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Assessing the efficacy of several impact-based mechanical techniques on fatigue behavior of additive manufactured AlSi10Mg



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ABSTRACT

Post-processing methods to reduce issues associated with the presence of internal and external anomalies are often necessary for obtaining adequate structural performance for additively manufactured products. However, the choice of the proper post-treatment and the corresponding parameters is still a challenge requiring adaption to the material type, geometry, size and undeniably costs. In this study, four different pure impact-based mechanical operations involving ultrasonic nanocrystal surface modification (UNSM), ultrasonic shot peening (USSP), severe shot peening (SSP), and severe vibratory peening (SVP) to investigate their efficacy on the fatigue behavior of hourglass AlSi10Mg specimens manufactured via laser powder bed fusion (LPBF) were considered. Experimental characterizations involving microstructural approach, porosity level and surface texture, hardness and residual stresses measurements, as well as tensile and fatigue testing, were conducted. The results exhibited considerable improvement in mechanical/physical performances leading to substantially enhanced fatigue performance of the mechanically treated specimens. Based on a cost-performance analysis, it was found that UNSM, while having reasonable cost, presented considerable improvement on fatigue behavior.

1. Introduction

Laser powder bed fusion (LPBF), a complex and influential additive manufacturing (AM) process, provides the manufactured parts with intrinsic features and dimensions. However, due to the complex manufacturing process including layer by layer melting and rapid solidification (high cooling rates of $\sim 10^6$ K/s [1,2]), these materials are majorly affected by multiple volumetric and surface anomalies [3–5]. In as-built configuration, these materials represent inhomogeneous microstructure [6], several types of volumetric defects (due to gas entrapment or joining failure, etc.) [7,8], hazardous residual stress [9, 10], and surface macro/micro distortions [11]. Completely unmelted powder aggregates, spatters and powder metal defects are known as the main sources of surface imperfections [12–15]. These defects potentially lower the performance of LPBF materials by means of wear, scratch and corrosion resistance as well as their fatigue behavior [16–21]. In particular, surface imperfections behaving like local stress concentration

regions lead to early crack nucleation and thus fatigue failure when subjected to cyclic loading [22,23]. Therefore, post-processing methods are being widely evaluated in the AM sector as they are essential to address these challenges [24,25].

Heat treatment (HT) can modulate some of the internal defects; in particular, homogenizing the microstructures and removing its anisotropy, besides releasing the intrinsic tensile residual stresses (TRS). Ductility of the LPBF materials can also be improved with HT [26,27], which may lead to enhanced fatigue behavior, specifically in the low-cycle fatigue regime [28,29]. Hot isostatic pressing (HIP) is also a commonly used post-treatment that besides homogenizing the microstructure of AM materials, can also reduce their porosity. This process simultaneously subjects the component to elevated temperature and isostatic gas pressure in a high-pressure containment vessel [30–32].

Among non-subtractive post-processing methods sourced to address the surface inadequacy, impact-based surface treatments have gained considerable attention. This category can include a wide range of treatments, such as shot peening (SP) [33–35], ultrasonic shot peening

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Abbrev	viations	LSP	Laser Shock Peening
		Μ	Machining
AM	Additive Manufacturing	Р	Polishing
CRS	Compressive Residual Stresses	SB	Sand Blasting
CSP	Conventional Shot Peening	SSP	Severe Shot Peening
ECP	Electro-Chemical Polishing	SVP	Severe Vibratory Peening
EL	Elongation To Fracture	SP	Shot Peening
HT	Heat Treatment	TRS	Tensile Residual Stresses
HIP	Hot Isostatic Pressure	UTS	Ultimate Tensile Strength
IPF	Inverse Pole Figure	UNSM	Ultrasonic Nanocrystal Surface Modification
KAM	Kernel Average Misorientation	USSP	Ultrasonic Shot Peening
LP	Laser Polishing	VF	Vibratory Finishing
LPBF/L	B-PBF Laser Powder Bed Fusion	YS	Yield Strength
LRM	Laser Re-Melting		

(USP) [36], severe vibratory peening (SVP) [37] and ultrasonic nanocrystal surface modification (UNSM) [38–40]. These operations all follow the same concept of mechanically inducing surface plastic deformation; they remove the surface irregularities, homogenize the surface morphology and can induce grain refinement and high compressive residual stresses (CRS) on (and in the vicinity of) the surface [41–45]. However, despite the common physics, these treatments differ in the type of the equipment, have dissimilarities in the range of treatable geometries and require different exposure times.

