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ARTICLE

# Effect of Addition of Er-TiB<sub>2</sub> Dual-Phase Nanoparticles on Strength-Ductility of Al-Mn-Mg-Sc-Zr Alloy Prepared by Laser Powder Bed Fusion

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**Abstract:** A dual-phase synergistic enhancement method was adopted to strengthen the Al-Mn-Mg-Sc-Zr alloy fabricated by laser powder bed fusion (LPBF) by leveraging the unique advantages of Er and TiB<sub>2</sub>. Spherical powders of 0.5wt% Er-1wt% TiB<sub>2</sub>/Al-Mn-Mg-Sc-Zr nanocomposite were prepared using vacuum homogenization technique, and the density of samples prepared through the LPBF process reached 99.8%. The strengthening and toughening mechanisms of Er-TiB<sub>2</sub> were investigated. The results show that Al<sub>3</sub>Er diffraction peaks are detected by X-ray diffraction analysis, and texture strength decreases according to electron backscatter diffraction results. The added Er and TiB<sub>2</sub> nano-reinforcing phases act as heterogeneous nucleation sites during the LPBF forming process, hindering grain growth and effectively refining the grains. After incorporating the Er-TiB<sub>2</sub> dual-phase nano-reinforcing phases, the tensile strength and elongation at break of the LPBF-deposited samples reach 550 MPa and 18.7%, which are 13.4% and 26.4% higher than those of the matrix material, respectively.

**Key words:** Al-Mn-Mg-Sc-Zr alloy; laser powder bed fusion; nano-reinforcing phase; synergistic enhancement

## 1 Introduction

Aluminum alloys are critical structural materials renowned for their high strength-to-weight ratio, superior impact resistance, excellent ductility, and corrosion resistance, making them widely used in the aerospace industry<sup>[1-2]</sup>. Laser powder bed fusion (LPBF) technique significantly improves material utilization, reduces manufacturing cycles, and lowers costs. A distinctive feature of LPBF is its ability to rapidly melt metal powders via a laser beam, followed by immediate solidification, achieving cooling rates of 10<sup>4</sup>–10<sup>6</sup> K/s. This rapid cooling characteristic, driven by large temperature

gradients, leads to significant refinement in the alloy's microstructure<sup>[3]</sup>. Moreover, under such fast-cooling conditions, the ultimate solid solubility of alloying elements increases markedly, enabling the formation of a supersaturated solid solution with a solubility far exceeding the equilibrium solid solubility of the alloying elements, and thereby improving the mechanical properties of the components.

Currently, most researches on LPBF-fabricated aluminum alloys are confined to Al-Si and Al-Mg-Si systems. Although these alloys can effectively reduce product mass, their strength and toughness are insufficient to meet the final

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application requirements as key materials for complex functional components in aerospace vehicles<sup>[4]</sup>. The 2xxx and 7xxx series aluminum alloys have wide solidification intervals and high hot cracking susceptibility, posing significant challenges to forming during laser additive manufacturing<sup>[5]</sup>. In comparison, the Al-Mn-Mg-Sc-Zr has relatively balanced overall performance. However, its tensile strength is still rather low, failing to meet the performance requirements of the aerospace field. This is attributed to its relatively high laser reflectivity and thermal conductivity, necessitating higher laser energy input. Consequently, this leads to poor forming stability and considerable fabrication difficulties<sup>[6-7]</sup>.

Li et al<sup>[8]</sup> prepared TiB<sub>2</sub>/AlSi<sub>10</sub>Mg specimens by LPBF, which significantly improved the tensile strength and tensile strain of the formed parts. The tensile strength reached 530±16 MPa, and the elongation reached 15.5%±1.2%, making this alloy an aluminum matrix composite formed by LPBF with relatively high comprehensive mechanical properties. Shi et al<sup>[9]</sup> investigated the influence of element Er on the microstructure of aluminum alloys. Through optimization of process parameters and adjustment of heat treatment, they increased the tensile strength from 415.2 MPa to 524.4 MPa. Martin et al<sup>[10]</sup> from HRL Laboratory (USA) coated 7075Al powder particles with nanoparticles such as TiB<sub>2</sub>, ZrH<sub>2</sub>, and WC. The addition of these nanoparticles inhibited the hot cracking tendency during the LPBF process, enabled the transformation from columnar to equiaxed grains, and reduced material anisotropy. Wang et al<sup>[11]</sup> prepared Ti+B<sub>4</sub>C/AA7075 composite powder by high-energy ball milling, increasing its tensile strength from 200.1 MPa to 336.9 MPa.

Studies have demonstrated that the rare earth element Er has a strong modifying effect. Similar to Al<sub>3</sub>Sc, the primary phase Al<sub>3</sub>Er has good interfacial coherence with the Al matrix, enabling it to act as a non-homogeneous nucleation core to greatly improve the nucleation rate. In addition, fine Al<sub>3</sub>Er particles can pin dislocations and subgrain boundaries, hindering dislocation aggregation and migration at subgrain boundaries, thus delaying the nucleation and growth of subgrains<sup>[12]</sup>. The laser reflectivity of ceramic particles is much lower than that of aluminum alloys, so TiB<sub>2</sub> is used as a reinforcing phase to strengthen aluminum alloys. When uniformly attached to the surface of the aluminum matrix,

TiB<sub>2</sub> improves the laser absorption capacity of aluminum alloys, resulting in better LPBF formability<sup>[13]</sup>.

