An assessment of structural enthalpy and crystallization pathways of Mg₆₅Zn₃₀Ca₅ bulk metallic glass and amorphous films

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ABSTRACT

The structural nature and thermal stability of amorphous alloys is highly dependent on the method by which they are produced, i.e. their relaxation rate upon cooling. Both bulk samples and metallic glass films of $Mg_{65}Zn_{30}Ca_5$ were produced by copper mold casting and direct current (DC) magnetron sputtering onto aluminium substrates, respectively. Comparisons between structural enthalpy, crystallization pathways, relaxation and crystallization kinetics of the bulk samples and films were examined by elevated temperature XRD and DSC. Compared with equivalent experiments on the bulk alloy, results for the thin films show distinct differences in structural enthalpy and deviations from the expected crystalline phase evolution, displaying minor peak shifts, failure of some phases to evolve, and variations in the evolution rates.

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1 INTRODUCTION

The structural nature and thermal stability of amorphous alloys is highly dependent on the method by which they are produced, i.e. their relaxation rate upon cooling. Both bulk samples and metallic glass films of Mg₆₅Zn₃₀Ca₅ were produced by copper mold casting and direct current (DC) magnetron sputtering onto aluminium substrates, respectively. Comparisons between structural enthalpy, crystallization pathways, relaxation and crystallization kinetics of the bulk samples and films were examined by elevated temperature XRD and DSC. Compared with equivalent experiments on the bulk alloy, results for the thin films show distinct differences in structural enthalpy and deviations from the expected crystalline phase evolution, displaying minor peak shifts, failure of some phases to evolve, and variations in the evolution rates.

Key sources [1, 2] [3]

2 METHOD

2.1 Master alloy

The master alloy of Mg₆₅Zn₃₀Ca₅ was produced using high-purity elements of Mg (99.85 wt%), Zn (99.995 wt%), and Ca (99.8 wt%). The alloy was prepared by induction melting in boron nitride coated graphite crucibles, purged with Ar (99.997 vol.% purity) five times, and protected with a circulating Ar atmosphere. Alloy homogeneity was ensured by heating and cooling through a cycle of 700°C, 385°C, 650°C, 385°C, 650°C to a casting temperature of 500 °C and 450°C for for injection and gravity casting respectively. Bulk amorphous Mg₆₅Zn₃₀Ca₅ rod of 2.5mm diameter and plates of thickness of XXµm were produced by copper mold injection casting. The 25.4mm diameter targets were prepared from a cylindrical copper mold gravity castings sectioned to thicknesses of 3.25mm. All samples and targets were stored under Ar (99.997 vol.% purity) when not being examined or used.

2.2 DC magnetron sputtering

Films were produced from an in-house DC magnetron sputtering facility with Ar working gas (99.997 vol.% purity). The power was 15W, typical voltage of 290-350V, nominal chamber pressure of 1 bar, substrate temperature of 25 °C, and Ar flow of 3.01 SCCM. Films

were deposited directly onto to Al DSC lid substrates. Depositions were for a period of 35 minutes. Deposition rate was estimated at 1.2nm/s.

2.3 DSC characterization

Isochronic DSC (204 F1 Phoenix, Netzsch, Selb, Germany) was carried out in aluminium crucibles under a protective Ar (99.997 vol.% purity) atmosphere. Scans were performed at heating rates of 5 to 100*K*/*min*.

For annealed XRD the samples were heat treated in the DSC by heating to the desired temperature at 20K/min followed by quenching to room temperature.

2.4 XRD characterization

Annealing XRD (PANalytical, Empyrean, Cu K_{α} X-ray source, $\lambda = 1.541\text{Å}$) was performed at room temperature. (Generator Voltage 45, Tube Current 40, Scan Step Size 0.0262606, Time per Step 397.29).

Dynamic XRD (Bruker, D8, Cu K_{α} X-ray source, $\lambda = 1.541\text{Å}$) was performed by raising temperature at a rate of 20K/min and performing scans in situ. The first scan was performed at 35 °C, then 75 °C, after which temperature was raised in 5K increments. The 2θ scans from $31-60^{\circ}$ were completed within 1092sec (18min, 12sec) to minimise the effects of recrystallisation during the experiment. (Generator Voltage 45, Tube Current 100, Scan Step Size 0.02, Time per Step 134.4).

3 RESULTS

deposition were for 35 minutes, and saw an average temperature rise of 3-4 °C during the deposition. Nominal film thickness was about $2.5\mu m$ giving a deposition rate of 1.2nm/s.

Relaxed differential scanning calorimetry (DSC) of bulk material taken at different heating rates was used to establish the fragility of the Mg₆₅Zn₃₀Ca₅ system. From the equations ... a fit of $\beta^{-1} = 1.338E - 16e^{5274(\frac{1}{T-T_0})}$ with Adj. $R^2 = 0.972$ was established. This give a $D^* = 20.4$ which using $D^* = 590/(m-16)$ Shuai2014 [4, 5] gives a fragility m = 44.9.