AlSi10Mg parts can be fabricated by LPBF at bulk volume density without solidification cracks owing to near eutectic composition of this alloy. The occurrence of quite small cellular structure during LPBF process in this alloy offers acceptable combination of ductility and strength [46–50]. Numerous studies have revealed the effect of various post-treatments and mechanical performance of AlSi10Mg manufactured by LPBF. The list of the applied surface post-processing methods (with and without material removal) and the material properties involving surface roughness, residual stress, hardness, corrosion and fatigue behavior are presented in Table 1.

As shown in Table 1, the post-treatments can effectively decrease the surface roughness or modify the surface morphology, enhance the mechanical performance, such as wear and corrosion resistance and improve fatigue behavior. For example, application of USP on LPBF AlSi10Mg with a 17 kHz frequency and a power of 1000 W and an amplitude of 80 μ m resulted in remarkable porosity reduction and surface hardening as well as induced high surface CRS compared to the prior stress state, leading to notable corrosion resistance improvement [54]. In another study, SP process applied by Almen intensity, 10A [0.001 inch] and 100% coverage via steel media, led to surface roughness reduction for LPBF AlSi10Mg specimens decreasing R_a from 9.0 to 4.5 μ m [58]. The combined effect of reduced surface morphology, surface hardening as well as maximum in depth CRS up to -155 MPa (on the contrary as-built state has 70 MPa tensile stresses), led to remarkable fatigue strength enhancement.

In our prior investigations, we surveyed the impact of multiple post operations of conventional shot peening (CSP) [60,61], SVP [37] and LSP [62] combined with HT on fatigue performance of notched LPBF AlSi10Mg specimens. The results demonstrated noteworthy fatigue life enhancement up to 233, 235 and 243 times better against AB specimen at a stress of 110 MPa via hybrid treatments of HT + CSP, HT + SVP and HT + LSP, respectively. Herein, differentiating from the previous studies, we specifically focus on different peening-based technologies to evaluate their effects when applied at the same energy level, to compare their efficacy, advantages, and limitations. In particular, severe shot peening (SSP), USP, SVP and UNSM are applied individually and combined with HT on LPBF AlSi10Mg specimens. Detailed experiments involving monotonic tensile tests, microstructural characterization, surface morphology and roughness measurements, porosity, hardening and internal residual stresses evaluations and fatigue behavior determination were conducted on all specimens to assess the efficacy of the applied surface post-treatments.

2. Experimental procedures

2.1. Material and specimens preparation

AlSi10Mg powder produced by gas atomization with the nominal size interval of 20–70 μ m and average diameter of 50 μ m was utilized for manufacturing of LPBF specimens as demonstrated in Fig. 1a. Table 2 presents powder chemical composition. The specimens were produced vertically as illustrated in Fig. 1b via an SLM 500 HL equipment by using Yttrium fiber lasers by a spot diameter of 78 µm and a power of 350 W. The rotation degree of 67° was utilized between the adjacent layers and contouring as depicted in Fig. 1c. The schematic view of the cylindrical hourglass fatigue test specimens is shown in Fig. 1d. In addition, miniature flat tensile test specimens were prepared later by wire cutting out of the fatigue specimens as shown in Fig. 1e. The shape of the miniature specimens follows the specifications presented by Cosa et al. [63] (Fig. 1f). Fig. 1g and h shows the schematic illustration of the manufacturing process of the specimens, layer-by-layer via LPBF technology and the variety of their possible internal and surface anomalies, respectively.

2.2. Post-processing methods

Different impact-based mechanical surface operations involving SSP, USP, SVP and UNSM as well as their combination with HT were applied to hourglass LPBF AlSi10Mg specimens. T6 type thermal treatment was applied to the specimens (merely half of the whole) aiming to homogenize the overall structure and vanish the TRS of the as-built state. T6 HT was applied according to durations and temperature zones demonstrated in the prior work [60], with an initial temperature towards 520 °C, solution based HT for 1 h, water quench for 1 h, subsequent to reaching temperature to 160 °C and aging for 6 h at 160 °C, followed by water quenching.

