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## **OPEN** Evidence for the transition from primary to peritectic phase growth during solidification of undercooled Ni-Zr alloy levitated by electromagnetic field

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The Ni<sub>83.25</sub>Zr<sub>16.75</sub> peritectic alloy was undercooled by electromagnetic levitation method up to 198 K. The measured dendritic growth velocity shows a steep acceleration at a critical undercooling of  $\Delta T_{crit}$  = 124 K, which provides an evidence of the transition of the primary growth mode from Ni<sub>7</sub>Zr<sub>2</sub> phase to peritectic phase Ni<sub>5</sub>Zr. This is ascertained by combining the temperature-time profile and the evolution of the solidified microstructures. Below the critical undercooling, the solidified microstructure is composed of coarse Ni<sub>7</sub>Zr<sub>2</sub> dendrites, peritectic phase Ni<sub>5</sub>Zr and eutectic structure. However, beyond the critical undercooling, only a small amount of Ni<sub>7</sub>Zr<sub>2</sub> phase appears in the solidified microstructure. The dendritic growth mechanism of Ni<sub>7</sub>Zr<sub>2</sub> phase is mainly governed by solute diffusion. While, the dendritic growth mechanism of Ni<sub>5</sub>Zr phase is mainly controlled by thermal diffusion and liquid-solid interface atomic attachment kinetics.

Peritectic solidification is frequently encountered among metallic alloy systems, such as Fe-Ni, Fe-Al, Al-Ni, Ni-Zr, etc.<sup>1-5</sup>, which is of great importance in preparing various commercial component materials. Recently, a transition of the primary growth mode from primary phase to peritectic phase during the solidification of undercooled peritectic alloys has been paid considerable attention<sup> $\delta$ -10</sup>. The underlying reason is that the transition of the primary growth mode may result in the formation of phase-pure peritectic phase in the final solidified microstructure, and thus improve the performance of peritectic alloys<sup>7</sup>. Tourret et al.<sup>4,5</sup> investigated the multiple phase transformations of Al-Ni peritectic alloys by electromagnetic levitation and gas atomization methods, and found that peritectic phase Al<sub>3</sub>Ni preferentially grows when the droplet diameter is 10µm for gas atomized Ni-80 at% Al hyperperitectic alloy, which is attributed to the competition between the cooling kinetics and the diffusion kinetics. Phanikumar et al.<sup>11</sup> found that once the undercooling (unless stated otherwise, any mention of undercooling in the paper refers to nucleation undercooling) exceeds a critical value of about 110 K, the solidified microstructure consists of only peritectic phase in Fe-25%Ge peritectic alloy processed by an electromagnetic levitator. Leonhardt et al.<sup>12</sup> reported that a transition of the primary growth mode from primary bcc-Mo to peritectic  $\sigma$ -phase was revealed if the undercooling of Fe<sub>47</sub>Mo<sub>53</sub> alloy is beyond 345 K. However, although some significant progresses have been reported, direct experimental evidence of the transition of the primary growth mode from the primary phase to peritectic phase is rather limited. Previous studies revealed this transition of primary growth mode mainly by the evolution of solidified microstructures. Actually, dendritic growth is the major growth mode in undercooled melts, the velocity of which can provide valuable insight into the transition of primary growth mode. Nevertheless, few investigations of dendritic growth kinetics in undercooled peritectic alloys have been reported.

