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Microhardness, structure and phase state of silumin modified by coating with yttrium oxide when electroexplosive alloying were analyzed. Appropriate processing modes were estimated according to the results of research. The study on the structure and phase composition was carried out for the processing mode, which increases microhardness of the surface layer more than twice. A complex of research procedures was used to analyze the structure and phase composition. As revealed in the study, the coatings with a thickness of  $50 - 80 \,\mu\text{m}$  are highly rough and porous, they have a sub-micro- and nano-scaled multiphase structure with particles of silicon,  $Y_2O_3$ ,  $YSi_2$  and  $Y_2Si_2O_7$  as strengthening phases, and alloying elements (silicon, yttrium, and oxygen) don't tend to spread homogenously in them.

Key words: Al-Si alloy, yttrium oxide, structure, microhardness, wear resistance.

#### INTRODUCTION

To date, alloys of non-ferrous metals are widely used instead of steel alloys. Their physical and mechanical properties are similar to those of steel, some of them are even better. It is said that the most prospective alloys are those on the base of aluminum and silicon (silumin). Since silumin has good plasticity, the alloy can follow the most complex shapes, filling casts evenly. As a consequence, casting of silumin is simplified, reducing, therefore, production costs [1-3]. The main advantage over other aluminum alloys, including other types of silumins is quite good casting properties, first of all high fluidity. These casting alloys are suitable for casting of thinwalled, complex and leak-proof products, which are resistant to vibration and impact loads [4, 5].

The dominating trend to make motors economically and ecologically compatible is the reason for attempts of designers, who modify the rod and piston group, e.g. reduce its weight, and change their geometry, making the shape of pistons more complex, this way. However, the loads on the rod and piston group are not reduced, since the capacity of motors is the same, deteriorating significantly the resource, as a consequence. Therefore, it is urgently needed to find new and efficient methods to improve properties of silumins, which involve modifying, micro-alloying, and surface treatment [8-16]. The promising procedure of outer energy impacts, conditioning serious changes in the structure, phase composition, physical and mechanical properties of the surface layers of metals and alloys is electroexplosive alloying. Using this method, coatings with high quality, functional characteristics, and sufficient adhesion with the substrate as well, can be produced. Due to this technique it is possible to produce coatings from conductor explosion products, and form composite surfaces with qualities better than those of the initial material [17].

To sum up, this work aims at modification of surface layers in alloy Al-Si with yttrium oxide particles in electroexplosive alloying with the consequent analysis of the structure, phase composition and mechanical properties of produced coatings.

#### **EXPERIMENTAL**

Plates of eutectic Al-Si alloy (silumin) (11,1 Si, 0,58 Mg, 2,19 Cu, 0,92 Ni, 0,25 Fe, 0,029 Mn, 0,047 Ti, 0,005 Cr, balance Al (wt. %)) were analyzed. The samples under consideration were  $20 \times 20 \times 10$  mm and oriented perpendicular to the plasma jet axis. Electroexplosive alloying was carried out using a laboratory pulse-discharging electroexplosive equipment EVU 60/10 [15]. Electrical current with a high density (~ $10^{10}$  $A/m^2$ ) passes through the conductor when a capacity storage discharges, resulting this way, in electrical explosion. We used aluminum foils to explode conductors and Y<sub>2</sub>O<sub>3</sub> as a weighted portion of powder. The treatment was carried out in three processing modes, which differ in discharge voltage, weights of foils to be exploded and powder portions. All processing modes are listed in Table 1.

The measurement of micro-hardness was used to describe mechanical properties of the surface layers. The difference in micro-hardness before and after treatment can be considered to indicate strengthening of the sur-

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Processing mode	Weight of alu- minum foil / g	Weight of powder $Y_2O_3 / g$	Discharge voltage / kV	
1	0,0589	0,0589	2,6	
2	0,0589	0,0589	2,8	
3	0,0589	0,02945	2,6	
4	0,0589	0,02945	2,8	
5	0,0589	0,0883	2,6	
6	0,0589	0,0883	2,8	

Table 1 Processing modes of electroexplosive alloying

face layers in metals and alloys. Micro-hardness was determined using the device to measure micro-hardness HVS-1000 and Shimadzu DUH-211S to estimate ultramicro-hardness according to Vickers method.

