

## Thermodynamic properties and microstructural characteristics of binary Ag-Sn alloys

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Received December 20, 2012; accepted January 6, 2013

The liquidus and solidus temperatures and enthalpy of fusion for Ag-Sn alloys are systematically measured within the whole composition range by differential scanning calorimetry (DSC). The measured enthalpy of fusion is related to Sn content by polynomial functions, which exhibit one maximum value at 52 wt%Sn and two minimum values around 21 wt%Sn and 96.5 wt%Sn, respectively. The liquidus slope, the solidification temperature interval, the solute partition coefficient and the entropy of fusion are calculated on the basis of the measured results. The undercoolability of those liquid Ag-Sn alloys solidifying with primary (Ag) solid solution phase is stronger than the other alloys with the preferential nucleation of  $\zeta$  and  $\varepsilon$  intermetallic compounds. Morphological observations reveal that peritectic reactions can rarely be completed, and the peritectic microstructures are always composed of both primary and peritectic phases.

### Ag-Sn alloy, enthalpy of fusion, undercoolability, peritectic solidification

**Citation:** Zhai W, Wei B B. Thermodynamic properties and microstructural characteristics of binary Ag-Sn alloys. *Chin Sci Bull*, 2013, 58: 938–944, doi: 10.1007/s11434-013-5695-4

In recent years, Ag-Sn alloys have aroused great scientific research interest because the Ag rich Ag-Sn alloys show excellent electronic conductivity and the Sn-rich Sn-Ag alloys are ideal substitutes for lead-free solders [1–5]. Many investigations reveal that the thermodynamic properties of liquid alloys and the final solid microstructures affect their performance [6–13]. However, there are few reports on these subjects for Ag-Sn alloys, and therefore, efforts should be made to the thermal properties and solidification microstructures of different Ag-Sn alloys in order to explore and understand their physical properties.

The present work focuses on the following three aspects. Firstly, the liquidus temperature and enthalpy of fusion for Ag-Sn alloys are fundamental thermodynamic parameters in determining other thermal properties. Although the enthalpy of fusion for binary alloys can be roughly estimated by Neumann-Kopp's rule from the values of the two pure components, this method usually brings in large discrepancy. Hence, the enthalpy of fusion for Ag-Sn alloys versus

composition should be measured experimentally. Secondly, the solidification microstructure is mainly undercooling-dependent, and hence, much work has been done to prevent the heterogeneous nucleation of liquid alloys by various denucleation techniques. One may neglect the fact that the undercoolability of liquid alloys also relies on the alloy composition, and different alloys may be undercooled by varying degrees even under the same external condition. From this point of view, it is essential to study the intrinsic undercoolability of binary Ag-Sn alloys with different compositions. Thirdly, for Ag-Sn alloys, there are two peritectic reactions and one eutectic transformation in the binary phase diagram. However, the growth morphologies of Ag-Sn peritectic and eutectic alloys are not well known and should be investigated.

The differential scanning calorimetry (DSC) is an efficient technique for quantitative thermal properties [14,15], which also provides the essential information on the liquid to solid phase transition characteristics [16]. In present work, the enthalpy of fusion for Ag-Sn alloys within the entire composition range is measured by DSC method. The un-

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dercoolability of liquid Ag-Sn alloys with different compositions is also studied. Meanwhile, the thermal characteristics and the final solidification microstructures of typical peritectic and eutectic Ag-Sn alloys are investigated on the basis of DSC calorimetric analyses.