In this research, a dual-phase synergistic reinforcement approach using Er and TiB<sub>2</sub> was employed to enhance the Al-Mn-Mg-Sc-Zr alloy, leveraging the respective strengthening advantages of Er and TiB<sub>2</sub>. It is expected that the dual-phase reinforcement of rare earth elements and ceramic particles will provide a new insight into the formation of novel aluminum matrix composites by LPBF.

## 2 Experiment

The powder used in this experiment was Al-4.88Mn-1.66Mg-0.69Sc-0.33Zr, developed by Suzhou Beifeng Technology, with a particle size range of 15–53 μm. The selected Er powder had a particle size of 1 μm and the TiB<sub>2</sub> powder had a particle size of 100 nm. 1wt% TiB<sub>2</sub> particles and 0.5wt% Er powder were mixed with the Al-Mn-Mg-Sc-Zr alloy by vacuum homogenization method. The vacuum homogenizer ZYMC-200V was used. The morphology and particle size of the mixed composite powder are shown in Fig.1a and Fig.1b, respectively.

The LPBF device used in this study was SLM100-N2, independently developed by Xi'an Jiaotong University. Cubic sample of 8 mm×8 mm×8 mm was printed under the constant parameters: laser power of 400 W, scanning speed of 1000 mm/s, hatch spacing of 100 μm, and layer thickness of 30 μm. In the LPBF process, the oxygen content in the forming chamber was controlled within 10<sup>5</sup> μg/L to prevent high-temperature oxidation of the material.

JEM-IT500 tungsten filament scanning electron microscope (SEM), manufactured by JEOL Ltd, Japan, was used to observe the microscopic morphology of the samples in detail. Phase analysis of the samples was performed using a transmission electron microscope (TEM), an energy dispersive spectrometer (EDS), and a Bruker D8 ADVANC X-ray diffractometer (XRD, Germany) within scanning range of 10°–90°. The acquired diffraction data were processed and analyzed using MDI JADE6.0 software. The grain size, orientation, and texture of the samples were analyzed by electron backscatter diffraction (EBSD) using a JSM-7900F field emission SEM manufactured by JEOL Ltd, Japan. The EBSD data were then processed using Channel 5 software. The mechanical properties of the samples were tested by

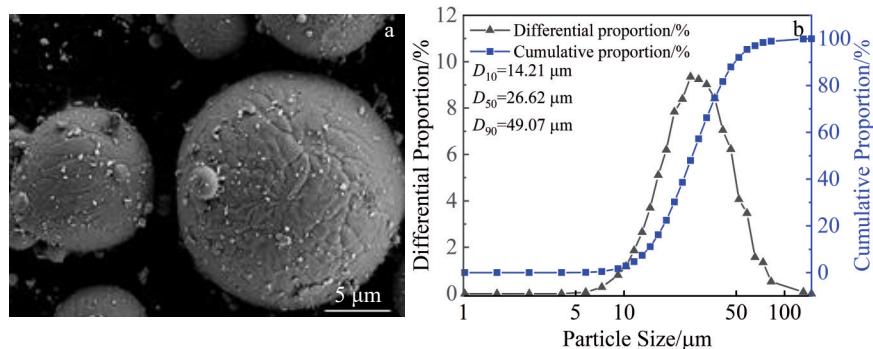


Fig.1 Morphology (a) and particle size distribution (b) of Er-TiB<sub>2</sub>/Al-Mn-Mg-Sc-Zr powder

tensile test on an Instron M4206 machine at a strain rate of 1 mm/min, according to GB/T 228.1-2010 standard.

### 3 Results and Discussion

#### 3.1 Phase analysis

Fig. 2 shows XRD patterns of the Al-Mn-Mg-Sc-Zr alloy and Er-TiB<sub>2</sub>/Al-Mn-Mg-Sc-Zr composite samples. It can be seen that the diffraction peak of Al (111) is significantly enhanced and becomes more intense than that of Al (200) after the addition of the reinforcing phases. This phenomenon occurs because TiB<sub>2</sub> has excellent thermal conductivity, enabling more efficient heat transfer from the center of the molten pool to its surrounding areas, making the temperature distribution in the molten pool more uniform, and thus optimizing the temperature gradient<sup>[14]</sup>. The optimized temperature gradient will guide the grain growth along the direction of heat flow. Since the (111) crystal plane is the most densely packed plane of Al atoms, under the appropriate temperature gradient conditions, Al atoms are more inclined to nucleate on the surface of TiB<sub>2</sub> particles with the (111) crystal plane, thereby increasing the nucleation rate of the (111) crystal plane. In terms of phase composition, only the fcc-Al phase is observed in the Al-Mn-Mg-Sc-Zr alloy, while the diffraction peak of Al<sub>3</sub>Sc is detected in the Er-TiB<sub>2</sub>/Al-Mn-Mg-Sc-Zr alloy, indicating that Al<sub>3</sub>Sc phase particles are generated in the composite sample. Additionally, TiB<sub>2</sub> is detected, and no other Ti-containing compounds are detected, indicating that TiB<sub>2</sub> does not react with the Al matrix. The diffraction peak of Al<sub>3</sub>Er is also detected in the composite, which indicates that

the reaction of Al and Er generates the Al<sub>3</sub>Er phase.