Scanning electron microscopy (SEM) with the help of SEM-515 Philips, tooled with micro-analyzer EDAX ECON IV was the technique to analyze the element and phase composition, and defect substructure of the modified layer. The phase composition of modified layers, that is, qualitative and quantitative characteristics of various phases in case there are any, their content, dispersion, structure and chemical composition were assessed using electron diffraction microscopy and X-ray spectrometry (diffractometer XRD-7000s, Shimadzu). The defect structure of samples was studied via STEManalysis of thin foils (JEM-2100F, JEOL). The data on thin structure of the material were used to classify morphological characteristics of the structure.

All data obtained were processed statistically, taking into account at least 10 measurements in each mode.

### **RESULTS AND DISCUSSION**

Micro-hardness in the sprayed layer and in the substrate of samples treated in electroexplosive alloying was measured in different points from the treated surface. Micro-indentation was carried out on crosscut sections. Analyzing the change in micro-hardness as far from the treated surface, it was identified that processing modes 2 and 5 (Figure 1a) provide optimal characteristics (thickness and micro-hardness of the alloyed layer) in the treated surface.

As seen in Figure 1(a), micro-hardness of the material is maximal in the surface layer and exceeds that of the initial material more than twice. As far from the modified surface, the drop of micro-hardness is registered and at a depth of  $\approx 90 \ \mu m$  micro-hardness equals that of silumin in the initial state. The sprayed layer, processed in Mode 2 is 50  $\mu m$  thick (Figure 1b), and in Mode 5 – 80  $\mu m$ .

As revealed, the optimal processing modes are Mode 2 and Mode 5. For the further analysis samples processed according to mode 2 were used to identify their phase composition and structure.

The structure of silumin volume after electroexplosive alloying was analyzed by the method of sections using optical (Figure 1b) and scanning electron microscopy (Figure 2). Studying the data in Figure 2, one can





**Figure 1** a) Micro-hardness profile of silumin after electroexplosive alloying (the load on indenter 50 mN). b) The coating of the system Al-Y<sub>2</sub>O<sub>3</sub> produced by the method of electroexplosive alloying (zooming 20 X optical microscopy – processing mode 2.

note that the thickness of the modified layer varies in the range  $30 - 50 \mu m$ , and the level of porosity is high. The pores are throughout the modified layer; their sizes are ones to tens of micrometer.

The phase composition of silumin modified via electroexplosive alloying was studied by the methods of Xray analysis. The data on the phase analysis of the material are given in Table 2.

Considering the results given in Table 2, first, a rather high amount of silicon is registered in the surface layer of



Figure 2 SEM - The structure of silumin section after electroexplosive alloying.

Phase	Content, rel. / %	Parameter of the lattice / nm	D (CSZ) / nm	∆d/d / 10 <sup>-3</sup>	
Al	68,2	0,40485	75,01	0,24	
Si	30,1	0,54231	16,4	0,80	
Y <sub>2</sub> O <sub>3</sub>	1,7	1,06010	16,6	7,88	

Table 2 The results of X-ray analysis of eutectic silumin after electroexplosive alloying

CDZ - coherent scattering zone

silumin that might be caused by emission of some aluminum layer when electroexplosive alloying. Secondly, there is  $Y_2O_3$  phase, maybe, due to particles of initial powder of yttrium oxide, penetrating into the surface layer of silumin when electroexplosive alloying. The findings of X-ray micro-spectral analysis, in particular, the element composition in the surface layer of silumin after electroexplosive alloying are given in Table 3.