## 1 Experimental procedure

Nineteen Ag-Sn alloys with different compositions were investigated, which are listed in Table 1 and illustrated in the binary Ag-Sn phase diagram [17] shown in Figure 1. Each sample had a mass of about 150 mg and was prepared

**Table 1** Thermodynamic properties of Ag-Sn alloys measured by DSC method

Alloy composition	Liquidus temperature $T_L$ (K)	Fusion enthalpy $\Delta H_f$ (kJ mol <sup>-1</sup> )	Fusion entropy $\Delta S_f$ (J mol <sup>-1</sup> K <sup>-1</sup> )
Ag-2.5%Sn	1223	8.81352	7.20648
Ag-5%Sn	1205	8.14955	6.76312
Ag-10%Sn	1153	7.58853	6.58155
Ag-14.2%Sn	1107	7.09429	6.40857
Ag-21%Sn	997	6.22016	6.23887
Ag-23%Sn	976	6.31741	6.45952
Ag-25%Sn	960	6.48410	6.75427
Ag-27%Sn	934	6.59270	7.05857
Ag-32%Sn	888	7.75715	8.73553
Ag-37%Sn	841	8.30077	9.87011
Ag-42%Sn	807	8.78680	10.88823
Ag-47%Sn	781	9.23046	11.81877
Ag-52%Sn	756	9.53129	12.60753
Ag-60%Sn	726	8.84006	12.17640
Ag-65%Sn	700	8.57640	12.25200
Ag-75%Sn	663	7.91837	11.94324
Ag-80%Sn	630	7.60435	12.07039
Ag-90%Sn	573	7.03490	12.27731
Ag-96.5%Sn	494	6.83622	13.83849

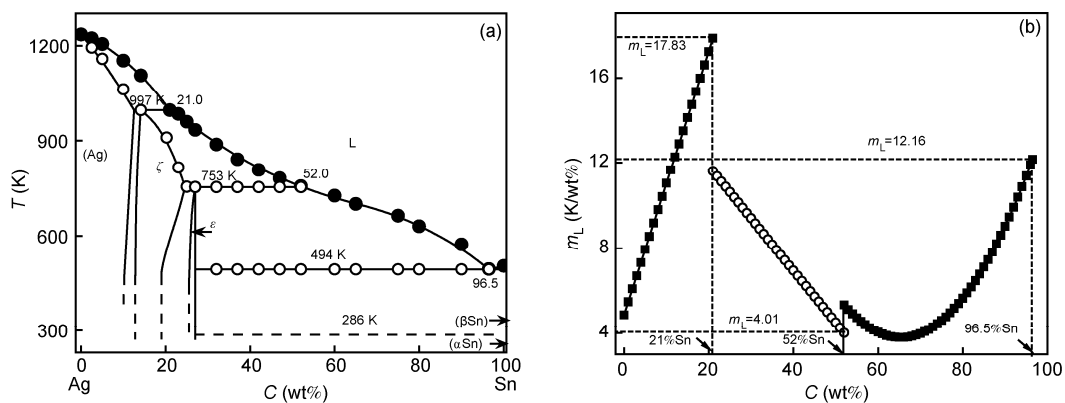
from high purity elements of Ag (99.999%) and Sn (99.999%) by laser melting under the protection of argon gas. The DSC experiments were carried out with a Netzsch DSC 404C differential scanning calorimeter. The calorimeter was calibrated with the melting points and the enthalpy of fusion for high purity In, Sn, Zn, Al, Ag, Au and Fe elements. The measuring accuracies of temperature and fusion of enthalpy are  $\pm 1$  K and  $\pm 3\%$  respectively, as verified by the measurements with pure Ag and Sn elements. Before each DSC experiment, the alloy specimen was placed in an Al<sub>2</sub>O<sub>3</sub> crucible. The chamber was evacuated and then back-filled with pure argon gas. The DSC thermal analyses were performed at different scan rates of 5 and 40 K/min, and the maximum heating temperatures were 100 K higher than the liquidus temperatures. Each specimen was heated, isothermally held at predetermined temperature, and then cooled at given scan rate for 2–3 cycles while kept in the DSC calorimeter, and the DSC profiles obtained in the last cycle was applied for further analyses. After the DSC experiments, the alloy specimens were polished and etched with a solution of 5 g FeCl<sub>3</sub>+1 mL HCl + 99 mL H<sub>2</sub>O, and were analyzed with an optical microscope.

