THERMAL STABILITY OF Cu$_{60}$Zr$_{30}$Ti$_{10}$ at. % BULK METALLIC GLASS

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Abstract

Cu$_{60}$Zr$_{30}$Ti$_{10}$ at. % metallic glass was prepared in two forms: as an ingot of diameter 3 mm and length of 26 mm and as a thin ribbon wide 5 mm and thick 35 μm. Chemical composition of the alloys were ascertain by Scanning Electron Microscopy (SEM) coupled with EDX analysis. Phase composition was evaluated by X-ray diffraction (laboratory and high-energy XRD at synchrotron). Thermal stability of ingot and ribbon samples were investigated by Differential Scanning Calorimetry (DSC) – where thermal characteristics such as glass transition temperature, crystallisation temperature and subsequent supercooled liquid region were compared for the two materials. Based on performed in-situ high temperature X-ray diffraction experiment we were able to determine changes in phases composition of the Cu$_{60}$Zr$_{30}$Ti$_{10}$ at. % amorphous alloys upon an annealing.

Keywords: bulk metallic glasses (BMG), Cu$_{60}$Zr$_{30}$Ti$_{10}$, suction casting, melt spinning, thermal stability

1. INTRODUCTION

Metallic glasses exhibit superior properties (high hardness and strength values, excellent corrosion resistance, large elongation limit, good combination of magnetic properties, etc.) which renders them unique for engineering and high-tech applications. Among metallic glasses, there is a group of noncrystalline solids called Bulk Metallic Glasses (BMG), which can be obtained by continuous cooling from the liquid state with section thickness of at least 1 mm (nowadays there’s a tendency among researchers to designate a glass as BMG with the requirement of the minimum diameter value to be 10 mm) [1]. To form a BMG, an alloy must fulfill 3 empirical rules formulated by Inoue [2]. Several exceptions to these empirical rules have been noted and one of these exceptions is connected to the ternary system of Cu-Zr-Ti. Cu – based glassy alloys have been synthesised in recent years in various forms (ribbons, bulk rods) and maximum diameter of the Cu – based BMG alloys was reported to be 12 mm in Cu$_{42}$Zr$_{42}$Ag$_{16}$ alloy [5]. Great ductility, which is uncommon for the vast majority of metallic glasses, predetermines Cu-based glasses for applications where deformation can be converted into measurable units such as fluid flow rate or electric signals and in this way they can be applied for different kinds of construction materials or sensors [6]. Cu – based systems (Cu-Zr-Ti, Cu-Hf-Ti, Cu-Zr-Ti-Hf) can be produced in a fully glassy state with a critical thickness of 4 mm. They can be used for structural applications - they have excellent mechanical properties with a tensile strength of about 2100 MPa, compressive plastic strain of 0,8 - 1,7 %, compressive fracture strength of 2200 MPa (Cu-Zr-Al) and a plastic strain of 0,2 % [7]. Das et al. has reported work – hardenable ductile bulk metallic glass of the composition of Cu$_{47.5}$Zr$_{47.5}$Al$_{5}$, which exhibits hight strength of up to 2265 MPa together with extensive work hardening and a large ductility of 18% measured by uniaxial compression tests. This remarkable ductility is attributed to special microstructural features at the atomic scale, which enable easy and homogeneous nucleation of the shear bands and continuous multiplication during deformation [8]. Ternary system of the composition of Cu$_{60}$Zr$_{30}$Ti$_{10}$ at.% belongs to the family of Cu – based bulk glassy alloys formed by Inoue et al. and is one of the exceptions to the Inoue empirical rules for high glass-forming ability (nanocrystalline microstructure of the BMG). This bulk metallic glass exhibits a large compressive plastic
strain of 1.6% with a high tensile and compressive strength of 2000 MPa. The supercooled liquid region was reported to be 37 °C. Since the reduced glass transition temperature (T_g = T_f / T_g) is relatively high (it was reported to be 0.62) and good glass forming ability is excepted, nanocrystalline microstructures with a size of 3 – 5 nm were always observed from the as-quenched state. Park et al. reported single phase amorphous state of the Cu60Zr30Ti10 ribbon. These observations suggest that the crystallisation occurs when cooling rate is not fast enough to suppress the crystallisation and relatively high plastic strain observed from the as-cast bulk Cu-Zr-Ti alloy is probably due to in-situ formed nanocrystals [10].

The main aim of this paper is to present the results of our own work on the ternary alloy of the composition of Cu60Zr30Ti10 at.%. Equipment of the Laboratory of Progressive Alloys of Materials and the Laboratory of Thermal Analysis of Institute of Materials Research allowed us to pass through the entire process - from the preparation of the material to the characterization with focus on thermal stability of the melt-spun ribbon and as-quenched ingot.