During the SSP, the surface of the specimen is exposed to shot flow either using high values of Almen intensity (index of deformation capability) or surface coverage (associated with peening duration) as presented in Fig. 2a. In the USP treatment, larger impacting media under an ultrasonic frequency in a peening chamber as depicted in Fig. 2b. The SVP treatment, on the other hand, utilizes high-frequency vibration and thus induces severe plastic deformation as presented in Fig. 2c. Finally, during the UNSM treatment, high frequency elastic waves generated by ultrasonic generator are amplified by a WC horn, which is impacted to the surface of specimen using a defined scanning direction as shown in Fig. 2d. Here, to compare the impact of these mechanical operations on Post-processing

method Laser polishing

(LP)

Laser shock

peening (LSP)

Vibratory finishing

Ultrasonic shot

peening (USP)

Sand blasting (SB),

VF, Machining

Polishing (P)

Laser re-melting

(LRM)

M, P, electro-

chemical

(SP)

polishing (ECP),

Shot peening

(M) and

(VF)

Table 1

Summary of typical post-processing methods applied on LPBF AlSi specimens.

Category

operation

operation

operation

operation

No additional removing

No additional removing

Additional removing

No additional removing

No additional removing

operation for VF, M and

No additional removing

Additional removing

and no additional

SP

operation for M, P, ECP

removing operation for

operation

operation for SB and

additional removing

	Table 1 (continued)	
i10Mg	Post-processing method	Category ^a	Remarks
Ref.			Surface roug
			$SP + P: R_a =$
[51]			Surface rough
[]			$SP + ECP: R_a$
			 Increasing
			-++1- (-

Ref.

		method			
Remarks	Ref.			Surface roughness after $SP+P:R_a=2.15~\mu m$	
Surface roughness reduction	[51]			Surface roughness after SP + ECP: $R_a = 1.77\ \mu m$	
Surface roughness of asbuilt state: $R_a^{\ b} = 8 \ \mu m$				 Increasing fatigue strength (smooth sample) 	
Surface roughness of LP: $R_a = 0.25 \ \mu m$				Fatigue strength of as-	
 Sub-surface porosity reduction 	[52]			built state: 75 MPa Fatigue strength after SP:	
Porosity in as-built state: ~0.87%		SP and SB	No additional removing	 105 MPa Surface roughness reduction 	[58]
Porosity after LSP: $\sim 0.02\%$			operation	Surface roughness of as-	
Surface roughness reduction	[53]			built state: $R_a = 9.1 \ \mu m$ Surface roughness after M: $R_a = 6.5 \ \mu m$	
Surface roughness of asbuilt state: $S_a = 14 \ \mu m$				M: $R_a = 6.5 \ \mu m$ Surface roughness after SB: $R_a = 4.8 \ \mu m$	
Surface roughness of VF: $S_a = 4 \ \mu m$	[[4]			 Increasing surface hardness 	
Increasing surface hardness	[54]			Surface microhardness in as-built state: 131 Hv	
Surface microhardness of as-built state: 116 Hv Surface microhardness of				Surface microhardness after SP: 157 Hv	
USP: 142 Hv • Inducing CRS				Surface microhardness after SB: 145 Hv	
Surface residual stress of as-built state: 38 MPa				 Inducing CRS Surface residual stress in 	
Surface residual stress of USP: –155 MPa				as-built state: 49 MPa Surface residual stress	
 Increasing corrosion resistance 				after SP: –80 MPa Surface residual stress	
 Surface roughness reduction 	[55]			after SB: -85 MPa • Increasing fatigue	
Surface roughness of asbuilt state: $S_a=15.4\ \mu m$				strength (Smooth sample)	
Surface roughness after SB: $S_a = 8.3 \ \mu m$				Fatigue strength of as- built state: 50 MPa	
Surface roughness after VF: $S_a=2.3\ \mu m$				Fatigue strength after SP: 185 MPa	
Surface roughness after $M + P: S_a = 0.5 \ \mu m$		M	A 1 114 1	Fatigue strength after SB: 173 MPa	[[0]
 Increasing fatigue strength 		M, SP	Additional removing operation M and no	 Surface roughness reduction Surface roughness of as- 	[59]
Fatigue strength in as- built state: 50–62 MPa			additional removing operation for SP	built state: $R_a = 6.89 \ \mu m$ Surface roughness after	
Fatigue strength after SB: 152.5 MPa				SP: $R_a = 5.82 \ \mu m$ Surface roughness after M: R_a	
Fatigue strength after VF: 95 MPa				0.22 µm Surface roughness after	
Fatigue strength after M + P: 194 MPa • Surface roughness	[56]			 SP: R_a = 2.05 μm Increasing surface 	
reduction Surface roughness in as-	[30]			hardness Surface microhardness in	
built state: $R_a = 20.6 \ \mu m$ Surface roughness after				as-built state: 120 Hv Surface microhardness	
 SB: R_a = 10.8 μm Porosity reduction 				after SP: 154 Hv Surface microhardness	
Porosity in as-built state: 0.15 vol%				after M: 128 Hv Surface microhardness	
Porosity after LRM: 0.09 vol%				after M + SP: 145 HvInducing CRS	
 Surface roughness reduction 	[57]			Surface residual stress in as-built state: 11 MPa	
Surface roughness in asbuilt state: $R_a = 8.92 \ \mu m$				Surface residual stress after SP: -152 MPa	
Surface roughness after M: $R_a = 1.75 \ \mu m$				Surface residual stress after M: –79 MPa Surface residual stress	
Surface roughness after $M + P: R_a = 0.59 \ \mu m$		SP	No additional removing	 after M + SP: -140 MPa Surface morphology 	[60]
Surface roughness after SP: $R_a = 4.60 \ \mu m$		Jf .	operation	 Surface morphology modification 	[00]