## 2 Results and discussion

### 2.1 Liquidus and solidus temperatures, solidification temperature interval and solute partition coefficient

The measured liquidus temperatures of the selected alloys are marked in the Ag-Sn phase diagram shown in Figure 1(a) and listed in Table 1. Clearly, all the measured values agree well with the published phase diagram [17], which verifies the accuracy of DSC measurement. The relationship between measured liquidus temperature  $T_L$  and the Sn content  $C$  can be well fitted by the following three functions. If the Sn content ranges from 0–21wt%, where the (Ag) phase is the primary solid phase,

$$T_L = 1236 - 4.93C - 0.307C^2. \quad (1)$$



**Figure 1** Selected compositions and measured results of Ag-Sn alloys. (a) Measured liquidus and solidus temperatures of selected alloys illustrated in the Ag-Sn phase diagram; (b) liquidus slope versus Sn content.

In the Sn content range from 21wt% to 52wt%, in which the intermetallic compound  $\zeta$  phase solidifies preferentially from the alloy melt,

$$T_L = 1314 - 17.54C + 0.131C^2. \quad (2)$$

Once the Sn content increases from 52wt% to 96.5wt%, where the intermetallic compound  $\varepsilon$  phase is the primary solid phase:

$$T_L = 1731 - 39.77C + 0.551C^2 - 2.81 \times 10^{-3}C^3. \quad (3)$$

Based on eqs. (1)–(3), the liquidus slope, defined as  $m_L = -dT_L/dC$  can be calculated. If the Sn content ranges from 0 to 21wt%, the liquidus slope rises from 4.855 to 17.83 K/wt%:

$$m_L = 4.93 + 0.614C. \quad (4)$$

In the Sn content range from 21wt% to 52wt%, the slope decreases from 11.605 to 4.01 K/wt%:

$$m_L = 17.54 - 0.262C. \quad (5)$$

When the Sn content elevates from 52wt% to 96.5wt%, it goes up from 5.33 to 12.16 K/wt%:

$$m_L = 39.77 - 1.102C + 8.43 \times 10^{-3}C^2. \quad (6)$$

The solidus temperatures  $T_S$  can also be obtained by the DSC experiments, which are marked in Figure 2(a) by the hollow circles. On the basis of these measured liquidus and solidus temperatures, the solidification temperature interval  $\Delta T_0$  of (Ag),  $\zeta$  and  $\varepsilon$  phases is calculated, and the relationship between solidification temperature interval and alloy composition is plotted in Figure 2(a). If the Sn content is in the range from 0 to 14.12wt%, the solidification temperature interval for (Ag) phase is

$$\Delta T_0 = 0.40251 + 10.96968C - 0.22344C^2. \quad (7)$$

Once the Sn content rises from 23wt% to 52wt%, the solidification temperature interval for  $\zeta$  phase versus composition is

$$\Delta T_0 = 564.96627 - 17.73493C + 0.13275C^2. \quad (8)$$

Finally, if the Sn content locates between 52wt% and 96.5wt%, the solidification temperature interval for  $\varepsilon$  phase can be written as:

$$\Delta T_0 = 2492.29239 - 88.71705C + 1.17811C^2 - 5.45 \times 10^{-3}C^3. \quad (9)$$

Furthermore, the solute content  $C_L$  at liquidus temperature and  $C_s$  at solidus temperature for (Ag),  $\zeta$  and  $\varepsilon$  phases can be expressed by two functions,  $C_L = \Phi_1(T_L)$  and  $C_s = \Phi_2(T_s)$ , respectively. Thus, the variation of solute partition coefficient  $k_e = C_s/C_L$  with temperature for each phase can be derived. As illustrated in Figure 3(b), in the Sn content range from 0 to 21wt%, the solute partition coefficient for (Ag) phase is

$$k_e = -212737.30422 + 1168.29219C - 2.67104C^2 + 3.25 \times 10^{-3}C^3 - 2.22824 \times 10^{-6}C^4 + 8.13053 \times 10^{-10}C^5 - 1.2351 \times 10^{-13}C^6. \quad (10)$$

When the composition is between 21wt% and 52wt%Sn, the solute partition coefficient of  $\zeta$  phase is