2. EXPERIMENTAL PROCEDURE

Cu60Zr30Ti10 alloy was prepared by arc-melting of the pure elements under a zirconium – getter argon atmosphere. The alloys were remelted several times due to homogeneity (problem of evaporation was neglected because of low vapour pressure of the elements) and finally cylindrical rods of the diameter of 3 mm and the length of 26 mm were prepared in suction casting facility attached to the arc melter. There were several efforts until the ingot was quenched because of choosing the appropriate suction parameters like temperature of the water – cooled copper crucible, argon overpressure and optimal weight of the sample. The amorphous ribbon of the thickness of 35 μm was prepared by melt-spinning technique under an argon atmosphere. Chemical composition of both samples was verified by SEM coupled with EDX microanalyser and the quantitative analysis proved good agreement with specified composition. The amorphous nature of the specimens was verified by X-Ray diffraction using Philips X’Pert Pro diffractometer with Cu Ka radiation. In order to investigate phase composition of the material upon an annealing, in-situ hard XRD measurement was carried out in transmission (Debye–Scherrer) geometry at the BW5 wiggler beamline [15]. Monochromatic synchrotron radiation of the energy of 100 keV (λ = 0.0124 nm) was used. The beam of photons illuminating sample had cross-section of 0.5 x 0.5 mm, diffracted X-rays were recorded by a 2D image plate detector Perkin Elmer XRD 1621 (2048 pixels x 2048 pixels, size of a pixel: 200μm x 200μm). In this experiment the Cu60Zr30Ti10 amorphous ribbon was cut into small pieces and put into a quartz capillary of the inner diameter of 0.8 mm. The capillary was mounted into a Linkam THMS600 resistive heater furnace with a maximal operating temperature of 600 °C. The temperature in the sample’s vicinity was measured by a thermocouple. The samples were heated up from the room temperature up to their crystallisation point at a heating rate of 10 °C/min. After reaching the required temperature, the glasses were illuminated by the X-ray beam for 20 s and 2D XRD patterns were recorded by the detector. Additionally, a time of about 15 s was needed for data transfer from the detector to a computer, to relax the detector and to stabilise the temperature. The XRD intensity was radially integrated to 2θ by using the Fit2D software. From the diffraction patterns we identified phases using the CMPR [16] toolkit. Thermal stability and characteristic temperatures were examined also by differential scanning calorimetry (Perkin Elmer DSC 8500) at a heating rate of 10 °C/min in an argon atmosphere. The compositional analysis was performed by energy – dispersive X-ray spectroscopy analysis installed in a field emission scanning electron microscope (Jeol JSM – 7000 F). Mechanical (compressive) tests has been realized on cylindrical specimen of the diameter of 3 mm and the length of 6 mm at room temperature using 200 kN Zwick – Extensometer.
3. RESULTS

As it is evident from Fig. 1 the X-ray diffraction patterns of the ingot (a) as well as the ribbon (b) exhibit broad diffraction maxima – characteristic feature for the glassy structures. XRD measurement indicating that the Cu_{60}Zr_{30}Ti_{10} alloy can be quenched as a single amorphous phase material with the diameter of 3 mm and more.

![Fig. 1 X-Ray diffraction pattern obtained from the ingot (a) and the ribbon (b) samples. Measurements were carried out on diffractometer Philips X’Pert Pro with Cu-K radiation, position sensitive detector and the scan step of size 0.03342 degree](image)

Fig. 1

![Fig. 2 DSC curves normalised to sample weight for the as-cast ingot (red) and the as-spun ribbon (blue) measured at the heat rate of 10 K/min show clear endothermic glass transition and exothermic crystallisation peaks](image)

Fig. 2

DSC curves normalised to sample weight for the as-cast ingot (red) and the as-spun ribbon (blue) measured at the heat rate of 10 K/min show clear endothermic glass transition and exothermic crystallisation peaks.

Fig. 2 shows DSC curves of the as-cast rod and from the melt-spun ribbon obtained upon continuous heating with a rate of 10 °C/min, all performed in an argon protective atmosphere. Both the curves exhibit a distinct glass transition, followed by a supercooled liquid region and strong exothermic reactions caused by crystallisations. The values of glass transition temperatures and crystallisation temperatures slightly differ as can be seen on Fig. 2 and in the Table 1. Supercooled liquid region increased from the value of 32 °C for as-cast ingot to the value of 36 °C for as-spun ribbon. For the as-spun ribbon there are three exothermic peaks, the first with maxima at 489 °C (enthalpy of the exothermic reaction is 17.79 J/g), the second – twin peak has a maxima at 535 °C (enthalpy is 31.89 J/g) and the third one has a maxima at the position of 690 °C with the enthalpy of 3.79 J/g. In the case of as-cast ingot, the first exothermic peak has a maxima at the...
temperature of 481 °C (enthalpy is 10.49 J/g) followed by a twin peak with the maxima at 530 °C (enthalpy 13.81 J/g) and the third crystallisation peak with the maxima at 645 °C and the enthalpy of 3.76 J/g.