(continued on next page)

Table 1 (continued)

Post-processing method	Category ^a	Remarks	Ref.
Chemical polishing (CP) and ECP and SP	Additional removing operation for CP and ECP and no additional removing operation for SP	• Increasing surface hardness Surface microhardness in as-built state: 121 Hv Surface microhardness after SP: 155 Hv • Inducing CRS Surface residual stress in as-built state: -11 MPa Surface residual stress in as-built state: -11 MPa Surface residual stress after SP: -83 MPa • Increasing fatigue strength (Notched sample) Fatigue strength in as-built state: 6 MPa Fatigue strength in as-built state: 6 MPa • Surface roughness reduction Surface roughness after CP: $R_a = 3.19 \ \mu m$ Surface roughness after CP: $R_a = 2.96 \ \mu m$ Surface roughness after SP: $R_a = 4.61 \ \mu m$ Surface roughness after SP: $R_a = 4.61 \ \mu m$ Surface roughness after SP: $R_a = 3.29 \ \mu m$ • Increasing fatigue life in as-built state: 1.26×10^4 Fatigue life after CP: 1.41×10^5 Fatigue life after SP: 1.19×10^6 Fatigue life after SP: 1.19×10^6	[61]

***Sa: arithmetical mean height of an area.

^a Based on the classification of post operations of AM products presented in Ref. [25].

^b R_a: arithmetical mean of height.

specimens, equal level of kinetic energy was used for all operations by specifying the same range of Almen intensity that is majorly employed among SP operations. Each treatment was operated on an Almen strip A to reach Almen intensities of 10–12A [0.001 inch], by measuring the arc heights of the strips. Fig. 2e presents the measured arc heights in different exposure times for each considered treatment. The detail of parameters of the applied treatments are listed in Tables 3–6. Overall, a total of 10 sets of specimens involving AB, AB + SSP, AB + USP, AB + SVP, AB + UNSM, AB + HT, AB + HT + SSP, AB + HT + USP, AB + HT + SVP and AB + HT + UNSM were prepared in this study to examine the mere and hybrid influence of several post-processing treatments.

2.3. Experimental characterizations

Microstructural analyses were performed by using an optical microscope (Nikon Eclipse LV150NL) and a Zeiss Sigma 500 VP scanning electron microscope (SEM) equipped with electron back-scatter diffraction (EBSD) and energy dispersive spectrometry (EDS) detectors. EBSD results were processed by AZtecHKL software. Considering build direction, the specimens were cut through cross-sections and were

etched using Keller's reagent for 20 s. In addition, XRD analyses were carried out using Rigaku SmartLab XRD equipment with CuK α radiation, with around 40 kV voltage, and 30 mA current. The average width of at half height (FWHM) parameter was evaluated using Gaussian approach. Afterwards, the crystallite size was calculated via Williamson-Hall relation described in Ref. [64].