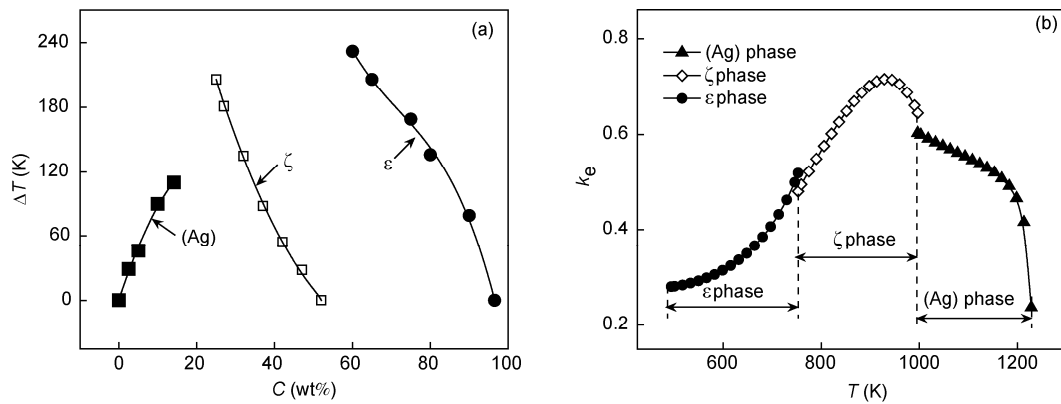
$$k_e = 15.50166 - 5.985 \times 10^{-2}C + 7.7419 \times 10^{-5}C^2 - 3.24246 \times 10^{-8}C^3. \quad (11)$$

If the Sn content is larger than 52wt%, the solute distribution of  $\varepsilon$  phase is

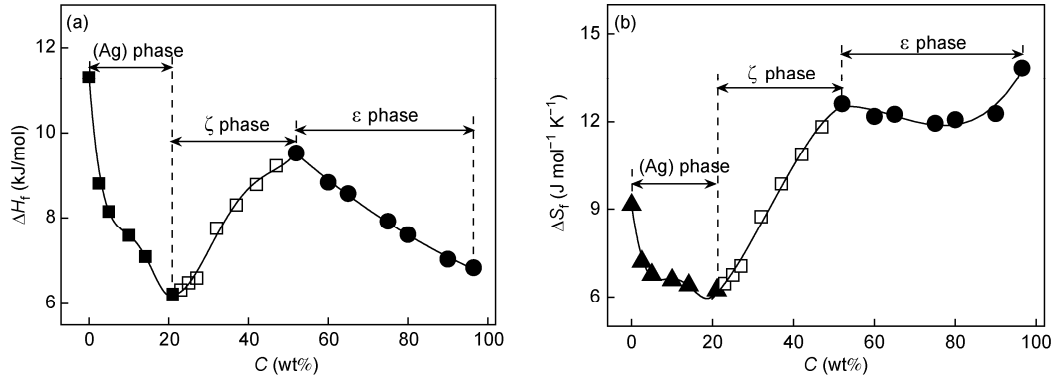
$$k_e = -1.58652 + 1.036 \times 10^{-2}C - 1.9597 \times 10^{-5}C^2 + 1.26733 \times 10^{-8}C^3. \quad (12)$$

## 2.2 Enthalpy and entropy of fusion

The enthalpy of fusion for Ag-Sn alloys versus Sn content in the whole composition range, covering all the heat absorption from solidus line to liquidus temperature, is determined by the solid-liquid transformation peaks during melting process at a scan rate of 5 K/min, as summarized in Table 1. According to Figures 1(a) and 3(a), the correlation



**Figure 2** Solidification temperature interval and solute partition coefficient of Ag-Sn alloys. (a) Measured solidification temperature interval; (b) solute partition coefficient for (Ag),  $\zeta$  and  $\varepsilon$  phases versus temperature.