**Table 1** Values of significant temperatures of the Cu_{60}Zr_{30}Ti_{10} ribbon and the ingot obtained from DSC curves at a heating rate of 10 °C/min in an argon atmosphere

<table>
<thead>
<tr>
<th>sample</th>
<th>T_g [°C]</th>
<th>T_{x1 onset} [°C]</th>
<th>T_{x1} [°C]</th>
<th>Enthalpy [J/g]</th>
<th>T_{x2} [°C]</th>
<th>Enthalpy [J/g]</th>
<th>T_{x3} [°C]</th>
<th>Enthalpy [J/g]</th>
<th>ΔT_x = T_x - T_g [°C]</th>
</tr>
</thead>
<tbody>
<tr>
<td>ribbon</td>
<td>443</td>
<td>479</td>
<td>489</td>
<td>17.79</td>
<td>535</td>
<td>31.89</td>
<td>690</td>
<td>3.79</td>
<td>36</td>
</tr>
<tr>
<td>ingot</td>
<td>435</td>
<td>467</td>
<td>481</td>
<td>10.49</td>
<td>530</td>
<td>13.81</td>
<td>645</td>
<td>3.76</td>
<td>32</td>
</tr>
</tbody>
</table>

The ribbon seems to be more glassy than the ingot because of higher value of the magnitude of heat produced by the exothermic transformations – it’s known that sample having a higher value of the magnitude of heat has more volume fraction of the amorphous phase. Uneven widths of the supercooled liquid regions and different values of the significant temperatures are presumabably features of unidentical glassy structure after casting / spinning of the rod and the ribbon. The observed differences indicate that structure of ingots contains some amount of nanocrystals (not detected by the XRD) acting as nucleation centers facilitating crystallisation of this alloy. This assumption need to by verified by transmission electron microscopy or by the other similar unambiguous technique.

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**Fig. 3** XRD patterns of the ribbon with phase evolution for various temperatures: in the glassy state, after 1st and after 2nd crystallisation

Fig. 3 shows XRD patterns taken from the ribbon samples at different temperatures. In the picture we can see 4 distinct curves belonging to 4 various temperatures – temperature at which the ribbon is still in the glassy state followed by 3 temperatures after 1st and 2nd crystallisation peaks. At the temperature of 150 °C, the ribbon is preserving the glassy structure manifested by the broad and diffuse peaks. Once the temperature increases and reaches the value of 514 °C, we can see distinct change of its character. We identified products of the glassy phase transformation as hexagonal Cu_{135}Zr_{0.82}Ti_{0.825} and two face centered cubic phases having significantly different lattice parameters - Cu_{0.92}Ti_{0.08} and a new – CuZr phases. As the temperature continuous to increase, above the second crystallisation peak 553 °C, hexagonal
Cu$_{1.35}$Zr$_{0.825}$Ti$_{0.825}$ phase begins to dissolve and transforms into the two fcc phases - Cu$_{0.92}$Ti$_{0.08}$ and a new - CuZr phase which. Space groups and lattice parameters of all the identified phases are listed in Table 2.

**Table 2** Lattice parameters of the identified phases in Cu$_{60}$Zr$_{30}$Ti$_{10}$ at. % ribbon

<table>
<thead>
<tr>
<th>Formula</th>
<th>Space Group</th>
<th>a [Å]</th>
<th>c [Å]</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cu$<em>{1.35}$Zr$</em>{0.825}$Ti$_{0.825}$</td>
<td>P63/mmc (194)</td>
<td>5.1775</td>
<td>8.3673</td>
<td>[18]</td>
</tr>
<tr>
<td>Cu$<em>{0.92}$Ti$</em>{0.08}$</td>
<td>Fm-3m (225)</td>
<td>5.08</td>
<td>-</td>
<td>[19]</td>
</tr>
<tr>
<td>CuZr</td>
<td>Fm-3m (225)</td>
<td>3.65</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

The room temperature compressive stress-strain curve obtained from ingot of nominal composition of Cu$_{60}$Zr$_{30}$Ti$_{10}$ at. % is shown in Fig. 4. The ingot exhibits an elastic regime of 7.5 % before yielding, which occurs at about 1640 MPa stress. After yielding the stress increases with increasing strain up to the fracture, which takes place at 1790 MPa stress and a strain of 8.4 %. This results in a plastic strain of 0.9 %.

![Room temperature compressive stress-strain curve](image)

**Fig. 4** Room temperature compressive stress – strain curve for the ingot of nominal composition of Cu$_{60}$Zr$_{30}$Ti$_{10}$ with aspect ratio of 2 (6mm length and 3 mm diameter) prepared from the cast rod

CONCLUSION

In this work thermal stability, phase analysis and compression characteristics of Cu$_{60}$Zr$_{30}$Ti$_{10}$ at. % BMG were examined. Thermal parameters of the ingot and the ribbon were evaluated and compared. Characteristic temperatures differ manifesting unequal glassy structure of the samples. Phase evolution of the ribbon was described in relation to distinct temperatures at which the sample was heated. Hexagonal Cu$_{1.35}$Zr$_{0.825}$Ti$_{0.825}$ and two cubic - Cu$_{0.92}$Ti$_{0.08}$ and new - CuZr phases with enhanced lattice parameters were identified. The yield strength, the elastic and plastic deformation and the compression fracture strength were evaluated.

ACKNOWLEDGEMENTS

REFERENCES