#### 3.2 Microstructure analysis

According to EBSD image of the Al-Mn-Mg-Sc-Zr alloy (Fig. 3a), it can be seen that the Al-Mn-Mg-Sc-Zr alloy prepared by LPBF is mainly composed of columnar grains growing along the forming direction, with small equiaxed grains present at the molten pool boundaries. Fig. 3b shows that the columnar grain region is significantly reduced after the addition of the reinforcing phases. The columnar crystal size decreases, and the equiaxed grain region increases significantly and becomes denser near the molten pool boundary. This is because TiB<sub>2</sub> and Er are uniformly distributed in the matrix to form a dispersed phase, which inhibits the grain growth<sup>[15]</sup>. Al<sub>3</sub>Er phase is formed in the LPBF process, exhibiting good interfacial coherence with the matrix. The melting point of Al<sub>3</sub>Er is much higher than that of the matrix powder, so Al<sub>3</sub>Er remains solid once formed, and acts as a heterogeneous nucleation site during melt crystallization, preventing the flow of the molten pool, hindering grain growth, and thus further enhancing the grain refinement effect.

It can be seen from Fig. 4a and 4b that both alloys exhibit a certain degree of texture. Specifically, the orientation strength of Al-Mn-Mg-Sc-Zr alloy in the [001] direction reaches 4.80 multiple of random distribution (mrd). After the addition of Er and TiB<sub>2</sub>, the maximum texture strength is lower than 2.99 mrd, the texture is weakened, and the polar intensity distribution is very uniform. This indicates that the addition of nanoparticles can eliminate the anisotropy caused by temperature gradient during the LPBF process, thereby enhancing the

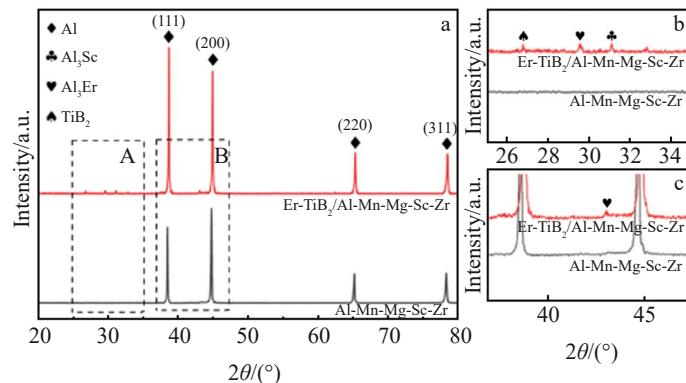


Fig.2 XRD patterns of formed samples: (a) overall patterns; (b) magnified patterns of area A; (c) magnified patterns of area B

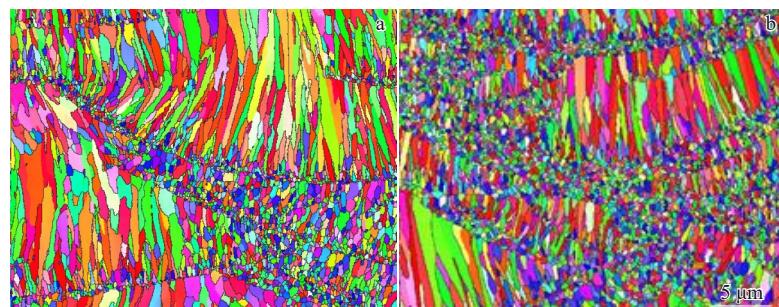


Fig.3 EBSD images of Al-Mn-Mg-Sc-Zr (a) and Er-TiB<sub>2</sub>/Al-Mn-Mg-Sc-Zr (b) samples

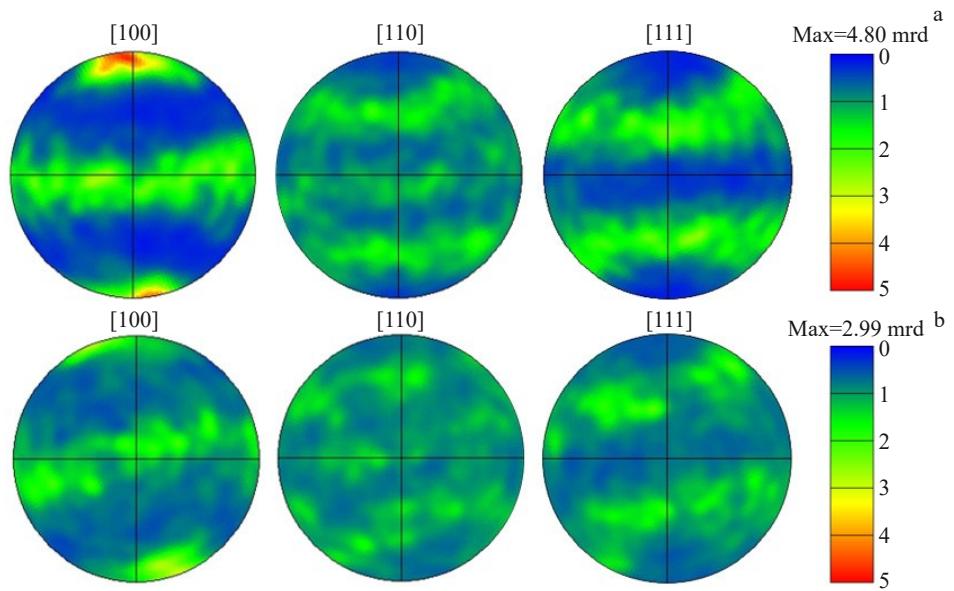


Fig.4 EBSD pole diagrams of Al-Mn-Mg-Sc-Zr (a) and Er-TiB<sub>2</sub>/Al-Mn-Mg-Sc-Zr (b) samples

multi-directional grain growth. Consequently, the material will display more uniform mechanical behavior under stress in dif-

ferent directions, and it will be less prone to brittle fracture<sup>[9]</sup>.