**Figure 3** Measured enthalpy and entropy of fusion for Ag-Sn alloys versus Sn content. (a) Enthalpy of fusion; (b) entropy of fusion.

between enthalpy of fusion and composition is found to be closely related to the primary solid phase during the solidification of liquid Ag-Sn alloys. In the Sn content range from 0 to 21 wt%, the (Ag) phase is the corresponding primary solid phase, and the enthalpy of fusion of Ag-Sn alloys in this range decreases monotonically with increasing Sn content:

$$\Delta H_f = 11.3 - 1.20119C + 0.154343C^2 - 8.92 \times 10^{-3}C^3 + 1.78134C^4. \quad (13)$$

When Sn content is in the range of 21wt%–52wt%, the intermetallic compound  $\zeta$  phase solidifies preferentially from the liquid alloys. In this region, the enthalpy of fusion rises with the increase of Sn content and reaches a maximum value around 52wt%. The relationship between enthalpy of fusion and the Sn content can be written as:

$$\Delta H_f = 29.88249 - 3.0211C + 0.13449C^2 - 2.44 \times 10^{-3}C^3 + 1.59593 \times 10^{-5}C^4. \quad (14)$$

As for the composition range from 52wt% to 96.5wt% Sn, the intermetallic compound  $\varepsilon$  phase precipitates primarily from the alloy melt, and the enthalpy of fusion experiences monotonically decreasing with the increase of Sn content till 96.5wt%, which can be expressed as:

$$\Delta H_f = 14.96711 - 0.12845C + 4.54699 \times 10^{-4}C^2. \quad (15)$$

The entropy of fusion for Ag-Sn alloys  $\Delta S_f$  can also be calculated directly from the measured enthalpy and liquidus temperature by:

$$\Delta S_f = \Delta H_f / T_L. \quad (16)$$

The calculated results on entropy of fusion versus composition are also presented in Table 1 and shown in Figure 2(b). In the Sn range from 0 to 21wt%, the entropy of fusion decreases with the increase of Sn content, and can be expressed as:

$$\Delta S_f = 9.12565 - 1.07759C + 0.16992C^2 - 1.003 \times 10^{-2}C^3 + 2.11749 \times 10^{-4}C^4. \quad (17)$$

If Sn content locates between 21wt% and 52wt%, the entropy of fusion increases with the rise of Sn content, and their relationship can be written as:

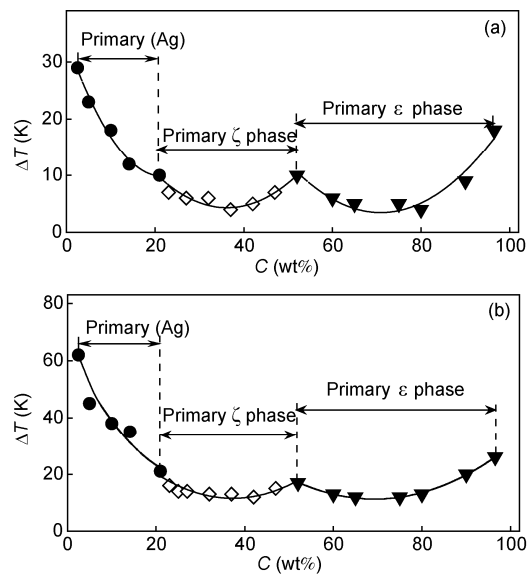
$$\Delta S_f = 7.82743 - 0.40294C + 0.01926C^2 - 1.87824 \times 10^{-4}C^3. \quad (18)$$

Once the Sn content is larger than 96.5wt%,

$$\Delta S_f = -9.61007 + 1.08715C - 0.01735C^2 + 8.89497 \times 10^{-5}C^3. \quad (19)$$

### 2.3 Undercoolability of liquid Ag-Sn alloys

The undercooling ( $\Delta T = T_L - T_{S1}$ ) of different Ag-Sn alloys obtained in the DSC calorimeter at scan rates of 5 and 40 K/min is measured. Here,  $T_{S1}$  is the initial solidification temperature of a specific Ag-Sn alloy upon cooling. Figure 4(a) illustrates the undercooling distribution at a scan rate of 5 K/min. The undercooling versus composition can be divided into three regions, which are 0–21wt% Sn, 21wt%–



**Figure 4** Measured undercoolings of liquid Ag-Sn alloys versus Sn content at different scan rates. (a) 5 K/min; (b) 40 K/min.

