Laser additive manufacturing of intermetallic alloys for high-temperature applications

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Abstract

Intermetallic alloys like e.g. Iron-Aluminides, Titanium-Aluminides or Molybdenum-Silizides are prospective materials for high-temperature applications. For additive manufacturing (AM) intermetallic structural materials are particularly challenging due to their high melting points, oxygen susceptibility and low temperature brittleness. The feasibility of manufacturing intermetallic Mo-Si-B alloys with the laser additive manufacturing process of direct energy deposition (DED) is demonstrated and recent results in characterizing rapidly solidified material with respect to correlations between process, composition and microstructures are presented. The possibility to dope the material with Yttrium oxide (Y₂O₃) for dispersion is successfully demonstrated. Current challenges, e.g. homogenous distribution of alloying elements and applicability are addressed.

Keywords

Mo-Si-B alloys, laser additive manufacturing, high-temperature mechanical properties, oxide dispersion strengthening

1. Introduction

Hard to manufacture but high performing intermetallic alloys provide dormant potential to be used in demanding environments like e.g. high temperature, high stress or oxidizing atmospheres. Currently, Ni-based alloys are state of the art for applications in aerospace and energy production markets. Mo-Si-B alloys are promising substituents. Generally, Molybdenum-alloys are not oxidation resistant, but with high contents of Silicon and Boron, the formed intermetallic phases allow the formation of a thin oxide protection layer on the material surface [1]. Especially near-eutectic compositions in the Mo-Si-B system show a good oxidation resistance in temperature ranges between 600 and 1300°C [2]. Both high temperature strength and creep performance emphasize these materials’ significance as prospective structural material for e.g. turbine blades [3-7]. Reasons for the outstanding properties are the intermetallic phases Mo₃Si and Mo₅SiB₂, which finely distribute in the Molybdenum solid solution (Mo₃ss) matrix phase, improving creep and oxidation resistance to a high extend. A homogenous phase distribution is crucial to gain the desired properties. In [1, 8], Mo-9Si-8B (at%) containing about 50 vol % of matrix α-Mo phase and 50 vol % of Mo₃Si and Mo₅SiB₂ provides a good balance of mechanical properties. The same alloy produced by casting shows a yield strength of 700 MPa at 1200°C and a maximum compression strength of 750 MPa and thus exceeds high temperature properties of state of the art Ni-based alloys [13]. The coarse microstructure and phase distribution are recognized as a
limiting issue for the oxidation resistance. The high melting point of Mo-Si-B materials, dependent on the composition equal to or higher than 2000°C[2-5, 9], causes the fabrication mainly by powdermetallurgical methods consisting of multiple steps. The possibility to produce fine grained and dense samples is given, but so far only on laboratory scale. [2, 6, 8, 10-14]

When the application of innovative materials is hindered by the limits of conventional manufacturing, additive manufacturing (AM) can thrive. Materials with high brittleness are challenging candidates for machinability of any kind. Additive production techniques like the powder-bed based Laser Powder Bed Fusion (LPBF) or he powder-nozzle based Direct Energy Deposition (DED, also known as Laser Metal Deposition LMD) provide unique possibilities: The high intensity of the laser beam enables rapid melting even of high melting materials, and the solidification can be used to specifically design desired microstructures, e.g. directed or fine grained. The near net shape manufacturing allows part production without any or at least reduced post subtractive processing. Crack formation can be prevented by the usage of adequate preheating systems. These possibilities to manufacture demanding materials are up to know only used to a very small extend. Nevertheless, the high melting points are challenging for additive manufacturing processes as well. Following [16, 17], the high-temperature resistant eutectic Mo-17.5Si-8B and alloys of very similar compositions are quite a suitable candidates for AM. Besides excellent mechanical properties, the melting point of approx. 2000°C is the lowest in the Mo-Si-B system when meeting the eutectic composition. The printability of near-eutectic Mo-Si-B compositions with DED and the resulting microstructures were presented in [18]. Dense Mo-13.5Si-7.5B samples were consolidated by DED from pre-alloyed gas atomized powders. The microstructure was found to consist of Mo₆₃ matrix with embedded binary Mo₃Si-Mo₅SiB₂ and ternary Mo₆₃-Mo₃Si-Mo₅SiB₂ regions. The present work is based on this very first study and contains results of printing Mo-16.5Si-7.5B with DED. Aspects of the manufacturing process itself, microstructure characterization and creep properties are included.

