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THE STRUCTURE OF AMORPHOUS ALLOYS SYNTHESIZED BY MECHANICAL ALLOYING-
NON GLASS FORMING SYSTEMS

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Abstract - Preliminary results are reported of the amorphization of Nb-Sn and
Nb-Ge alloys by mechanical alloying of the elemental powders. Nb-25
at.% Sn, Nb-25 at.% Ge, and Nb-27 at.% Ge alloys were made amorphous. Dif­
ferential scanning calorimetry (DSC) indicated a crystallization peak at
995 K for Nb-25 at.% Sn but > 1000 K for the Nb-Ge amorphous alloys. A
large endothermic DSC peak and the diffraction patterns of the as-mechani­
cally alloyed vs. heated Nb-27 at.% Ge amorphous alloy suggest a large
change in the structure of the amorphous phase.

I - INTRODUCTION

Amorphous alloys have been prepared by a variety of methods. These methods may be
classified according to the nature of the precursor to the amorphous structure -
e.g. synthesis from the liquid state, vapor state, or aqueous solution. Synthesis
of the amorphous structure from the crystalline solid has been given increasing
attention in recent years. Techniques in this regard include ion implantation /1/,
ion bombardment /2/, neutron-irradiation /3/, ion mixing of metallic bilayers /4/,
and solid-state interdiffusion /5/. Koch et al /6/ have demonstrated that mechani­
cally alloying of elemental Ni and Nb powders by high energy ball milling for extend­
ted times leads to the formation of an amorphous alloy powder. Mechanical alloy­
ing (MA) is a high-energy ball milling technique for producing composite metal
powders with controlled microstructures by the repeated cold welding and fracture
of powder particles. If MA is continued to the point where the starting powders
are mixed to dimensions of a few atomic spacings, then an "alloy" of the elemen­
tal powders can be formed. The composition range over which the amorphous structure
can be attained has been extended by MA over that produced by rapid solidifi­
cation in both the Ni-Nb system (P. Y. Lee and C. C. Koch, unpublished research,
North Carolina State University) and the Ni-Ti system (R. B. Schwarz and coworkers,
Argonne National Laboratory, private communication).

Both of the above alloy systems are "easy glass formers". That is, the amorphous
structure can be produced by "conventional" rapid solidification processing (quench
rate - 106 K/s) over fairly wide ranges of composition. While additional studies
of such glass forming systems prepared by MA are being conducted in our labora­tory
we are also studying the feasibility of producing the amorphous structure by MA
in non glass forming systems. The two systems picked for this study are Nb-Sn and Nb-Ge. Alloys with the compositions Nb3Sn and Nb3Ge have been made amorphous by vapor quenching methods, e.g. /7/. However, no amorphous phases have been produced in these systems by rapid solidification methods at usual quenching rates (10^6 K/s) - that is, they are non glass forming.

This paper presents the results of preliminary studies of the feasibility of synthesizing the amorphous structure by MA of Nb-Sn and Nb-Ge powders. The stability of the amorphous phases so formed was also examined.

II - EXPERIMENTAL

The MA was carried out on pure elemental powders of Nb (Alfa Products, 99.8% - 325 mesh), Sn (Alfa Products, 99.5%, - 325 mesh), and Ge (Alfa Products, 99.999%, - 325 mesh). Mechanical alloying was performed in a Spex Mixer/Mill Model 8000. Martensitic stainless steel balls (AISI 440C), which were 7.9 mm in diameter, were used in the cylindrical hardened tool steel vial (76 x 57 mm diameter). The elemental powders were loaded in the vial under air with a ball to powder weight ratio of 6:1. In some cases the vial was opened after a given time and a small sample of powder was removed for x-ray diffraction analysis. In other cases MA was continued to a final processing time before the vial was opened. Recently several experiments were conducted with a slit in the vial cap to allow for the insertion of a thermocouple. This provided a continuous source of air (oxygen) to the powders. The oxygen that is certainly incorporated in the MA powder has not yet been analyzed for. However, based on previous results on MA of Mn60 Nb40 in air /6/, it is expected that at least 3 wt.% oxygen may be contained in the present powders.