As a typical peritectic alloy system, Ni-Zr binary alloy system has aroused particular interests due to its good glass forming ability in a wide compositional range<sup>13-15</sup> as well as its abundant intermetallic compounds<sup>16-18</sup>. Ni<sub>83.25</sub>Zr<sub>16.75</sub> is a peritectic composition in Ni-Zr alloy system, whose primary phase and peritectic phase are

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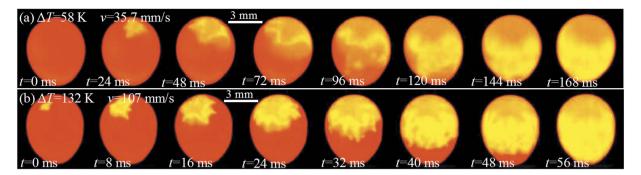


Figure 1. High-speed camera images of recalescence front of undercooled Ni<sub>83.25</sub>Zr<sub>16.75</sub> peritectic samples at different undercoolings. The red part is the undercooled liquid, and the yellow part is the recalescing solid. The time interval between two adjacent images is 24 ms and 8 ms, respectively. (a)  $\Delta T = 58$  K; (b)  $\Delta T = 132$  K.

both intermetallic compounds. The solidification behavior for this type of peritectic alloy is complicated but also considerably important for deep and comprehensive understanding of peritectic solidification under highly undercooled condition. Therefore, the objective of this work is to investigate the transition of the primary growth mode from  $Ni_7Zr_2$  phase to peritectic phase  $Ni_5Zr$  during the solidification of undercooled  $Ni_{83,25}Zr_{16,75}$  peritectic alloy by the measured dendritic growth velocity and solidified microstructures. Meanwhile, the dendritic growth kinetics of  $Ni_7Zr_2$  phase and  $Ni_5Zr$  phase is also studied to reveal the evolution of solidified microstructure.

#### **Results and Discussion**

The high-speed camera technique can allow the visualization of the propagation front, which is a feasible approach to investigate the rapid solidification process. Figure 1 shows a few snapshots of rapid solidification front in undercooled  $Ni_{83.25}Zr_{16.75}$  peritectic melts at different undercoolings, which were captured by a Red-lake HG 100 K high-speed camera with the resolution of 24 bits (color) pixel depth. The yellow area corresponds to the solid due to the released heat, and the red area corresponds to the undercooled liquid. Noticeably, solidification starts at the upper surface of the sample and proceeds to the lower part. The propagating front appears ambiguous for low undercooling, and gives distinct feature of dendritic structure for high undercooling. The dendritic growth velocity can be determined from the sequence of projected images captured by the high-speed camera, which is coincident with the value measured by a photoelectric detector.

The results of dendritic growth velocity with different undercoolings in Ni<sub>83.25</sub>Zr<sub>16.75</sub> melts are presented in Fig. 2(a), which was measured by a photoelectric detector. The maximum undercooling obtained in the present work is about 198 K. It is obvious that the measured dendritic growth velocity continuously increases with the enhancement of the undercooling. When the undercooling is smaller than a critical value of  $\Delta T_{crit} = 124$  K, the growth velocity appears sluggishly. Once the undercooling exceeds the critical value of  $\Delta T_{crit} = 124$  K, the growth velocity increases rapidly. More importantly, a steep rise of the growth velocity is observed at the critical undercooling of  $\Delta T_{crit} = 124$  K, which jumps from 61 mm/s to 88 mm/s. Such a phenomenon implies that a transition of the primary growth mode from Ni<sub>7</sub>Zr<sub>2</sub> phase to peritectic phase Ni<sub>5</sub>Zr occurs. The equilibrium solidification of Ni<sub>7</sub>Zr<sub>2</sub> phase is replaced by the direct growth of peritectic alloy, the liquidus temperature of Ni<sub>7</sub>Zr<sub>2</sub> phase is just higher than that of Ni<sub>5</sub>Zr phase by about 39 K. If the undercooling of the melt goes through the peritectic temperature  $T_p$ , the peritectic phase Ni<sub>5</sub>Zr becomes a metastable phase and may form directly from the undercooled melt despite the low thermodynamic driving force. Our previous study<sup>3</sup> has revealed that the peritectic phase Ni<sub>5</sub>Zr can form directly from the undercooled melt by completely suppressing the growth of Ni<sub>7</sub>Zr<sub>2</sub> phase if the droplet diameter is less than a critical value in the drop tube experiments.