Oxide dispersion strengthening (ODS) is known to increase the materials strength especially at high temperatures and improve creep behavior. The feasibility to apply ODS in additive manufacturing of Fe-, Ni-, Ti-based and y-TiAl alloys and the positive effect on mechanical properties has been demonstrated for various material systems [19-24]. It leads to increased hardness and strength. Nevertheless, the effect in structural intermetallic materials for high temperature applications is still widely unexplored. In this work, we attempt to apply ODS to the DED process of Mo-Si-B.

2. Materials and Experimental

Gas atomized powder material produced by Nanoval GmbH & Co. KG, Germany was used in the size of 45-90 µm. The compositions determined by EDX of the two batches used in this study were Mo-16.5Si-7.5B and Mo-13.5Si-7.5B (at.-%).

The DED process was performed using a machine with a 3-axis gantry system. A 2 kW diode laser system was used as the laser beam source. The powder was fed through a coaxial powder feed nozzle. The beam diameter was set to 0.6 mm. The parameters used in this study were a laser power of 70 W, a scanning velocity of 500 mm/min and a layer height of 0.15 mm. The preheating temperature was approximately 600°C, reached via an inductive preheating system. The movable coil mounted on a vertical access allowed constant conditions in the respective processing plane. Shielding to prevent oxidation was realized by use of an
Argon filled gas chamber and oxygen contents < 100 ppm. An additional inert gas stream of 15 l/min was chosen to additionally protect the melt pool locally.

Oxide dispersion strengthened samples were produced in the same manner using Mo-Si-B-powder doped with nano-sized (approx. 45 nm) \( \text{Y}_2\text{O}_3 \) particles (abcr GmbH, Germany) via a shortened milling process in a planetary mill. The milling procedure was performed in a planetary mill (Pulverisette 4 Classic line, Fritsch GmbH, Germany) with zirconia grinding bowls and yttrium stabilized zirconia (YSZ) grinding balls (diameter: 5 mm). The process included three milling intervals of 20 min each followed by pausing for 10 min to cool down the powder material. The milling atmosphere was argon.

Image analysis of the DED microstructure was performed using a SEM LEO 1455 EP (Carl Zeiss AG, Germany). The relative density of specimens was determined by computer-aided light microscopical analysis. EBSD phase analysis was performed at FZ Jülich by E. Wessel. Micro-hardness was tested as HV0.3 using a semi-automatic Vickers tester (LECO Instrumente GmbH, Germany) with a test load of 300 g and a testing time of 20 s. The test indentation distance was 0.5 mm. Creep tests were performed with a Zwick Roell Z100 machine including a Maytec furnace in pure argon atmosphere at 1093°C.

3. Results

3.1 DED of Mo-Si-B

Typical defects for LMD of intermetallics are cracking, ablation as a consequence of internal stresses and porosity. The brittleness of Mo-Si-B is high due to present \( \text{Ti}_2(\text{Mo}_5\text{SiB}_2) \) and \( \text{Mo}_3\text{Si} \) phases at room temperature. Therefore, preheating and slow cooling rates are required after the process to prevent thermally induced cracking. The final deposition strategy elaborated includes bidirectional scanning and slow cooling by a stepwise reduction of the induction powder over several minutes. Nevertheless, all samples detach from the substrate during the cooling stage. The size of the produced samples is approx. 20 mm length, 4 mm width and 10 mm height (Fig. 1, A). Pore formation could not be suppressed completely, but significantly reduced by iterative adapting of the scanning strategy. The relative density determined by optical measurements in polished cross sections (Fig. 1, B) is >98%.

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**Fig. 1:** Additively manufactured (DED) Mo-Si-B specimen: a) macro photograph of a sample b) LOM micrograph of the polished cross section c) LOM micrograph (etched) with drawn layers

SEM micrographs show homogeneous dendritic structure over the entire height with refined features in the solidification start region of each track (Fig. 1, C) triggered by different
solidification speeds which are typical for AM processes. These regions are typically richer in Molybdenum (MoSS). EBSD analysis reveals the phases MoSS, Mo3Si and T2 (Fig. 2). While T2 and Mo3Si are known to improve hardness, creep resistance and oxidation resistance the MoSS phase increases the room temperature toughness.

![EBSD Analysis](image)

**Fig. 2**: EBSD of Mo-Si-B AM sample having a three-phase microstructure

Hardness measurements result in values of 1300-1400 HV0.3, which is in the expected range of the material [5, 19].