Three conditions were used in terms of the average temperature of the vial during MA. If no attempt were made to remove the heat generated by the kinetic energy of the balls etc., the vial temperature, as measured by either external or internal thermocouples, would reach approximately 65°C after extended periods. Forced convection for heat removal with a fan lowered the average temperature to about 35°C. In order to obtain additional cooling a flow of liquid nitrogen was impinged upon the vial during MA. The average temperature attained was not as reproducible as the above conditions due to difficulties in controlling the liquid nitrogen flow on the rapidly moving vial. Temperatures of -150°C ± 5°C were observed.

The structures of the MA powders were followed by x-ray diffraction measurements using cylindrical samples (powder uniformly coated on a glass fiber) in a 57.3 mm diameter Debye-Scherrer camera. The x-ray radiation used was CuKα (λ = 0.154 nm). Several samples were also examined in a Philips APD 3600-02 automated powder diffractometer.

The powder morphology as a function of MA time was followed by optical and scanning electron microscopy. Microhardness measurements were made on metallographically mounted powders using a Buehler Micromet microhardness tester.

Initial studies of the amorphous phase stability were carried out by heating the amorphous powders in a Perkin-Elmer differential scanning calorimeter (DSC II) at 20 K/min.

III - RESULTS

A. Amorphous Phase Formation

1. Nb-Sn System

The only alloy studied to date in the Nb-Sn system is Nb-25 at.% Sn. It was necessary to cool the vial during MA to prevent excessive cold welding of the Sn powders to the vial wall and the balls. Because of this experimental inconvenience more compositions have been examined in the Nb-Ge system.
The progress of structural changes during MA was observed in Nb-25 at.% Sn by x-ray diffraction measurements taken as a function of time of MA. The diffraction patterns changed with MA time in the following manner: first the Nb and Sn lines broadened; then the Sn lines could not be resolved; the peak position of the (110) Nb line moved to lower angles and the line width increased; finally, a diffraction pattern typical of the amorphous structure developed. There is an almost linear decrease in (110) peak $2\theta$ up to about 14 hours of MA, (at $-15^\circ$C) after which time $2\theta$ stays constant and the amorphous diffraction pattern develops. Similar results were obtained previously for the Ni$_{60}$Nb$_{40}$ alloy /6/. The changes in $2\theta$ must reflect the "alloying" stage of the process as the dimensions of the component powders are reduced to the atomic level. After "alloying" is complete the transformation to the amorphous structure occurs.

The time at which the amorphous structure forms varied with temperature as is indicated in Table 1. The time for attainment of the amorphous phase appears to go through a maximum at 35°C and MA at $-15 \pm 5^\circ$C gives the shortest time. The same temperature dependence has also been observed for amorphization time in the Nb-Ge alloys to be discussed below.

<table>
<thead>
<tr>
<th>Alloy Composition</th>
<th>Phase Formed by M.A.</th>
<th>Time for Amorphization by M.A. (hours)</th>
<th>Ave. Temp.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nb-25 at.% Sn</td>
<td>Amorphous</td>
<td>18.5</td>
<td>-15 ± 5°C</td>
</tr>
<tr>
<td></td>
<td></td>
<td>27</td>
<td>35°C</td>
</tr>
<tr>
<td></td>
<td></td>
<td>23</td>
<td>65°C</td>
</tr>
<tr>
<td>Nb-25 at.% Ge</td>
<td>Amorphous</td>
<td>10</td>
<td>-15 ± 5°C</td>
</tr>
<tr>
<td></td>
<td></td>
<td>11</td>
<td>65°C</td>
</tr>
<tr>
<td>Nb-27 at.% Ge</td>
<td>Amorphous</td>
<td>7</td>
<td>-15 ± 5°C</td>
</tr>
<tr>
<td></td>
<td></td>
<td>18</td>
<td>35°C</td>
</tr>
<tr>
<td></td>
<td></td>
<td>10</td>
<td>65°C</td>
</tr>
<tr>
<td>Nb-38 at.% Ge</td>
<td>NbGe$_3$</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Nb-59 at.% Ge</td>
<td>NbGe$_2$</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Nb-67 at.% Ge</td>
<td>NbGe$_2$</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

2. Nb-Ge System

Due to the brittle mechanical behavior of Ge, excessive cold welding was not a problem in MA of the Nb-Ge alloys. In all cases the starting material consisted of the elemental Nb and Ge powder. Of the five compositions studied to date, only the 25 at.% Ge and 27 at.% Ge alloys could be made amorphous by MA (see Table 1). The 38, 59, and 67 at.% Ge compositions formed the structures, listed in Table 1, which are consistent with the phases found in the equilibrium diagram at those compositions.