Fig. 5 shows TEM analysis results of 0.5wt% Er-1wt%

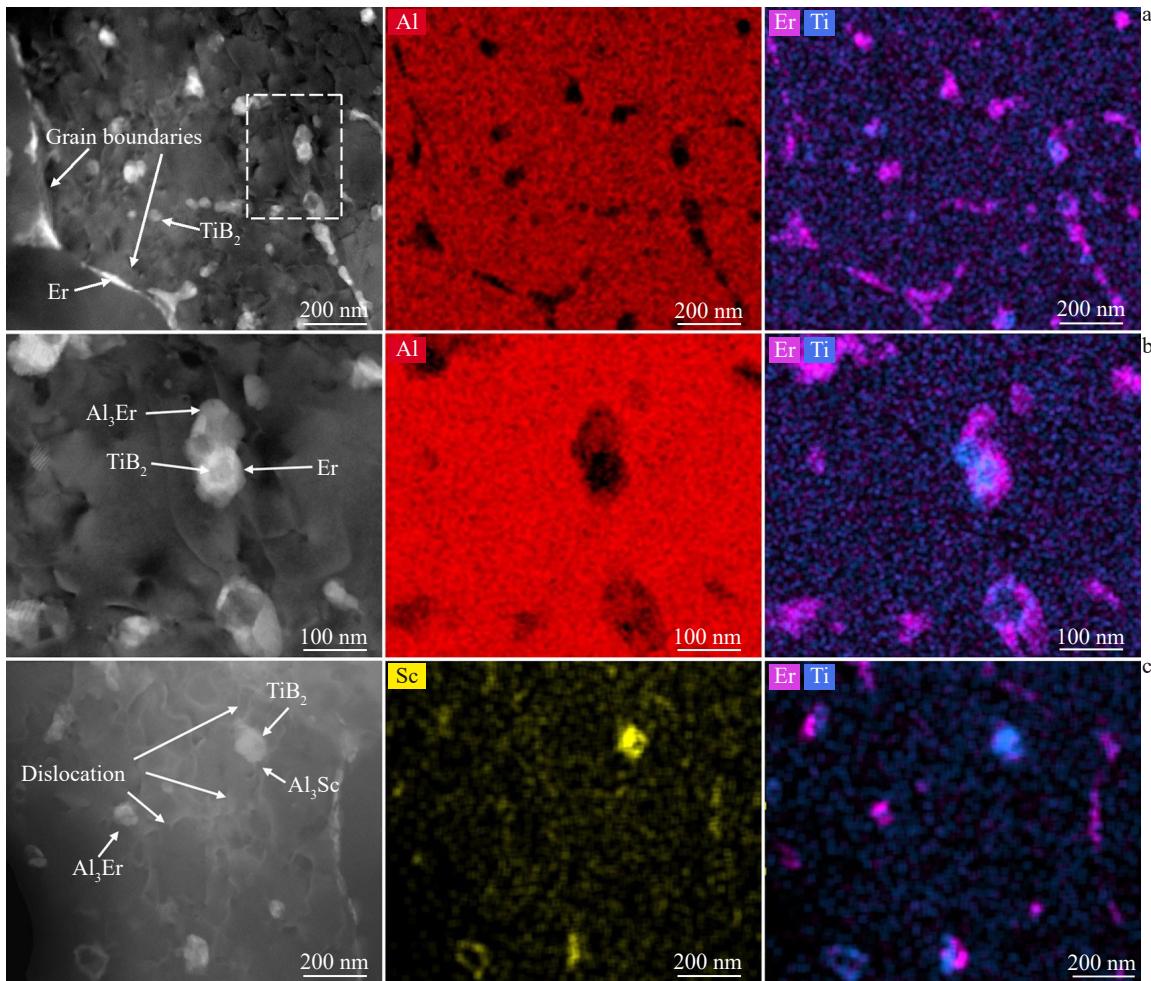


Fig.5 TEM images and corresponding EDS mappings of Al<sub>3</sub>Er, TiB<sub>2</sub>, and Er (a, c); magnified TEM image and corresponding EDS mappings of dotted area in Fig.5a (b)

