Phase Formation in Contacting Zinc Melting with Indium and Stannic in the Presence of Strontium Impurity and Microhardness of Derived Alloys

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Abstract — Phase formation during contact melting, including in the presence of electric transfer, in the system indium-zinc and stannic-zinc with the introduction of a small admixture of strontium was studied. The formation of two-component intermetallic compounds has been established. An attempt has been made to explain the obtained results.

Keywords — contact melting, electrotransport, X-ray phase analysis, intermetallide

I. INTRODUCTION

It is known that various metallic admixtures can significantly change the properties of alloys and composite materials, and sometimes even lead to the formation of new substances. In this regard, it seems relevant to study the effect of small additives on the phase transition between two solid dissimilar materials, known as contact melting (CM). In addition, if impurities are chemically active, they will form new compounds. Considering all the above, we continued the study of metal CMs in the presence of alkaline-earth additives. In this work, zinc with a small admixture of strontium was used.

II. METHODS AND MATERIALS

CM of zinc solid solutions with indium and stannic was carried out in unsteady-diffusion mode in a thermostat filled with silicone oil PPHMS-2/5L [1]. Temperature experiments were 147ºС and 202ºС. The experiments lasted 4 hours.

Also during the experiments, the effect of electromigration was studied [2,3]. For this, in one case, no electric current passed through the samples, and a constant electric current with a density of 0.5 A / mm² was passed through the other two. In this case, the direction of the current in them was the opposite. This condition is due to the fact that due to the difference in the effective charges [4] of different metals, depending on the polarity of the electromagnetic field, contact melting is accelerated or slowed down. The result is determined by the so-called “integral criterion of electric transfer” given in [1-5].

An X-ray phase analysis of the diffusion zones obtained was performed. The studies were carried out using a D2 Phaser X-ray diffractometer. The radiation wavelength was 1.54 Å, the width of the emitted beam was 0.8 mm. Also, the microhardness of the contact zones was determined in the
studied systems. The measurements were carried out on a PMT-3 microhardness meter according to the method described in [6].

III. RESULTS

Zinc-indium system

Tables 1 and 2 show the intermetallic compounds found in the diffusion zones during contact melting in the system (Zn + 0.5 at.% Sr) – In and (Zn + 0.74 at.% Sr) – In, respectively, as well as the parameters of their crystal lattices.

### TABLE I. INTERMETALLIDES AND PARAMETERS OF THEIR CRYSTAL LATTICES IN THE SYSTEM (Zn + 0.50 at.% Sr) – In

<table>
<thead>
<tr>
<th>Intermetallic</th>
<th>Conductance, A/mm²</th>
<th>Crystalline lattice</th>
</tr>
</thead>
<tbody>
<tr>
<td>SrZn₁₁</td>
<td>+</td>
<td>Tetragonal (10.749 Å x 6.899 Å)</td>
</tr>
<tr>
<td>SrIn₁₁</td>
<td>-</td>
<td>Tetragonal (8.738 Å x 16.442 Å)</td>
</tr>
</tbody>
</table>

Note that in the system (Zn + 0.5 at.% Sr) – In in all cases, the compound SrZn₁₁ was formed, which, however, was absent in the Sr-Zn state diagram [7]. Apparently, it was already present in the solid solution of zinc with strontium and did not completely dissolve upon contact melting. With the accelerating direction of the current, there is also intermetallic compound SrIn₁₁. It is observed in the equilibrium melting diagram of the In-Sr system. The formation of this substance may be due to more intensive leaching of the SrZn₁₁ compound to an area rich in indium. As a result, some SrZn₁₁ could disintegrate, and strontium was captured by indium to form a new compound.

In the system (Zn + 0.74 at.% Sr) – In (table 2), a slightly different picture is observed. Only strontium compounds with zinc are present. If in two versions of the experiment the compound SrZn₁₁ was again detected, then the SrZn₁₁ intermetallic was formed at the current slowing down direction. The second substance, in contrast to the first, is available on the Sr-Zn state diagram [7].

Figure 1 shows the results of measuring the microhardness of the contact layer of the system (Zn+0.5 at.% Sr) - In obtained in the absence of electric current.

Low values of the microhardness of the interlayer (within 40-60 MPa) compared with the microhardness of zinc (about 500 MPa) indicate an extremely low content of zinc in the resulting alloy. In this case, the zone adjoining the solid solution (Zn + 0.5 at.% Sr) is harder than the rest due to the smaller amount of indium in it. However, the further course of the microhardness is difficult to explain only by a decrease in the zinc content in the alloy, since at first this value drops rather sharply and then slightly increases. It can be assumed that in the compound zone the intermetallic compounds were unevenly distributed, which led to such non-monotonic microhardness behavior.

![Microhardness (Hμ) along the contact layer obtained at CM in the system (Zn + 0.50 at.% Sr) – In; d is the distance from the measured point to the boundary with strontium solid solution in zinc](image1)

Figure 2 presents the results of measuring the microhardness of the contact layer of the system (Zn + 0.74 at.% Sr) – In, obtained in the absence of electric current.