To verify the transition of the primary growth mode, the temperature-time curves during undercooling and rapid solidification of  $Ni_{83.25}Zr_{16.75}$  peritectic melts at two different undercoolings are illustrated in Fig. 2(b). The recalescence behavior is characterized by a steep temperature rise detected by the pyrometer. For low undercooling of 58 K, the temperature of the melt after recalescence rises nearly to the liquidus temperature  $T_L$ , which gives an indication of growth of primary  $Ni_7Zr_2$  dendrites. However, for high undercooling of 160 K, the recalescence process is observed to stop below the peritectic temperature  $T_P$ . This may give an evidence for a change of growth mode that the growth of  $Ni_7Zr_2$  phase is replaced by the growth of peritectic phase  $Ni_5Zr$  if the undercooling exceeds the critical value of 124 K.

The microstructures of Ni<sub>83.25</sub>Zr<sub>16.75</sub> peritectic samples solidified at different undercoolings are shown in Fig. 3, in which both the Ni<sub>7</sub>Zr<sub>2</sub> phase and Ni<sub>5</sub>Zr phase have been marked. The solidified microstructure consists of Ni<sub>7</sub>Zr<sub>2</sub> dendrites, peritectic phase Ni<sub>5</sub>Zr and inter-dendritic eutectic microstructure for low undercooling of  $\Delta T = 9$  K, as illustrated in Fig. 3(a). Apparently, the Ni<sub>7</sub>Zr<sub>2</sub> phase exhibits coarse and developed dendrites, which is enwrapped by peritectic phase Ni<sub>5</sub>Zr. Figure 3(b) is an enlarged view of the inter-dendritic eutectic microstructure of (Ni) and Ni<sub>5</sub>Zr phases. With the enhancement of undercooling, the fragment of Ni<sub>7</sub>Zr<sub>2</sub> dendrites occurs which seems like the primary Ni<sub>7</sub>Zr<sub>2</sub> dendrite trunks have partially been transformed to peritectic phase, and the fragmented zone is marked as a box, as shown in Fig. 3(c). However, when the undercooling is beyond the critical value of  $\Delta T_{crit} = 124$  K, a significant change of microstructure takes place. The solidified microstructure for a

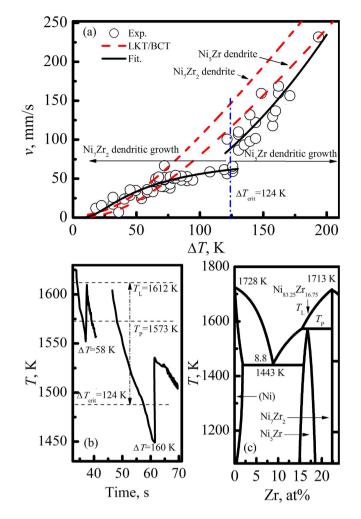
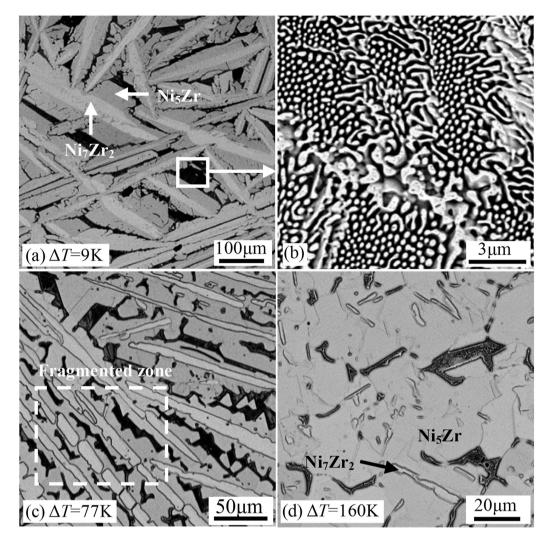


Figure 2. (a) Dendritic growth velocity versus undercooling. (b) Temperature-time recalescence characteristics at different undercoolings. (c) The left part of Ni-Zr binary phase diagram<sup>24</sup>.