$\text{TiB}_2/\text{Al}-\text{Mn}-\text{Mg}-\text{Sc}-\text{Zr}$  alloy. It can be seen from Fig. 5a that Er and  $\text{TiB}_2$  are distributed both inside the grains and at the grain boundaries. As shown in Fig. 5b, Er and  $\text{TiB}_2$  are co-distributed in the composite matrix, and  $\text{TiB}_2$  is attached to Er-containing structure, which has a stabilization effect on the Er while stabilizing the matrix structure. The spherical  $\text{Al}_3\text{Er}$  phases can be found, indicating that element Er exists in the matrix not only as individual atoms but also in the form of  $\text{Al}_3\text{Er}$  phase. In Fig. 5c, a large number of dislocations are accumulated around the  $\text{Al}_3\text{Er}$  phases. This phenomenon originates from the incorporation of element Er into the matrix during fabrication, coupled with thermal stress created inside the material during the LPBF process. When the  $\text{Al}_3\text{Er}$  phase is formed, the matrix around the  $\text{Al}_3\text{Er}$  phase experiences greater stress in the thermal cycling process due to the relatively large difference in thermal expansion coefficients between  $\text{Al}_3\text{Er}$  and the matrix. These thermal and residual stresses lead to the generation and proliferation of dislocations, which are accumulated around the  $\text{Al}_3\text{Er}$  phases. This is because of the dislocation-pinning effect of  $\text{Al}_3\text{Er}$  phase, making it difficult for the dislocation to escape from the  $\text{Al}_3\text{Er}$  phases. It is found that the element Sc forms  $\text{Al}_3\text{Sc}$  phases, with extensive dislocations accumulated around them during the formation process. This suggests that both  $\text{Al}_3\text{Sc}$  and  $\text{Al}_3\text{Er}$  can accumulate and pin dislocations. Furthermore, EDS mapping shows that  $\text{TiB}_2$  is distributed adjacently to Sc-rich region, indicating that  $\text{TiB}_2$  also has a certain stabilization effect on Sc.

Fig. 6 shows the favorable interface matching among Er,  $\text{TiB}_2$ , and the Al matrix. It can be seen from Fig. 6a that the Er reacts with Al matrix to form  $\text{Al}_3\text{Er}$ . The lattice constant of  $\text{Al}_3\text{Er}$  is 0.4215 nm, while that of the Al matrix is 0.4043 nm, resulting in a lattice mismatch of only 4.1%. As shown in Fig. 6d, the lattice constant of  $\text{TiB}_2$  is 0.3028 nm, with a lattice

mismatch of only 25.2% relative to the Al matrix. This demonstrates that Er and  $\text{TiB}_2$  can achieve favorable interface matching with the Al matrix through direct and indirect interactions, respectively. Such good interface compatibility enhances the heterogeneous nucleation efficiency of the reinforcing phases, especially for Er and  $\text{TiB}_2$ , which is crucial for improving material properties<sup>[16]</sup>. Additionally, the dispersed distribution of Er and  $\text{TiB}_2$  within the matrix complicates the interface network, thereby contributing to enhancement of the mechanical properties of the material.

### 3.3 Mechanical properties

Tensile tests were performed on the two alloys, and the results are shown in Fig. 7. Compared with  $\text{Al}-\text{Mn}-\text{Mg}-\text{Sc}-\text{Zr}$  alloy, the Er- $\text{TiB}_2/\text{Al}-\text{Mn}-\text{Mg}-\text{Sc}-\text{Zr}$  alloy exhibits a 13.4% increment in tensile strength and a 26.4% increment in elongation. This is because the addition of Er forms many nano-sized  $\text{Al}_3\text{Er}$  phases in the aluminum matrix. These fine compounds will hinder the movement of dislocations inside the aluminum matrix and induce the accumulation of dislocations around them, further restricting the movement of other dislocations. Additionally,  $\text{Al}_3\text{Er}$  also acts as a heterogeneous nucleating agent during the solidification process of the molten pool, promoting the nucleation and growth of  $\alpha$ -Al grains and thereby achieving grain refinement. The addition of  $\text{TiB}_2$  particles optimizes the temperature gradient, improves the laser absorption efficiency of aluminum matrix, promotes the precipitation of secondary phase, and facilitates the heterogeneous nucleation, collectively contributing to the formation of an excellent microstructure. Therefore, the mechanical properties are greatly improved. Based on this study, the properties of the current aluminum alloy materials are compared. The fracture morphologies of the two alloys are depicted in Fig. 8. The

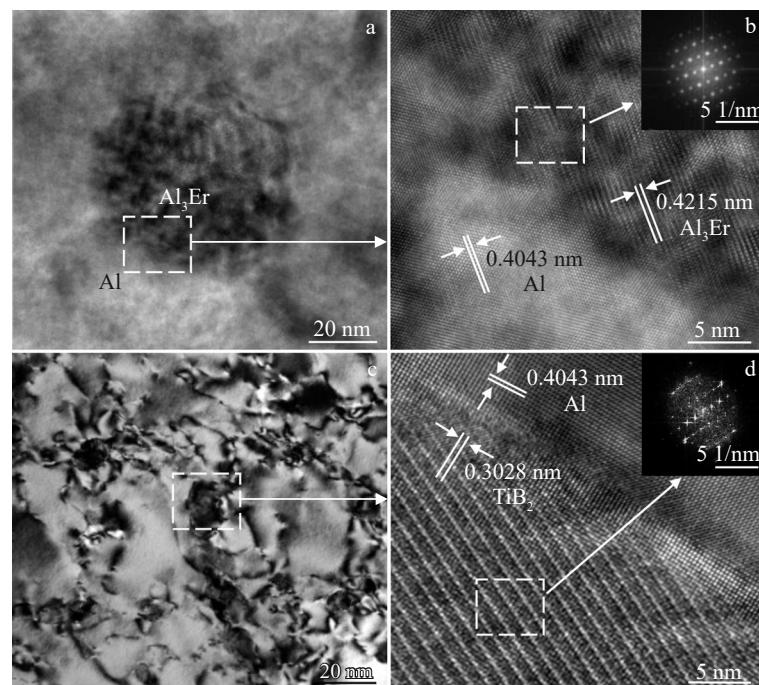


Fig.6 Lattice matching between Er and Al interface (a-b) as well as  $\text{TiB}_2$  and Al interface (c-d)

