

SURFACE HIGH SPEED STAINLESS STEEL ALLOYING WITH COPPER **Yu.F. IVANOV, E.A. PETRIKOVA, A.D. TERESOV, N.A. PROKOPENKO, M.S. PETYUKEVICH**Institute of High Current Electronics SB RAS, Tomsk, Russia*

The using copper as an alloying element, the addition of which in small concentrations to low-carbon steel instead of expensive elements - niobium, titanium and vanadium, leads to the appearance of high corrosion and mechanical characteristics associated with the formation of Fe-Cu precipitates in the bulk of the material [1-3]. It has been established that these precipitates are nanosized particles of a saturated (more than 1% at.) solid solution of copper in iron, while in the equilibrium state the maximum solubility of copper in iron does not exceed 0.38% at. In this case, one should speak about the properties of the material as a function of near-surface transition states. Nanosized copper-enriched particles in α -Fe formed during cooling provide high ductility and fracture toughness and cause dispersion strengthening of the steel.

The formation of the “film (Cu)/(steel 321) substrate” system was carried out on the QUINTA installation [4] by sputtering copper films 0.5 μm thick onto steel specimens. High-speed alloying of steel with copper was carried out by irradiating the “film (Cu)/(steel 321) substrate” system with a pulsed electron beam using a SOLO setup [4]. The irradiation mode corresponded to the liquid-phase alloying of the steel surface layer with copper.

Studies performed by scanning electron microscopy showed that at an electron beam pulse duration of 50 μs (15 J/cm², 15 pulses, 0.3 s⁻¹) a nanocrystalline structure with a crystallite size of (80-120) nm is formed on the specimen's surface. At an electron beam pulse duration of 200 μs (30 J/cm², 15 pulses, 0.3 s⁻¹), regions with a lamellar structure are formed on the specimens surface. A structure of cellular crystallization is observed in the bulk of the plates. The cell sizes vary within (0.58-0.81) μm . X-ray microanalysis revealed a decrease (more than 4 times) in the concentration of copper in the steel surface layer with an increase in the duration of exposure to the electron beam from 50 μs to 200 μs .

Using X-ray phase analysis methods, it was shown that at an electron beam pulse duration of 50 μs , a solid solution of copper in a crystal lattice based on γ -Fe and a Fe_{0.5}Cu_{0.5} phase with a bcc crystal lattice are formed in the surface layer. With an increase in the pulse duration to 200 μs , a two-phase structure is formed in the surface layer - γ -Fe and a phase of Fe_{0.7}Cu_{0.3} composition, which has an fcc crystal lattice.

Cu as a separate phase is not detected. An increase in the electron beam pulse duration from 50 μs to 200 μs leads to an increase in the crystal lattice parameter of γ -Fe from 0.35191 nm to 0.35300 nm. Taking into account the ratio of the sizes of the atomic radii of Fe ($R(\text{Fe}) = 0.126$ nm) and Cu ($R(\text{Cu}) = 0.128$ nm), we can conclude that the process of replacing iron atoms in the crystal lattice of the γ -phase by copper atoms increases with an increase in the duration of the electron beam pulse, which leads to an increase in the lattice parameter.

Thus, a two-stage mechanism of solid solution decomposition in the “film (Cu) / (steel 321) substrate” system irradiated with a pulsed electron beam was revealed, as a result of the performed studies. At an electron beam treatment duration of 50 μs , the formation of nanosized particles of the Fe_{0.5}Cu_{0.5} phase, which has a bcc crystal lattice, is observed. With an increase in the duration of exposure to the electron beam to 200 μs , the formation of Fe_{0.7}Cu_{0.3} composition phase, which has an fcc crystal lattice, is recorded in the steel surface layer.

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