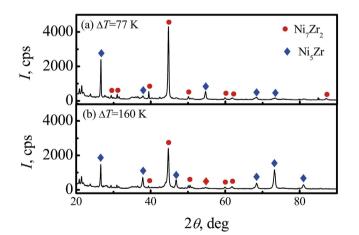
high undercooling of  $\Delta T = 160$  K is composed of a small amount of Ni<sub>7</sub>Zr<sub>2</sub> phase, predominant Ni<sub>5</sub>Zr phase and eutectic microstructure, as presented in Fig. 3(d). It is evident that the amount of Ni<sub>7</sub>Zr<sub>2</sub> phase is very low, and the Ni<sub>7</sub>Zr<sub>2</sub> phase seems to be decomposed and nearly disappears.

To further confirm the variation of phase constitution, X-ray diffraction (XRD) patterns of samples solidified at two different undercoolings of 77 K and 160 K are shown in Fig. 4. It can be seen that the main peaks of  $Ni_7Zr_2$  phase decrease sharply with the increasing undercooling. On the contrary, the peaks of  $Ni_5Zr$  phase increase with the enhancement of undercooling. This indicates that the volume fraction of  $Ni_5Zr$  phase at high undercooling is larger than that at low undercooling.

According to the above microstructures presented in Fig. 3, two growth modes can be concluded. If the undercooling is smaller than the critical value of  $\Delta T_{\text{crit}} = 124$  K, the Ni<sub>7</sub>Zr<sub>2</sub> phase is preferred to primarily nucleate and grow into the manner of dendrites during the rapid solidification of the undercooled melt, which results in a sudden temperature rise due to the released heat of crystallization. Subsequently, with the decrease of temperature, the peritectic phase Ni<sub>5</sub>Zr starts to nucleate at the surface of the Ni<sub>7</sub>Zr<sub>2</sub> dendrites when the temperature drops below the peritectic temperature T<sub>P</sub> In this case, the primary Ni<sub>7</sub>Zr<sub>2</sub> dendrites, peritectic phase Ni<sub>5</sub>Zr and liquid phase will contact with each other at a triple junction, which is the requirement of peritectic reaction. Then, peritectic reaction of  $Ni_7Zr_2 + L \rightarrow Ni_5Zr$  takes place and peritectic phase  $Ni_5Zr$  grows along the surface of  $Ni_7Zr_2$ dendrites to form a thin peritectic layer. Peritectic reaction, which is governed by local short range diffusion of the solute in the melt ahead of the primary Ni<sub>7</sub>Zr<sub>2</sub> dendrites and peritectic phase Ni<sub>5</sub>Zr, can proceed rapidly at the initial stage of the peritectic growth process. Whereas, once the primary  $Ni_7Zr_2$  dendrites are enwrapped by peritectic phase Ni<sub>5</sub>Zr, peritectic phase Ni<sub>5</sub>Zr will separate the primary phase Ni<sub>7</sub>Zr<sub>2</sub> and liquid phase, leading to the disappearance of the triple junction and the cease of peritectic reaction. After which, the peritectic phase Ni<sub>5</sub>Zr grows into the primary Ni<sub>7</sub>Zr<sub>2</sub> dendrites by peritectic transformation. Since the peritectic transformation is controlled by long range solid-state diffusion, it proceeds sluggishly. Unfortunately, the cooling rate in the experiments is about 15 K/s, which results in that peritectic transformation could not proceed completely<sup>11</sup> and only a small amount of primary dendrites could transform to peritectic phase by peritectic transformation. Therefore, the primary phase is always retained in the final microstructures after peritectic solidification, as shown in Fig. 3(a-c). Due to the existence of the Ni<sub>7</sub>Zr<sub>2</sub> phase, the composition of residual liquid deviates from



**Figure 3.** Solidified microstructures of undercooled Ni<sub>83.25</sub>Zr<sub>16.75</sub> peritectic alloy at undercooling of (**a**,**b**)  $\Delta T = 9$  K, (**c**)  $\Delta T = 77$  K, and (**d**)  $\Delta T = 160$  K.