Porosity measurments were conducted via X-ray computed tomography (Nikon XTH 225ST micro-CT) at 190 kV voltage, 40 µA current and 3900 ms exposure time. Specimens' surface morphology was analyzed using a Zeiss EVO50 SEM. Roughness investigation around the highest stress concentration zone (middle of the specimens) was conducted via Alicona Infinite Focus microscope. ISO 25178-2 standard [60, 65] was followed to obtain the values of roughness parameters of arithmetic roughness average of the surface (Ra) and the root mean square of the surface roughness (R_q). Microhardness tests were carried out on polished longitudinal yz-plane (considering z axis for build direction) via Leica WMHT30A equipment (microvickers) with the load of 0.25 N and the load application duration of 15 s considering 50 μ m spacing between indentations. Three paths were used on each specimen from surface through 740 µm interior. X-ray diffraction (XRD) was performed to obtain the variation of residual stress using AST X-Stress 3000 CrKa radiated XRD equipment. Measurements were performed from surface towards the inside of the specimen for a depth of $700 \,\mu m$ to assess the perpendicular path with respect to the build direction. ECP was employed for thinner layer removal in each step using acetic (94%) and perchloric acid (6%) solution at 40 V.

The tensile tests were performed via electromechanical universal testing equipment of MTS QTEST/100 at an operation speed of 1.5 mm/ min at 25 °C using a miniature MTS 632.29 extensometer with the gage length of 5 mm. The utilized extensometer had the strain measurement capacity of -10%-30%, which was extensively convenient for the subsize tensile test specimens for dilatometry. Fatigue behavior of specimens was assessed using an Italsigma rotating bending fatigue test equipment at a constant stress of 110 MPa, the stress ratio of R = -1 and the angular speed of 2500 rpm. The run-out limit was determined as 1.5×10^7 cycles. Three specimens were tested for each condition and the average fatigue lives were used for all comparisons. In addition, fractography assessment was conducted to the failed specimen by using Zeiss EVO50 SEM.

3. Results

3.1. Tensile behavior

Fig. 3a and b shows the comparison of stress-strain curves for all the specimens. Histograms in Fig. 3c and d shows the average data of the three tested specimens. Tensile test outputs revealed a considerable distinction between the quasi-static tensile properties pre and post HT (AB vs. AB + HT) demonstrating a notable enhancement of elongation to fracture (EL) along with reduction of yield (YS) and ultimate tensile strength (UTS). EL was raised from 3% up to 14% after HT, however, YS and UTS diminished from 273 to 385 MPa down to 201 and 264 MPa, respectively.

The mechanical surface treatments, on the other hand, increased both YS and UTS while slightly diminishing the EL, as summarized in Table 7. As similarly reported for conventional and AM materials subjected to mechanical surface treatments, changes in YS were more significant than that of the UTS [66–70]. This strengthening effect induced by the peening based surface treatments can be associated with surface grain refinement and the presence of CRS [71].

3.2. Microstructural analyses

Fig. 4a depicts the optical microscopy (OM) investigations of AB/AB + HT specimens in both transverse and longitudinal cross-sections. In the AB specimen inhomogeneous microstructure can be observed with

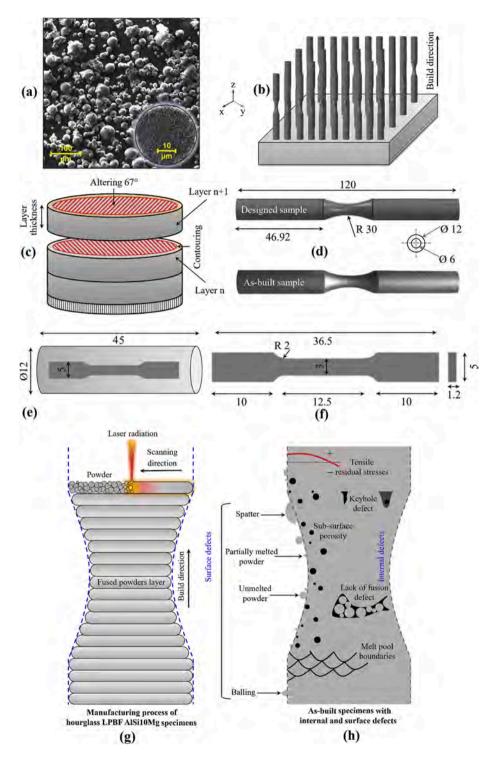


Fig. 1. (a) Morphology of the employed gas atomized AlSi10Mg powder and corresponding microstructure. (b) Cylindrical specimens fabricated along vertical build direction. (c) Schematic view of 67° rotation between adjacent layers for contouring. (d) Shape of the cylindrical hourglass fatigue test specimens. (e) Preparation of miniature tensile test specimens by wire cutting of the cylindrical parts of the fatigue test specimens and (f) shape and dimensions of the subsize tensile test specimens. (g) Schematic illustration of the manufacturing stages of the hourglass specimens layer-by-layer via LPBF technology, and (h) the possible internal and surface anomalies of these specimens. All dimensions are in mm.