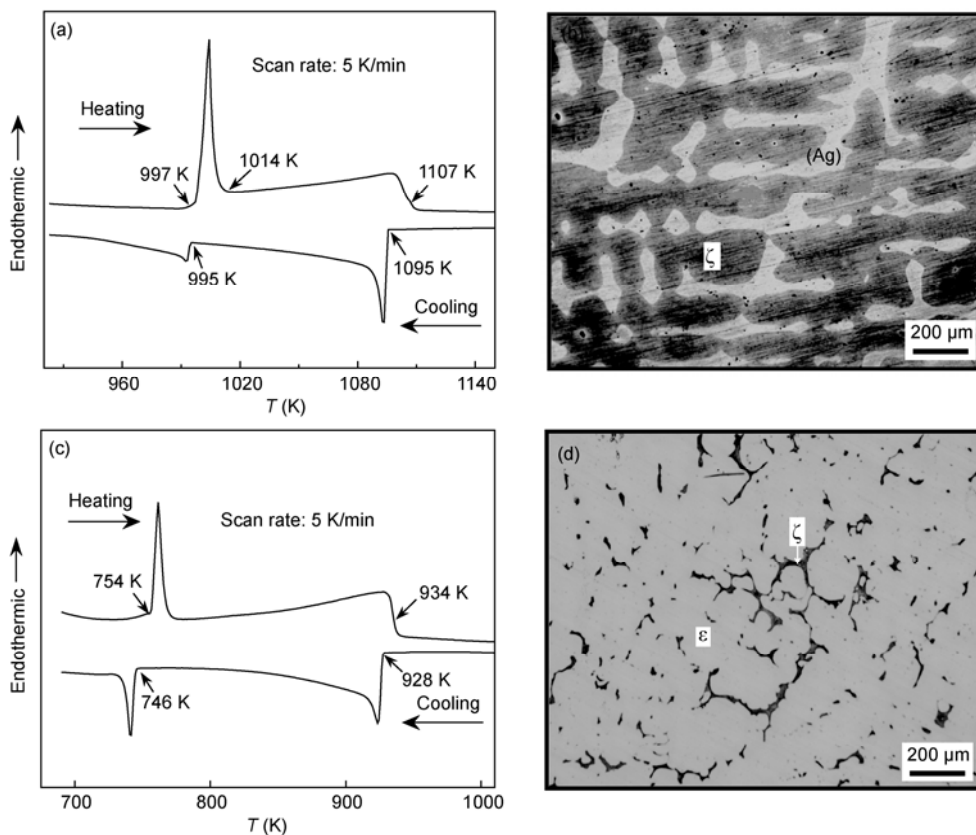
52wt% Sn and 52wt%–96.5wt% Sn. In the first region, the (Ag) phase always solidifies primarily in all the alloys, and the undercooling decreases from 29 to 10 K with the increase of Sn content. Then, the undercooling drops dramatically to about 5–10 K in the alloys of the second region. As for these alloys, their solidification processes initiate with the nucleation of intermetallic compound  $\zeta$  phase. In the third region, where the intermetallic  $\varepsilon$  phase nucleates preferentially from all the liquid alloys, the undercooling shows an increasing tendency from about 5 to 18 K with the increase of Sn content. If the cooling rate goes up to 40 K/min, as presented in Figure 4(b), the undercooling distribution shows the same changing tendency as that at 5 K/min. These results suggest that the undercoolability achieved in the DSC experiments is strongly dependent on the primarily nucleating solid phases and follows the relationship below:

$$\Delta T_{(\text{Ag})} > \Delta T_{\varepsilon} > \Delta T_{\zeta}. \quad (20)$$

The undercoolability of those liquid alloys solidifying with primary (Ag) solid solution phase is stronger than the other alloys with the preferential nucleation of intermetallic compounds  $\zeta$  and  $\varepsilon$  phases. Furthermore, it also needs to be mentioned that the undercooling level of all these alloys rises accordingly with the increase of cooling rate from 5 to 40 K/min. This indicates that faster cooling rate facilitates the higher undercooling of liquid Ag-Sn alloys.