**Figure 4.** XRD patterns of samples solidified at undercoolings of (a)  $\Delta T = 77$  K and (b)  $\Delta T = 160$  K to reveal the volume change of the two phases.

the initial composition of the melt and moves to eutectic zone according to the phase diagram in Fig. 2(c). Hence, the residual liquid solidifies as eutectic when the temperature drops below the eutectic temperature, which is presented in Fig. 3(b).

Once the undercooling exceeds the critical value of  $\Delta T_{crit} = 124$  K, there exists only a small amount of Ni<sub>7</sub>Zr<sub>2</sub> phase in the solidified microstructure, as illustrated in Fig. 3(d). There are two possible solidification paths for the formation of such a microstructure. The first possibility is that only a small amount of Ni<sub>7</sub>Zr<sub>2</sub> phase forms first but peritectic phase Ni<sub>5</sub>Zr prefers to grow from the undercooled melts. Thus, the microstructure is composed of predominant peritectic phase  $Ni_5Zr$  and a small amount of  $Ni_7Zr_2$  phase. The second possibility is that  $Ni_5Zr$ phase primarily grows in the undercooled melt. Since the atoms of different species have to sort themselves onto proper lattice place during the growth of intermetallic compounds, the growth velocity of Ni<sub>5</sub>Zr phase is sluggish. The released heat of crystallization results in a steep rise of temperature and a decrease of interface undercooling. Thus, the growth of peritectic Ni5Zr phase cannot proceed to completion and a small amount of liquid remains in the inter-dendritic region. The undercooling of residual liquid after recalescence is smaller than 124 K according to the temperature data in Fig. 2(b). In this case, Ni<sub>7</sub>Zr<sub>2</sub> is preferred to grow from the residual undercooled liquid. Actually, due to less released heat and high cooling rate during the solidification of Ni<sub>7</sub>Zr<sub>2</sub> phase, the second recalescence is difficult to distinguish from the undulations in the pyrometer signal in Fig. 2(b). The second possibility is more likely to occur according to our previous study<sup>3</sup>. We suggest that peritectic phase Ni<sub>5</sub>Zr preferentially grows when the undercooling is larger than 124 K. It is verified again that the equilibrium solidification of  $Ni_7Zr_2$  phase is replaced by the direct growth of peritectic phase  $Ni_5Zr$  when the undercooling is beyond the critical value of 124 K.

A heat flux from the melt to the surrounding is necessary during the solidification, which is dominated by cooling rate. If the cooling rate of undercooled Ni<sub>83.25</sub>Zr<sub>16.75</sub> melt is sufficiently high, the crystallization heat would be rapidly transferred to the surrounding during the growth of peritectic phase Ni<sub>5</sub>Zr. This will result in the continuous growth of peritectic phase Ni<sub>5</sub>Zr and the formation of phase-pure peritectic phase Ni<sub>5</sub>Zr microstructure. In the case of electromagnetic levitation, heat is mainly transferred by flowing helium gas. To obtain high cooling rate and verify the speculation, a Ni<sub>83.25</sub>Zr<sub>16.75</sub> sample was undercooled up to 160 K and then quenched on a Cu-substrate. The cross-sectional micrographs of different zones in the quenched sample are shown in Fig. 5. Figure 5(b) presents the microstructure away from the Cu-substrate, which consists of Ni<sub>7</sub>Zr<sub>2</sub> dendrites, peritectic phase Ni<sub>5</sub>Zr and eutectic. The microstructures on the Cu-substrate side are illustrated in Fig. 5(c,d). Obviously, the microstructure is composed of two regions. The upper one is characterized by the primary  $Ni_7Zr_2$ dendrites enveloped by peritectic phase Ni<sub>5</sub>Zr. The below one adjacent to the Cu-substrate consists of only peritectic phase  $Ni_5Zr$  with no primary phase  $Ni_7Zr_2$  because high cooling rate is obtained at the interface between the Cu-substrate and melt. To check the reproducibility of the observation, the quench experiments were performed on two samples and the results agree well. This suggests that peritectic phase Ni<sub>5</sub>Zr directly solidifies by completely suppressing the growth of the primary Ni<sub>7</sub>Zr<sub>2</sub> dendrites. Furthermore, this is an evident proof for the transition of the primary growth mode from Ni<sub>7</sub>Zr<sub>2</sub> phase to peritectic Ni<sub>5</sub>Zr phase when the undercooling exceeds the critical value of 124 K.