Analyzing the results given in Figure 3 and Table 3, it should be noted that after electroexplosive alloying the surface layer is formed with a high level of alloying elements distributed non-homogeneously, it is most visible in spreading of yttrium and oxygen atoms. To be more precise, there are zones with concentration of yttrium and oxygen exceeding the average one twice and more. These results confirm that in the plasma flow of powder particles there is an alloying material identified in some studies before. Furthermore, the surface layer with a high level of roughness containing a lot of micropores, micro-craters and micro-cracks is formed in electroexplosive alloying.

The defect sub-structure, element and phase composition of the surface layer in silumin after electroexplosive alloying were also analyzed using the methods of transmission electron diffraction microscopy of thin foils. As revealed, a structure with cellular crystallization of aluminum is formed in the surface layer because the modified layer gets cooled at a high-speed when electroexplosive alloying. The sizes of cells vary from 200 to 450 nm. On borders of the cells there are interlayers of the second phase. X-ray micro-spectrum analysis showed that the interlayers are formed by silicon and yttrium atoms.

The phase composition of silumin after electroexplosive alloying was studied by the methods of transmission diffraction electron microscopy using the procedure of dark-field analysis and indention of respective electron-diffraction micro-patterns (Figure 4).

Analyzing Figure 4, it is obvious that the surface layer of silumin after electroexplosive alloying has a structure of cellular crystallization. The cells are of a clearly oval form. Their sizes vary from 150 nm to 350 nm. According to the micro-diffraction analysis the cells are a solid solution based on aluminum.

In the volume of cells particles of the second phase are identified with the help of dark-filed analysis; the sizes of particles are in units of nanometers (Figure 4c).

With respect to electron-diffraction micro-pattern (XRD) (Figure 4b) we can say with a good reason that



Figure 3 SEM data on the structure of silumin surface treated by the electroexplosive method: a), b) see Table 3

Table 3	The element composition of the surface layer in
	silumin after electroexplosive alloying, identified
	by the methods of X-ray micro-spectral analysis of
	zones shown in Figure 3 / wt. %

Zone	Al	Si	Mg	Ti	Fe	Ni	Cu	Y	0	С
Figure 3a	47,2	3,0	0,6	1,0	0,7	1,3	1,8	16,2	10,8	17,4
Figure 3b	0,8	0,0	0,0	0,2	0,7	0,5	0,7	34,0	28,1	35,0

these particles are yttrium silicide of the composition  $YSi_2$ . The aluminum cells are distributed as interlayers of the second phase. As seen in the electron-diffraction micro-pattern (Figure 4b), the interlayers are formed by silicon and yttrium silicate of the composition  $Y_2Si_2O_7$ .

## CONCLUSION

The surface layer in eutectic silumin was modified with yttrium oxide using the method of electroexplosive alloying. The appropriate processing modes were revealed: Mode 2 (discharge voltage – 2,8 kV, weight of aluminum foil – 0,0589 g, weight of yttrium powder  $Y_2O_3 - 0,0589$  g.), and Mode 5 (discharge voltage – 2,6 kV, weight of aluminum foil – 0,0589 g, weight of yttrium powder  $Y_2O_3 - 0,0883$  g.). It was revealed that as a result of electroexplosive alloying of silumin a highlyporous surface layer is formed with a thickness from 50



Figure 4 XRD data of the silumin structure after electroexplosive alloying; a) bright field; b) electron-diffraction micro-pattern, arrows indicate reflexes, where dark fields were obtained; c) dark field obtained in reflexes 2 [111]AI + [112]YSi,.

to 80  $\mu$ m; alloying elements (Si, Y and O) don't tend to spread homogeneously in this layer, it also has sub-micro- and nano-scaled multiphase structure with particles of silicon, Y<sub>2</sub>O<sub>3</sub>, YSi<sub>2</sub> and Y<sub>2</sub>Si<sub>2</sub>O<sub>7</sub> as strengthening phases, its wear resistance and micro-hardness are high.

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