![Microhardness (Hμ) along the contact interlayer obtained at a CM in the system (Zn + 0.74 at.% Sr) – In; d is the distance from the measured point to the boundary with the solid solution of strontium in zinc](image2)

The microhardness of the alloy along the interlayer in this experiment is not explained by the usual arguments about the distribution of components along the zone. In the region lying near the solid solution (Zn + 0.74 at.% Sr), the microhardness lies within 50–60 MPa, and it does not decrease with the distance from the boundary. This is followed by a sharp decrease to about 30 MPa, and also does not decrease when approaching the indium sample. There is only an assumption that for some reason intermetallic compounds do not penetrate into the second half of the diffusion zone, creating a barrier for zinc too.

Zinc – stannic system
Tables 3 and 4 show the intermetallic compounds found in the diffusion zones during contact melting in the system (Zn + 0.5 at.% Sr) –Sn and (Zn + 0.74 at.% Sr) –Sn, respectively, as well as the parameters of their crystalline lattices.

**TABLE III. INTERMETALLIDES AND PARAMETERS OF THEIR CRYSTALLINE LATTICES IN THE SYSTEM (Zn + 0.5 AT.% Sr) –Sn**

<table>
<thead>
<tr>
<th>Intermetallic</th>
<th>Conductance, A/mm²</th>
<th>Crystalline lattice</th>
</tr>
</thead>
<tbody>
<tr>
<td>SrZn₁₁</td>
<td>0</td>
<td>Tetragonal (10.749 Å x 6.899 Å)</td>
</tr>
<tr>
<td>SrZn₁₃</td>
<td>+0.5</td>
<td>Cubical (12.242 Å)</td>
</tr>
<tr>
<td>SrSnₓ</td>
<td>-0.5</td>
<td>Monoclinal (12.175 Å x 4.062 Å x 5.163 Å)</td>
</tr>
</tbody>
</table>

**TABLE IV. INTERMETALLIDES AND PARAMETERS OF THEIR CRYSTALLINE LATTICES IN THE SYSTEM (Zn + 0.5 AT.% Sr) –Sn**

In the system (Zn + 0.5 at.% Sr) –Sn, by passing an electric current, a SrZn₁₁ compound is formed, which is absent in the Sr-Zn state diagram [7]. In the currentless variant of the experiment, the intermetallic compound SrZn₁₃ is found instead. There is a logical assumption that, in the presence of current, SrZn₁₃ decomposes with the formation of SrZn₁₁ and the release of free zinc. Check this in our experiments is not possible. At the same time, when the current is slowing down, the SrSnₓ compound also appears.

In the system (Zn + 0.74 at.% Sr) –Sn, the SrZn₁₁ intermetallic compound is present in the absence of current and its accelerating direction. In these, it is the only compound in the contact zone. When the current is slowing, it disappears. In this case, two other intermetallic compounds are formed - SrSnₓ and SrZnₓ, which are present on the Sr-Sn and Sr-Zn state diagrams [7]. Since both of these compounds are richer in strontium than SrZn₁₁, it can be assumed that an inversion of the effective charges of the components occurred, which led to a greater diffusion of strontium-containing substances towards tin and their decomposition in this area with the formation of new compounds. This assumption is also supported by the fact that in the system (Zn + 0.5 at.% Sr) –Sn, the SrSnₓ intermetallic compound was also synthesized under the moderating current.

Figure 3 presents the results of measuring the microhardness of the contact zone of the system (Zn + 0.5 at.% Sr) –Sn obtained in the absence of electric current.

Despite the relatively small variation in the values (maximum 40 MPa), the microhardness along this layer behaves rather unusual. The softest areas are near the boundaries, and if this is understandable in the case of stannic, the low values near the boundary with the solid solution of strontium in zinc are surprising. In the middle part, the microhardness generally gradually decreases with distance from (Zn + 0.5 at.% Sr), although there are also deviations here. This observation can be explained only by the concentration of zinc (for some reason) in the middle part of the layer.

The microhardness of this alloy with a distance from the boundary with a solid solution of strontium in zinc monotonously decreases, except for the region in the middle of the diffusion zone [8,9]. Apparently, in the currentless mode, the distribution of components in the interlayer, despite the presence of an intermetallic compound, is close to that dictated by their state diagram [10]. In comparison with the system (Zn + 0.5 at.% Sr) –Sn, the region near the solid solution is harder, and the region near stannic is softer.

**IV. CONCLUSION**

Studies have shown that contact melting of indium and stannic with a solid solution of strontium in zinc produces intermetallic compounds, some of which are absent in
equilibrium state diagrams. Electric current can influence their formation, and its direction is also important. A small difference in the content of strontium in the initial solution nevertheless has a noticeable effect on the resulting alloy. This can be seen in the flow of microhardness along the interlayers, which undergoes considerable leaps, and in different directions.

**Acknowledgment**

The study was carried out with the financial support of the Russian Foundation for Basic Research in the framework of research project No. 16-32-00666 mol_a.

**References**