Dendritic growth is the major growth mode in undercooled melts, which determines the evolution of the solidified microstructure. Meanwhile, dendritic growth is controlled by the temperature and concentration gradients, resulting from the heat and solute transport around the solid-liquid interface. To analyze the dendrites growth kinetics of Ni<sub>2</sub>Zr, and Ni<sub>5</sub>Zr, a LKT/BCT model<sup>19-21</sup> is adopted to describe the dendritic growth as a function of undercooling. The physical parameters used in the calculations are obtained by molecular dynamics simulation and linearly fitting the values of pure metals<sup>22</sup>, which are listed in Table 1. The calculated dendritic growth velocities of Ni<sub>7</sub>Zr<sub>2</sub> and Ni<sub>5</sub>Zr phase are shown in Fig. 2(a). Evidently, the calculated results of Ni<sub>7</sub>Zr<sub>2</sub> phase are in good agreement with the experimental results when the undercooling is smaller than 80 K. The dendritic growth mechanism of Ni<sub>7</sub>Zr<sub>2</sub> phase is mainly governed by solute diffusion. Similarly, the calculated dendritic growth velocity of Ni5Zr phase is also close to the experimental values. The initial composition of the melts is the same as that of peritectic phase Ni<sub>5</sub>Zr. Hence, if peritectic phase Ni<sub>5</sub>Zr preferentially grows, mass transport by segregation and constitutional effects can be neglected<sup>23</sup>, thus, the constitutional undercooling  $\Delta T_c = 0$ . The curvature undercooling is usually small due to the large curvature radius of dendrites, which also can be neglected. Therefore, the dendritic growth of Ni<sub>5</sub>Zr phase is controlled by thermal undercooling and kinetic undercooling. In other words, thermal diffusion and liquid-solid interface atomic attachment kinetics play a vital role in determining the growth velocity of Ni<sub>5</sub>Zr phase.

#### Conclusion

In summary, the dendritic growth in undercooled Ni<sub>83.25</sub>Zr<sub>16.75</sub> peritectic alloy was investigated by electromagnetic levitation method. The maximum undercooling achieved in the experiment is 198 K. The dendritic growth velocity shows a steep acceleration around a critical undercooling of  $\Delta T_{\rm crit}$  = 124 K, which gives the evidence of the transition of the primary growth mode from Ni<sub>7</sub>Zr<sub>2</sub> phase to Ni<sub>5</sub>Zr phase. This is ascertained by combining the temperature-time profile and the evolution of the microstructures. The solidified microstructure is composed of coarse Ni<sub>7</sub>Zr<sub>2</sub> dendrites, peritectic phase Ni<sub>5</sub>Zr and eutectic structure when the undercooling is less than the critical undercooling of  $\Delta T_{\rm crit}$  = 124 K. However, only a small amount of Ni<sub>7</sub>Zr<sub>2</sub> phase appears in the solidified microstructure once the undercooling exceeds the critical value of 124 K, which indicates that the peritectic phase Ni<sub>5</sub>Zr primarily solidifies. Furthermore, In the case of dropping the undercooled sample of 160 K onto a Cu-substrate, the microstructure of the quenched sample adjacent to the Cu-substrate consists of only peritectic phase Ni<sub>5</sub>Zr with no primary phase Ni<sub>7</sub>Zr<sub>2</sub>, which suggests that peritectic phase Ni<sub>5</sub>Zr directly solidifies by completely suppressing the growth of the primary phase Ni<sub>7</sub>Zr<sub>2</sub>. The dendritic growth mechanism of Ni<sub>7</sub>Zr<sub>2</sub> phase is mainly governed by solute diffusion. However, thermal diffusion and liquid-solid interface atomic attachment kinetics play a vital role in determining the growth velocity of Ni<sub>5</sub>Zr phase.

