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THE ELASTIC CONSTANTS OF MgAg AND MgCu₂ SINGLE CRYSTALS

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Abstract—The three independent elastic constants of MgAg and MgCu₂ have been determined by the pulse–echo method from 80° to 500°K. The temperature coefficients of all the elastic constants are normal but smaller than the average of their constituent elements. The anisotropy factor is high for MgAg and low for MgCu₂, as one would expect from their respective crystallographic configurations. The elastic behavior of MgCu₂ was found to be similar to that of the diamond structure.

INTRODUCTION

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SINCE many intermetallic compounds of many different crystal structures have been found in alloy systems, the basic reason for their stability has drawn a great deal of attention. However, measurements on elastic constants of these compounds, which may enable one to evaluate the extent of interatomic forces, have been relatively few. It was the intention of the present investigation to determine the elastic constants of two compounds, MgAg and MgCu₂, so that a better understanding about the nature of the interaction between neighboring ions in the crystals may be achieved.

MgAg has a CsCl-type structure. The elastic properties of this structure have been the most explored among all intermetallic compounds, both theoretically and experimentally. A metallic compound of this type is characterized by a high degree of anisotropy due to its low resistance to a (110) [110] shear. MgCu₂ is the proto-type cubic Laves phase. It is believed that its stability can be attributed to the fact that it is a close-packed arrangement of unequal spheres. If this interpretation is correct, we may expect that MgCu₂ will be fairly isotropic. Both compounds are stable and remain ordered at all temperatures up to the melting point.

EXPERIMENTAL PROCEDURE AND RESULTS

Single crystals of MgAg and MgCu₂ of stoichiometric compositions, received from DR. V. B. Kurfman of our laboratory, were grown by the Bridgman method in graphite molds from high purity metals. For each compound, two cylindrical specimens were cut in the [100] and [110] orientations with an electrical discharge machine. The surface finishing was done on metallographic polishing wheels without final etching. The dimensions of each specimen were about 3 cm dia. $\times 1$ cm thick, with misorientation $<2^{\circ}$ and parrallelism within ± 0.005 mm.

Sonic velocities were determined by the pulseecho technique using a 10 Mc/s rectified pulse of $1.5 \,\mu$ sec duration. The experimental set-up was similar to that of HUNTINGTON.⁽¹⁾ The basic components of the electrical system consisted of an Arenberg PG650C pulsed oscillator, a balancing network, and a WA600C wide-band amplifier, in conjunction with a Tektronix 545A oscilloscope. A 4× magnifying lens mounted in front of the cathode-ray tube screen helped to improve visibility of the echoes. The arrangement of the specimen holder is shown in Fig. 1. The temperatures from 80° to 500°K were measured with a copper-constant thermocouple embedded in the base of the specimen seat. To bond the X-cut or Y-cut quartz transducer to the specimen, Nonaq was used for low temperatures and Dow Corning 806A silicone resin for high temperatures. For the



FIG. 1. The sample holder.

latter, a thin layer of the resin was first applied to the top surface of the specimen and to the bottom surface of the transducer. These layers were cured separately for an hour at 100°C, then were matched together and baked at 250°C under spring loading for another hour. For MgCu₂ specimens, numerous echoes, ~100, were visible, but attenuation in MgAg was high, with only 8 or 9 echoes displayed.

The room temperature densities were taken as 6.042 g/cm^3 for MgAg and 5.76 g/cm^3 for MgCu₂.⁽²⁾ The linear coefficients of thermal expansion were determined experimentally. The average values between 0° and 250°C were found to be 22.8×10^{-6} (MgAg) and 18.7×10^{-6} cm/cm/°C (MgCu₂). The latter value is significantly smaller

than the previously reported one.⁽³⁾ For each compound, five velocities were measured and the elastic constants were computed with two internal checks. Transit-time corrections of $0.025 \,\mu\text{sec}$ and $0.04 \,\mu\text{sec}$ were subtracted from the observed delay times in calculating the longitudinal and transverse velocities respectively.

The adiabatic elastic constants of MgAg and MgCu₂ from 80° to 500°K are shown in Figs. 2 and 3, and listed in Tables 1 and 2. The estimated over-all accuracy is ± 1 percent for C_{11} or C_{44} , and ± 2 percent for C_{12} . They follow the normal



FIG. 2. The temperature dependence of elastic constants of MgAg.

features of temperature dependence with an approach to zero slope towards 0°K, and a negative slope at higher temperatures. The temperature coefficients are smaller than the average of the constituent elements. The low temperature data extrapolated to 0°K give the Debye temperatures at $315^{\circ} \pm 3^{\circ}$ K (MgAg) and $380^{\circ} \pm 4^{\circ}$ K (MgCu₂) by using BLACKMAN's formula.⁽⁴⁾

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DISCUSSION

As one can see from Fig. 2, with decreasing temperature MgAg exhibits a strong tendency to increase its anisotropy factor, C/C'. [Here $C = C_{44}$ and $C' = (C_{11} - C_{12})/2$.] This trend is even