## 2.4 DSC curves and microstructural features

In order to present a comprehensive survey on the phase transition characteristics, the DSC curves and growth morphologies for different types of Ag-Sn alloys are analyzed. Figure 5(a) and (b) show the DSC curves and solidified microstructure of peritectic Ag-14.12wt%Sn alloy. There are two endothermic events during its melting process. The first endothermic peak corresponds to the decomposition of solid peritectic  $\zeta$  phase into liquid and (Ag) phases, while the second one relates to the melting of (Ag) phase. The peritectic and liquidus temperatures of this alloy are 997 and 1107 K, respectively. During the cooling process, the primary (Ag) phase nucleates at 1095 K with a very sharp crystallization peak, and the following peritectic transition  $L+(\text{Ag}) \rightarrow \zeta$  at 995 K yields another relatively broad exothermal peak. As shown in Figure 5(b), the solidified microstructure consists of both the primary (Ag) dendrites and the peritectic  $\zeta$  phase, whose volume fractions are about 30% and 70%, respectively. In fact, under equilibrium condition, 100%  $\zeta$  phase is expected to be obtained at the end of solidification process. However, since the peritectic transformation is mainly controlled by atomic interdiffusion and it is very slow, the peritectic reaction can only occur to a limited extent even under slow cooling condition during the DSC experiments. Consequently, the microstructure is



**Figure 5** Thermal and structural analyses of two peritectic Ag-Sn alloys. (a) DSC curves of Ag-14.12wt%Sn alloy; (b) microstructure of Ag-14.12wt%Sn alloy; (c) DSC curves of Ag-27wt%Sn alloy and (d) microstructure of Ag-27wt%Sn alloy.

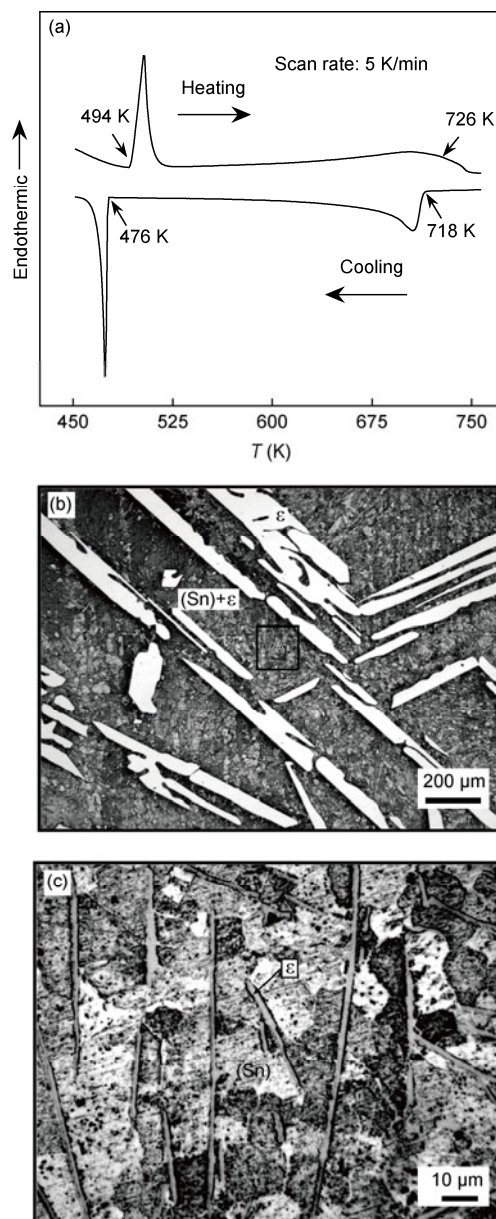
composed of peritectic  $\zeta$  phase and primarily precipitated (Ag) phase.