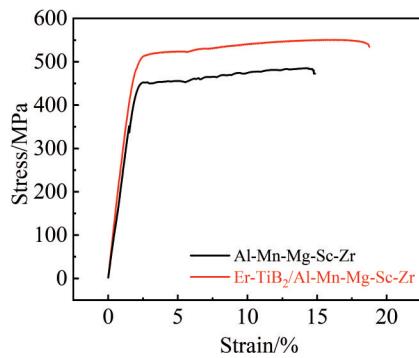


Fig.7 Stress-strain curves of Al-Mn-Mg-Sc-Zr and Er-TiB<sub>2</sub>/Al-Mn-Mg-Sc-Zr alloys formed by LPBF

fracture surface of the Al-Mn-Mg-Sc-Zr alloy exhibits a combination of cleavage and dimple features (Fig. 8a), indicating a mixed fracture mode encompassing both ductile and brittle characteristics. In the locally magnified image (Fig. 8b), the quantity and size of dimples are small, confirming a quasi-brittle fracture behavior with limited toughness. In contrast, the fracture surface of the Er-TiB<sub>2</sub>/Al-Mn-Mg-Sc-Zr alloy (Fig. 8c) is dominated by dimples without any cleavage facets. Higher-magnification observation

(Fig. 8d) reveals that these dimples are larger and deeper than those in the Al-Mn-Mg-Sc-Zr alloy, showing a ductile fracture mode, which reflects the excellent toughness of the alloy. These observations from the fracture morphology are consistent with the elongation results.

### 3.4 Strengthening mechanisms

#### (1) Hall-Petch strengthening

The Hall-Petch strengthening mechanism in materials is because the greater the number of grains, the more the grain boundaries. In the solidification process, adjacent grains will squeeze each other due to the interaction forces generated by grain growth, resulting in plastic deformation. Dislocations form inside the grains, slipping and multiplying along the crystal planes, and thereby creating a stress field near the grain boundaries. Grain refinement enhances material strength by increasing the number of grain boundaries, which effectively hinders dislocation motion<sup>[17]</sup>. The longitudinal sections of Al-Mn-Mg-Sc-Zr and Er-TiB<sub>2</sub>/Al-Mn-Mg-Sc-Zr alloys are shown in Fig.9.

#### (2) Solid solution strengthening

In the Al-Mn-Mg-Sc-Zr alloy, some Mn and Mg atoms will dissolve into the aluminum matrix, where they exert a pinning effect on dislocations, impeding the dislocation movement and thus increasing the alloy's strength. During the forming

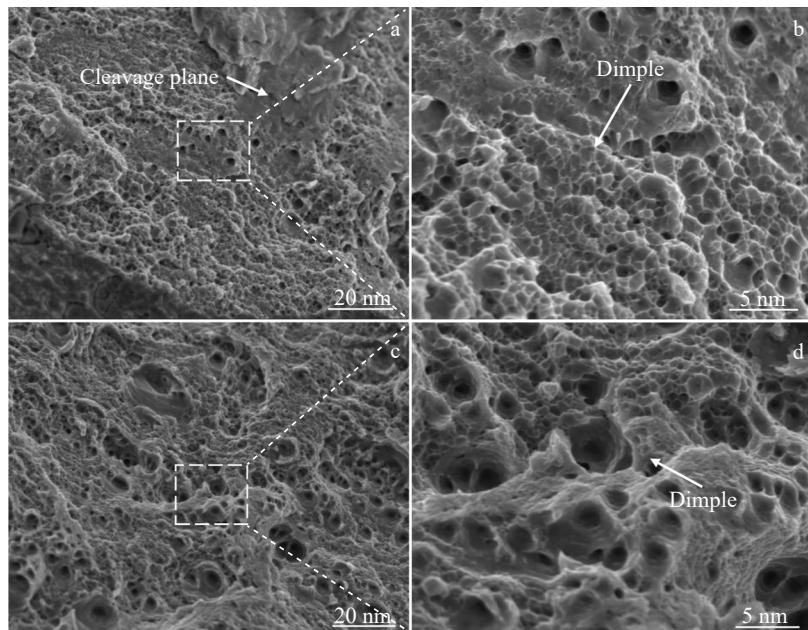


Fig.8 Fracture morphologies of Al-Mn-Mg-Sc-Zr (a-b) and Er-TiB<sub>2</sub>/Al-Mn-Mg-Sc-Zr (c-d) alloys formed by LPBF

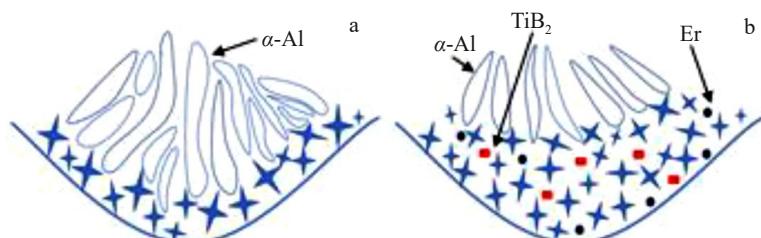


Fig.9 Schematics of longitudinal sections of Al-Mn-Mg-Sc-Zr (a) and Er-TiB<sub>2</sub>/Al-Mn-Mg-Sc-Zr (b) alloys formed by LPBF

process, the Sc, Er, and  $\text{TiB}_2$  will precipitate to form secondary phases that do not dissolve in the aluminum matrix. The content of the element Zr is very low, so only the solid solution strengthening of elements Mn and Mg is considered<sup>[18]</sup>.