B. Stability of the Amorphous Structure

Preliminary measurements of the stability of the amorphous phases in the Nb-25 at.% Sn, Nb-25 at.% Ge, and Nb-27 at.% Ge alloys were made by differential scanning calorimetry (DSC).

On heating in the DSC at 20 K/min the Nb-25 at.% Sn sample exhibited a large endothermic peak which started at 540 K, peaked at 660 K, then merged into the
exothermic crystallization curve at about 920 K. The exothermic peak occurred at 995 K, almost at the temperature limit of the DSC II. X-ray diffraction of the powder after the DSC run indicated the presence of the Nb₆Sn₅ and NbSn₂ phases.

Similar DSC experiments on the Nb-25 at.% Ge and Nb-27 at.% Ge amorphous powders also revealed broad endothermic peaks which started at about 600 K and peaked at about 780 K (25 at.% Ge) or about 860 K (27 at.% Ge). While the DSC traces of both these alloys became exothermic at temperatures above 900 K (920 K for 25 at.% Ge, 960 K for 27 at.% Ge) no exothermic peak could be obtained up to the temperature limit of the DSC (1000 K).

X-ray diffraction patterns for the amorphous Nb-27 at.% Ge powder in the as-MA condition and after heating in the DSC at 20 K/min to 1000 K, (and subsequent cooling to room temperature) are illustrated in Figures 1b and 1a respectively. Figure 1b shows a diffraction pattern which is typical of those we have observed for "amorphous" MA powders. The intensities of the first and second peaks are considerably lower than usually observed for liquid-quenched amorphous alloys. After the temperature excursion to 1000 K in the DSC the diffraction pattern (Figure 1a) more closely resembles that for a liquid-quenched alloy. That is, the intensities of both the first and second peaks have increased and the second peak shows the characteristic splitting. Very small Bragg peaks can be seen above the amorphous pattern as evidence of the start of crystallization, which is consistent with the beginnings of the exothermic reaction observed in the DSC experiment.

![Graph](image)

Figure 1 (a) Diffraction pattern of Nb-27 at.% Ge amorphous alloy produced by MA heated to 1000 K at 20 K/min in DSC. (b) Diffraction pattern of Nb-27 at.% Ge amorphous alloy as-mechanically alloyed.
IV. DISCUSSION

While the preliminary results presented above are not extensive enough to draw general conclusions regarding the amorphization tendency by MA for the Nb-Sn and Nb-Ge systems, some comments on possible mechanisms are in order. Crystalline solids have been amorphized by ion, neutron, and electron irradiation (e.g. /2/, /3/, /8/ respectively). A mechanism first proposed by Swanson et al /3/ has been used to explain amorphization by irradiation of crystalline solids. It involves the concept of a critical defect concentration introduced by irradiation such that spontaneous transformation to the amorphous state will occur when the free energy of the crystalline phase (GC) plus the free energy increase due to defects produced by irradiation (GD) is greater than the free energy of the amorphous phase (GA): that is, GC + GD > GA. A critical defect density may be the controlling mechanism for amorphization by mechanical alloying.

The mechanism presented for amorphization by solid state diffusion involves systems with large negative heats of mixing in the liquid (or amorphous) alloys and anomalously fast diffusion /5/. If solid state diffusion were the controlling mechanism in the amorphization of Nb-Sn and Nb-Ge alloys by MA then it might be expected that the time of MA for amorphization would decrease with increasing temperature. Our preliminary results do not indicate such a trend. Clearly more work is needed to determine the mechanism(s) responsible for amorphization by MA as a function of composition.

The above preliminary results on the occurrence and structure of the MA amorphous Nb-Sn and Nb-Ge alloys have provided several important questions to be resolved by further research.

ACKNOWLEDGEMENTS

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