The DSC profiles of peritectic Ag-27wt%Sn alloy are presented in Figure 5(c). There are also two endothermic and two exothermic peaks during heating and cooling processes. The peritectic and liquidus temperatures of this peritectic alloy are 754 and 934 K. During cooling, the first exothermic peak at 928 K relates to the nucleation and growth of primary  $\zeta$  phase. The second exothermic event at 746 K corresponds to the peritectic transition  $L+\zeta\rightarrow\epsilon$ . As illustrated in Figure 5(d), the peritectic  $\epsilon$  phase grows into equiaxed grains. It is worthwhile to note that there still remains some primary  $\zeta$  dendrites, which distribute at the grain boundaries of peritectic  $\epsilon$  phase. This demonstrates again that the peritectic transformation can hardly be completed even under near-equilibrium condition.

Figure 6(a) depicts the thermographs of Ag-60wt%Sn hypoeutectic alloy, whose eutectic and liquidus temperatures are 494 and 726 K, respectively. During the heating process, the melting of ( $\epsilon$ +Sn) eutectic structure yields a sharp endothermic peak whereas that of primary  $\epsilon$  phase produces a very broad peak. In the cooling period, the primary  $\epsilon$  phase solidifies preferentially from the liquid alloy at 718 K, and it grows into long faceted plate, as shown in Figure 6(b). This differs with its nonfaceted growth morphology in Ag-27wt%Sn alloy as the peritectic phase. When the temperature drops to 476 K, eutectic transformation  $L\rightarrow(\epsilon+\text{Sn})$  occurs. As presented in Figure 6(c), the eutectic structure is characterized by a small amount of needle shaped  $\epsilon$  phase distributed in the eutectic (Sn) phase matrix. This is because the volume fraction of eutectic (Sn) phase is much higher than that of eutectic  $\epsilon$  phase, and regular lamellar eutectic structure can not form in this case.

### 3 Conclusions

In summary, the liquidus and solidus temperatures and enthalpy of fusion for binary Ag-Sn alloys in the whole composition range are determined by DSC method. The relationship between enthalpy of fusion and Sn content can be fitted by polynomial functions, which show one maximum value around 52wt%Sn and display two minimum values around 21 wt%Sn and 96.5 wt%Sn, respectively. The liquidus slope, the solidification temperature interval, the solute partition coefficient and the entropy of fusion are calculated on the basis of the measured results. The undercoolability of liquid Ag-Sn alloys depends mainly on the primary solid phases, and shows an increasing tendency with the increase of cooling rate. The undercoolings of those liquid alloys solidifying with primary (Ag) solid solution phase are higher than the other alloys with the preferential nucleation of  $\zeta$  and  $\epsilon$  intermetallic compounds. The morphological observation reveals that the primary  $\epsilon$  phase in Ag-60wt%Sn



**Figure 6** DSC thermograph and microstructures of hypoeutectic Ag-60wt%Sn alloy. (a) DSC curves; (b) growth morphology of primary  $\epsilon$  phase; (c) microstructure of (Sn+ $\epsilon$ ) eutectic in Zone A marked in (b).

hypoeutectic alloy grows into faceted plate, and the ( $\epsilon$ +Sn) eutectic structure is characterized by needle shaped  $\epsilon$  phase distributed in the (Sn) matrix. The peritectic reactions can rarely be completed in the Ag-14.12wt%Sn and Ag-27wt%Sn peritectic alloys, and their solidification microstructures are composed of both primary and peritectic phases.

*This work was supported by the National Natural Science Foundation of China (50971105 and 51201136), Doctoral Fund of Ministry of Education of China (20126102120059), Aviation Foundation of China (2012ZF53069), Technology Foundation for Selected Overseas Chinese Scholar, ShaanXi Province, Northwestern Polytechnical University Excellent Personnel*

Supporting Project of Ao Xiang Star, and Fundamental Research Fund of Northwestern Polytechnical University (JC20110280).

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