### (3) Orowan strengthening

The obstruction of dislocation movement by dispersed nanoparticles in the material can enhance its strength, and this

strengthening mechanism is called dispersion strengthening (or Orowan strengthening). Due to the presence of Sc in both the base material and the composites, the Sc-containing phases are not the primary focus for the discussion of the Orowan strengthening mechanism in this research. Therefore, dispersion strengthening effect refers specifically to that caused by  $\text{TiB}_2$  and Er nanoparticles.  $\text{TiB}_2$  and Er achieve the strengthening effect by pinning dislocations within grains and at grain boundaries. The strengthening and toughening mechanisms of Er and  $\text{TiB}_2$  are shown in Fig.10<sup>[19]</sup>.

### (4) Load-bearing reinforcement

In the composite material, as depicted in Fig.11,  $\text{TiB}_2$  and Er are evenly distributed throughout the matrix. A strong interfacial bonding exists between the reinforcing phases and the matrix, enabling effective load transfer. Under tensile loading conditions, the softer Al matrix transfers load to the harder reinforcing phase particles, thereby enhancing the mechanical properties of the material. As shown in Fig.11, the cellular network structure is clear, continuous, and dense, with the reinforcing particles well-integrated into their surrounding matrix<sup>[9]</sup>.

### (5) Dislocation density reinforcement

As seen in Fig. 12, significant dislocation accumulation occurs in the Er- $\text{TiB}_2$ /Al-Mn-Mg-Sc-Zr alloy. The formation of dislocations in the alloy is attributed to not only the thermal stress induced by rapid cooling during the forming process, but also the mismatch in thermal expansion coefficients between the reinforcing phase and the matrix<sup>[20]</sup>.

## 3.5 Toughening mechanism

Based on the experimental results, the elongation at break

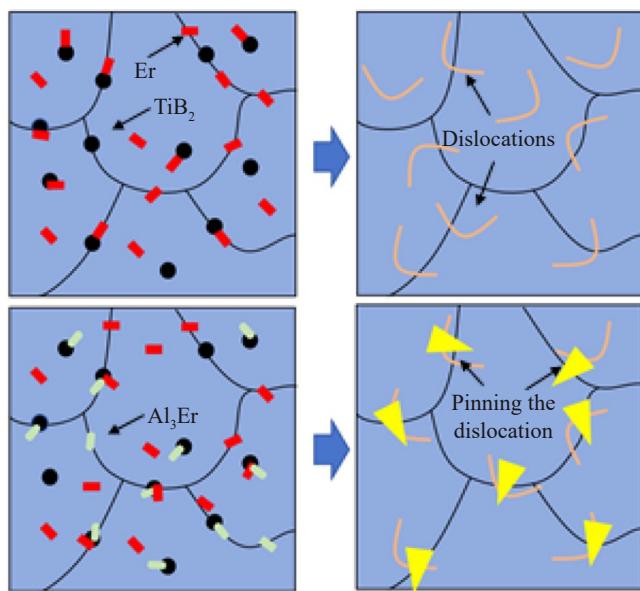


Fig.10 Strengthening and toughening mechanisms of Er- $\text{TiB}_2$ /Al-Mn-Mg-Sc-Zr alloy<sup>[19]</sup>

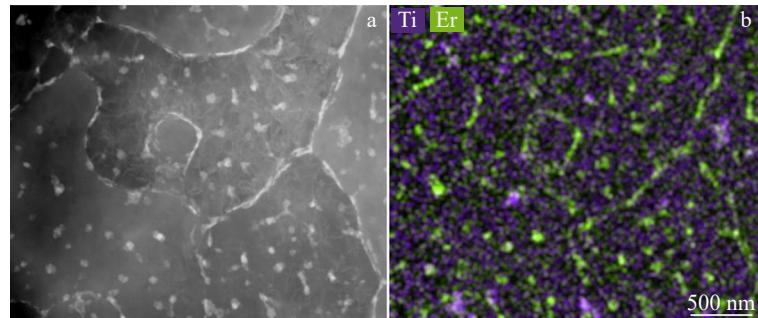


Fig.11 Interface bonding of Er- $\text{TiB}_2$ /Al-Mn-Mg-Sc-Zr alloy (a) and corresponding EDS mapping (b)

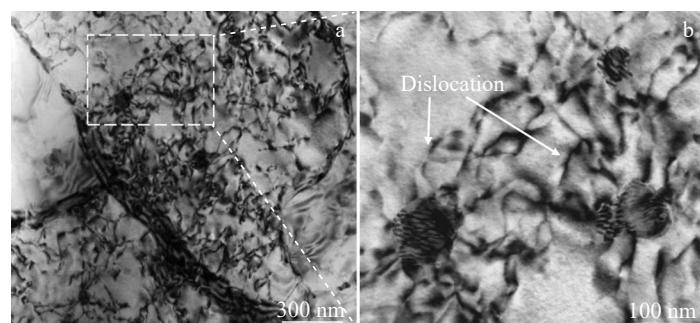


Fig.12 Internal dislocation distributions in grains of Er- $\text{TiB}_2$ /Al-Mn-Mg-Sc-Zr alloy

and the tensile strength of the dual-phase composite can reach a maximum of 18.7% and 550 MPa, respectively. Generally, the improvement in tensile strength is often accompanied by a sacrifice in toughness. For nanocomposites, however, the good interfacial bonding between the reinforcing phase and the matrix, as well as the grain refinement effect, can alleviate this trade-off.