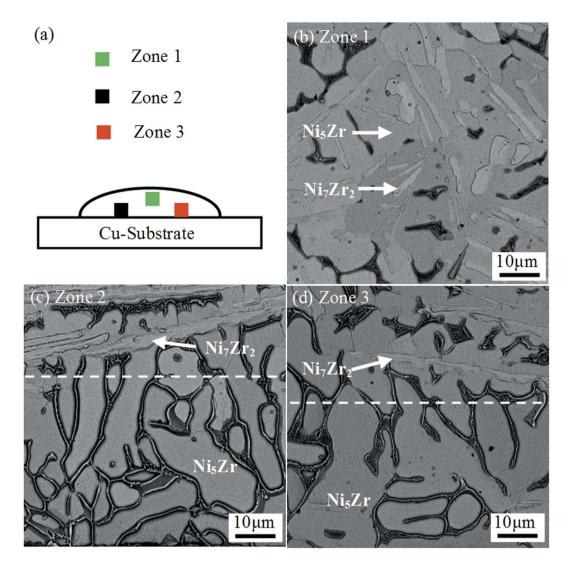


Figure 5. Cross-sectional SEM image of the sample dropped onto the Cu-substrate after undercooled to 160 K. (a) Schematic of the Cross-sectional view of the dropped sample. (b) Microstructure in Zone 1.
(b) Microstructure in Zone 2. (3) Microstructure in Zone 3.

Parameter	Unit	Ni <sub>7</sub> Zr <sub>2</sub>	Ni <sub>5</sub> Zr
Heat of fusion	$\Delta H$ (J/mol)	21000	18500
Heat capacity	$C_{\rm p}$ (J/mol)	39.4	39.8
Liquidus slope	m <sub>l</sub> (K/at.%)	8.6	0
Melt diffusion coefficient	$D_{\rm L} ({\rm m^2/s})$	$1.22 \times 10^{-7} \exp(-54951/RT)$	$1.22 \times 10^{-7} \exp(-54951/RT)$
Gibbs-Thomosion parameter	Г (mK)	$1.40 imes10^{-7}$	$1.41  imes 10^{-7}$
Velocity of sound	V <sub>0</sub> (m/s)	3000	3000

### Table 1. Physical parameters of $Ni_{83.25}Zr_{16.75}$ peritectic alloy<sup>22</sup>.

#### **Experimental Details**

Master alloys of Ni<sub>83.25</sub>Zr<sub>16.75</sub> peritectic alloy were prepared by 99.99% pure Ni and 99.9% pure Zr mixtures in an arc melting furnace. The samples of about 0.6 g were levitated and melted by an electromagnetic levitation facility, which was evacuated to  $10^{-5}$  Pa and backfilled with argon gas to 1 atm. The sample was cooled with flowing helium gas to achieve substantial undercooling. Its temperature was measured using a one-color Raytek Marathon MR1SCSF infrared pyrometer, which was calibrated by a PtRh30-PtRh6 thermocouple. The dendritic growth velocity was determined from the recalescence time measured by a photoelectric detector. The solidified samples and phase constitution were analyzed by a Phenom Pro SEM and a Rigaku D/max 2500 X-ray diffractometer (XRD).

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#### **Author Contributions**

P. Lü and H.P. Wang designed the experiments. P. Lü carried out the experiments and wrote the paper. H.P. Wang and K. Zhou revised this paper.

#### Additional Information

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