Heating Rate β <i>K/min</i>	T_g	T_{x1}	T_{x2}	T_{x3}	T_{x4}	T_{x5}
	107.1	150.4	102.4	201.0	240.2	262.4
100	136.1	152.4	193.4	201.8	240.2	262.4
80	132.0	151.6	194.4	201.9	238.2	260.3
60	129.6	151.6	190.0	197.8	232.9	259.0
40	126.6	151.0	189.0	200.0	226.4	254.7
30	126.2	147.4	187.0	198.4	221.0	251.1
20	125.1	147.3	188.4	197.0	216.0	246.8
15	123.8	146.0	186.2	195.6	212.2	243.9
10	123.5	142.3	183.4	192.9	207.4	239.8
5	120.5	139.2	179.7	187.5	199.8	232.7

Table 1: Bulk Mg₆₅Zn₃₀Ca₅ alloy onset temperatures for the various DSC heating rates β . All temperatures are in °C.

Heating Rate β <i>K/min</i>	T_g	T_{x1}	T_{x2}	T_{x3}	T_{x4}	T_{x5}
100	108.5	135.1	0.0	177.3	0.0	240.3
80	106.0	128.2	0.0	165.6	0.0	238.8
60	107.3	129.1	0.0	176.1	0.0	237.8
40	100.2	117.6	0.0	170.7	0.0	234.2
30	95.3	114.4	0.0	169.5	0.0	232.5
20	95.5	110.0	0.0	170.5	0.0	229.4
15	92.5	107.4	0.0	168.8	0.0	224.0

Table 2: Film Mg₆₅Zn₃₀Ca₅ alloy onset temperatures for the various DSC heating rates β. All temperatures are in °C.

SCOnsets

lkOnsets

4 DISCUSSION

The use of a 60K DSC heating rate compared to the more commonly used 20K rate [sources] shifts peaks for the bulk $Mg_{65}Zn_{30}Ca_5$ alloy about 8 - 15 degrees higher. This higher heating rates were used because crystallization events for the films were different to differentiation at the lower heating rate. Films show little shift to high temperature peaks with increases heating rates, but large shifts with relaxation. Bulk show the opposite behaviour, larger peaks shifts with higher heating rates and little shift with relaxation.

5 CONCLUSIONS

6 ACKNOWLEDGEMENTS

Yu Wang for his assistance with XRD experimentation and Rietveld refinement.

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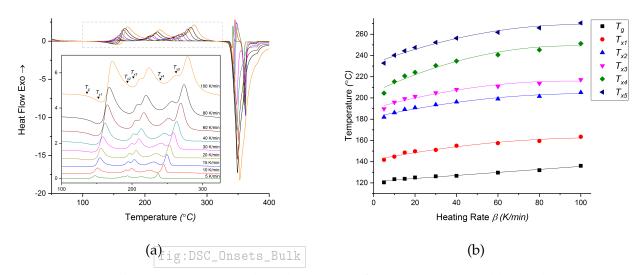


Figure 1: (a) Bulk $Mg_{65}Zn_{30}Ca_5$ relaxed at 120 °C for 10 minutes and heated at various heating rates. The insert stacks the DSC curves and labels the T_g and T_x es of the 100K/min sample. (b) The T_g s and T_x es of the bulk $Mg_{65}Zn_{30}Ca_5$ at all heating rates.

ate_Bulk

DSC_Bulk

ate_Film

DSC_Film

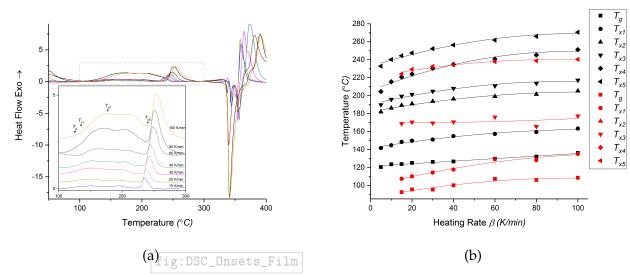


Figure 2: (a) Unrelaxed film $Mg_{65}Zn_{30}Ca_5$ heated at various heating rates. The insert stacks the DSC curves and labels the T_g and T_x es of the 100K/min sample. (b) The T_g s and T_x es of the bulk and film $Mg_{65}Zn_{30}Ca_5$ at all heating rates.