Grain refinement can enhance ductility. As the grains are refined, columnar grains are gradually transformed into equiaxed grains. The total number of grains increases, and consequently, the grain boundaries increase as well. The stress concentration borne by individual grains is reduced and can be released over time, which decreases the possibility of crack nucleation. Randomly oriented equiaxed grains are beneficial to the improvement of plasticity. During the tensile deformation of columnar grains, local strain makes it difficult to release stress concentration, increasing the likelihood of crack generation. In contrast, non-oriented equiaxed grains are subjected to uniform stress, reducing local strain. Meanwhile, the deformation coordination ability is enhanced, and the relatively low stress concentration can be released over time, thereby achieving more uniform plastic deformation and increasing the elongation at fracture. Therefore, grain refinement is one of the key reasons for the increase in elongation at break.

The distribution of the reinforcing phase is another important factor influencing the elongation at break. Experiments show that the reinforcing phase particles are discretely distributed both intragranularly and intergranularly, and these particles are well bonded to the matrix. The lattice constants of  $\text{Al}_3\text{Er}$  and the aluminum matrix are highly similar, and there is good atomic coordination between them. When the material is subjected to external forces, load can be more effectively transferred from the matrix to the  $\text{Al}_3\text{Er}$  phase. When the lattice constants of the precipitated phase are close to those of the aluminum matrix, the interfacial lattice mismatch is very small. A small lattice mismatch means minimal stress generated due to lattice distortion at the interface between the  $\text{Al}_3\text{Er}$  phase and the aluminum matrix. Such a low-stress interface can effectively prevent the initiation and propagation of cracks at the interface. Since the generation and propagation of cracks typically need to overcome specific energy barriers, the low-stress interface reduces the driving force for crack formation, thereby improving the toughness of the material. While causing a dislocation-pinning effect within the microstructure, the reinforcing phases are closely combined with the matrix, hindering the grain growth. Most cracks are suppressed by the reinforcing phases, and the fracture is mainly dominated by dislocation activity. Therefore, the increase in elongation at break of nanocomposites can be attributed to the inhibitory effect of the reinforcing phases on cracks.

#### 4 Conclusions

1) The addition of Er in the Al-Mn-Mg-Sc-Zr alloy leads to the formation of fine  $\text{Al}_3\text{Er}$  phases through reaction with the

matrix. The addition of  $\text{TiB}_2$  can optimize the influence of temperature gradients in the forming process and promote the precipitation of more secondary phase.

2) In the Er-TiB<sub>2</sub>/Al-Mn-Mg-Sc-Zr alloy, the grains exhibit obvious refinement, and the texture strength tends to weaken. This indicates that the addition of reinforcing phase refines the grains and reduces the anisotropy caused by temperature gradients during the LPBF forming process. Consequently, the multidirectional growth of grains is enhanced.

3) The Er-TiB<sub>2</sub>/Al-Mn-Mg-Sc-Zr alloy achieves a tensile strength of 550 MPa and an elongation at break of 18.7%. Compared with those of the Al-Mn-Mg-Sc-Zr alloy, both the strength and plasticity are significantly improved. The strengthening mechanisms underlying the enhanced mechanical properties of the Er-TiB<sub>2</sub>/Al-Mn-Mg-Sc-Zr alloy are analyzed.

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## 添加 Er-TiB<sub>2</sub>双相纳米颗粒对激光粉末床熔融制备 Al-Mn-Mg-Sc-Zr 合金 强度-韧性的影响

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**摘要:** 采用双相协同增强的方法, 利用Er和TiB<sub>2</sub>的特有优势对激光粉末床熔融(LPBF)制备的Al-Mn-Mg-Sc-Zr合金进行强化。采用真空均质技术制备了0.5wt% Er-1wt% TiB<sub>2</sub>/Al-Mn-Mg-Sc-Zr纳米复合材料的球形粉末, LPBF工艺制备的样品密度达到99.8%。探讨了Er-TiB<sub>2</sub>的强化机理和增韧机理。结果表明, XRD分析存在Al<sub>3</sub>Er衍射峰, EBSD结果显示双相纳米颗粒添加后合金组织强度下降。添加的Er和TiB<sub>2</sub>纳米增强相在LPBF形成过程中起非均相核作用, 阻碍晶粒长大, 可有效细化晶粒。添加Er和TiB<sub>2</sub>双相纳米增强相后, LPBF成形样品的抗拉伸强度和延伸率分别达到550 MPa和18.7%, 相较基体材料提高13.4%和26.4%。

**关键词:** Al-Mn-Mg-Sc-Zr合金; 激光粉末床熔融; 纳米增强相; 协同增强

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