### 3.2 Creep behaviour of DED manufactured Mo-16.5Si-7.5B

A deeper insight into the mechanical behaviour of the DED Mo-Si-B alloy is given by tests of the creep rate of the material. The creep properties of the DED Mo-16.5Si-7.5B (tested in as-built conditions) are presented in Fig. 3.
The alloy shows a creep behavior located between a powder metallurgy (PM) Mo-9Si-8B alloy, providing a microstructure of intermetallic Mo₃Si and Mo₅SiB₂ phases which are embedded in a continuous Mo₃ss network (Mo₃ss > 50%) [13], and the chemically very similar zone-molten (ZM) Mo-17.5Si-8B alloy. The chemical similarity of both alloys (DED Mo-16.5Si-7.5B, ZM Mo-17.5Si-8B) leads to comparable phase distributions, with microstructures consisting of more than 50% of intermetallic phases (Mo₃Si and Mo₅SiB₂) due to the higher content of Si and B. A difference can be found in the grain size of the alloys, where the creep behavior is improved due to the coarser grained microstructure in the ZM Mo-17.5Si-8B alloy [17]. On the contrary, the higher proportion of Mo₃ss phase in the PM Mo-9Si-8B lowers the creep resistance in comparison to the DED Mo-16.5Si-7.5B and ZM Mo-17.5Si-8B alloy, since the Mo₃ss phase provides the weakest creep strength of the three phases (Mo₃ss, Mo₃Si, Mo₅SiB₂) [25]. Relative to DED Mo-16.5Si-7.5B the creep response of the CMSX-4 alloy can be found in the same order of magnitude. The creep behavior of the DED manufactured Mo-16.5Si-7.5 alloy can be classified as competitive compared to other Mo-Si-B alloys having a three-phase microstructure and the single-crystalline Ni-based superalloy CMSX-4.

3.3 Oxide dispersion strengthening of DED manufactured Mo-13.5Si-7.5B

In Fig. 4 A-B, the successful attempt to dope DED Mo-13.5Si-7.5B (composition determined by EDX) with Y₂O₃ is presented. The sample is free of cracks after the build-up. Nevertheless, by reasons of mechanical preparation cracking occurs afterwards, visible in the cross section (Fig. 4, B). The sample shows an identical grain structure with and without dispersed oxide particles.
**Fig. 4:** Additively manufactured (DED), Y\textsubscript{2}O\textsubscript{3}-doped Mo-Si-B specimen: a) macro photograph of a sample b) cross section c) LOM micrograph

For comparison reasons, one reference sample without additional particles was produced out of the same batch (Fig. 5, B). SEM reveals finely dispersed Y\textsubscript{2}O\textsubscript{3}-particles (white spots) in the matrix phase of the doped sample (Fig. 5, A). The size of the particles is up to 500 nm. The distribution however is not homogeneous. Clustered regions are visible as well as individual particles, following no obvious rule of accumulation. Nevertheless, they seem to segregate at phase boundaries, which has to be characterized in depth by EBSD measurements.

**Fig. 5:** Additively manufactured (DED) Mo-Si-B specimen: SEM micrographs A) Y\textsubscript{2}O\textsubscript{3}-doped B) non-doped

When testing the samples against each other, the hardness is measured to be 1370±100 HV0.3 compared to 1330±60 HV0.3 in the non-doped sample. Therefore, no significant increase in hardness is found. Most likely the hardness induced by the intermetallic phases which predominate the microstructure is already so high, that under consideration of measurement deviations the additional gain is unrecognizable. Further investigations will include detailed analysis of the distribution of the particles and determination of creep behaviour in comparison with non-doped Mo-Si-B.
4. Summary and Conclusions

In this work, the feasibility to build crack free bulk DED-samples with a high density out of the intermetallic Mo-16.5Si-7.5B alloy is demonstrated. Bulk crack free samples can be built out of Mo-16.5Si-7.5B gas atomized powder. The density is in the range of 98-99%. The microstructure consists of the expected phases Mo\(_5\)Si, Mo\(_3\)Si and T\(_2\) with more than 50% intermetallic phase fraction. Furthermore, in accordance with literature and observed microstructure, the creep behaviour was tested to be in the range of a state of the art Ni-based super alloy. Oxide dispersion strengthening in order to increase the creep resistance was tested successfully. A first trial to dope Mo-13.5Si-7.5B with Yttrium-oxides was successfully performed, however leading to irregular distribution. Therefore further process adjustments, as well as creep tests of the material, are part of current research.

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