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Figure 3: Fitted fragility for the $Mg_{65}Zn_{30}Ca_5$ system obtained by DSC at various heating rates

 m_mValue

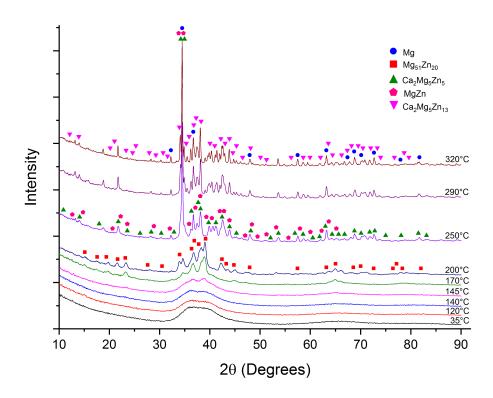


Figure 4: XRD pattern for Bulk Mg₆₅Zn₃₀Ca₅ heated through several crystallization peaks identified from DSC

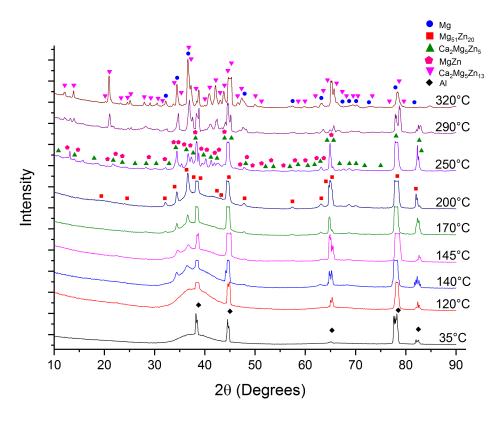
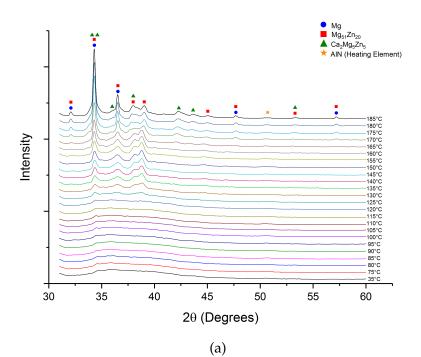


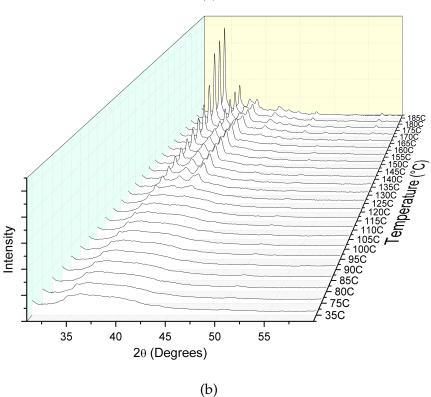
Figure 5: XRD pattern for Film $Mg_{65}Zn_{30}Ca_5$ heated through several crystallization peaks identified from DSC

ing_Film

ing_Bulk



amic_FullStack_Bulk



amic_WaterFall_Bulk

mic_Bulk

Figure 6: (a) Stacked X-ray diffraction (XRD) patterns from the incremental heating of bulk $Mg_{65}Zn_{30}Ca_5$. (b) Cascading XRD patterns from the incremental heating of bulk $Mg_{65}Zn_{30}Ca_5$.

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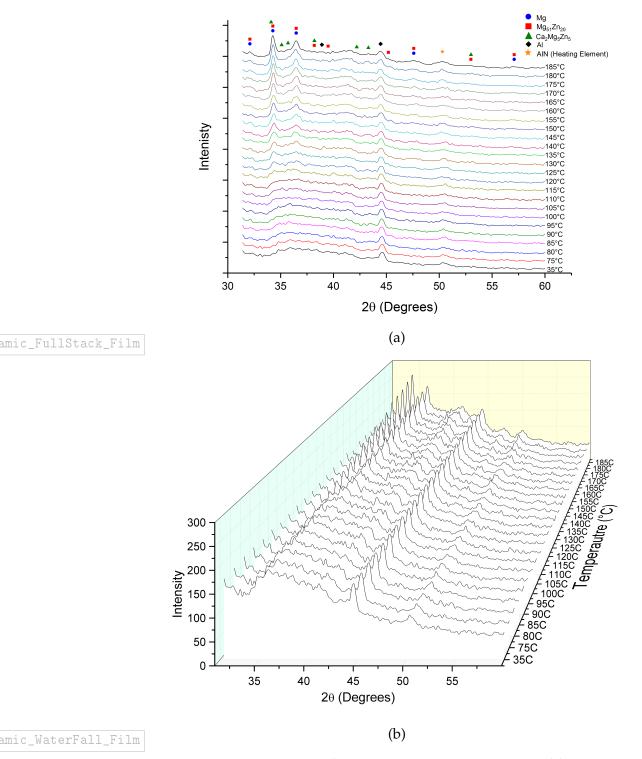


Figure 7: (a) Stacked XRD patterns from the incremental heating of film $Mg_{65}Zn_{30}Ca_5$. (b) Cascading XRD patterns from the incremental heating of film $Mg_{65}Zn_{30}Ca_5$.

mic_Film