1	a	Ы	le	2

AlSi10Mg powder chemie	cal composition.
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Element	Al	Si	Fe	Cu	Mn	Mg	Zn	Ti	Ni	Pb	Sn
AlSiMg10 (wt%)	Balance	9.87	0.07	< 0.01	0.002	0.35	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01

tracks of melt pool boundaries and hatching lines. While, in the AB + HT specimen, the boundaries of melt pool and hatch like traces are less noticeable resulting in considerable homogenized microstructure. Only few faded melt pool boundaries persist, which is likely phenomena in

LPBF AlSi10Mg heat treated at temperatures around 240–450 $^{\circ}$ C [72]. This homogenization leads to enhanced ductility after HT.

EBSD analyses were also performed on both AB/AB + HT specimens to obtain the maps of grain boundaries, inverse pole figure, grain size

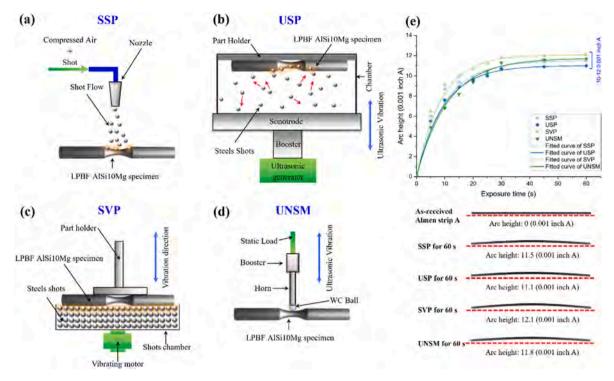


Fig. 2. Schematic illustration of different mechanical surface treatments including (a) SSP, (b) USP, (c) SVP and (d) UNSM, as well as (e) measured arc heights in different exposure times for all the considered treatments resulting in Almen intensity of 10–12 A [0.001 inch] in all of them.

Table 3

Parameters of the SSP treatment applied to the LPBF AlSi10Mg specimens.

Treatment	Shot material	Shot hardness (HRC)	Shot diameter (mm)	Projection pressure (Bar)	Peening duration (s)	Surface coverage (%)	Incident distance (mm)
SSP	Steel	48–50	0.43	3	1500	1000	120

Table 4

Parameters of USP treatment applied to the LPBF AlSi10Mg specimens.

Treatment	Shot material	Shot hardness (HRC)	Shot diameter (mm)	Shot number	Amplitude (µm)	Peening duration (s)	Incident distance (mm)
USP	Steel	48–50	0.58	250	50	300	50

Table 5

Parameters of	arameters of SVP treatment parameters for the LPBF AISITOM specimens.									
Treatment	Shot material	Shot hardness (HRC)	Frequency (Hz)	Shot type	Shot diameter (mm)	Lubrication state	Peening duration (min)			
SVP	Steel	52	50	100Cr6	1	Dry	30			

Table 6

Parameters of the UNSM treatment applied to the LPBF AlSi10Mg specimens.

Treatment	Impacting material	Frequency (kHz)	Amplitude (µm)	Linear speed (mm/min)	Static load (N)	Path Interval (µm)	Ball diameter (mm)
UNSM	WC	20	30	2000	50	65	2.5

distributions and strain contouring. Fig. 4b illustrates the grain boundary zones of both AB/AB + HT specimens with different misorientations of 2° (green lines) and 5° (black lines). Formation of finer equiaxed grains around the boundaries of melt pool (in both *xy*-plane and *yz*-plane) as well as epitaxial growth of columnar grains (in *yz*-plane), which is a common phenomenon in LPBF metallic materials, can be noted [73]. In addition, in the semi-visible melt pool boundaries of AB + HT condition, the elongation of columnar grains have also been reported by previous studies [74].

Fig. 4c indicates the inverse pole figure (IPF) maps and the oriented crystallography of the both AB/AB + HT specimens. The outputs exhibit random orientation of the grains with domination of (001) orientation in *xy*-planes which is parallel to the build direction in the longitudinal planes; this is because of the solidification along the direction of the heat transfer and the resulting epitaxial grain growth [75–78].

Fig. 4d demonstrates the corresponding grain size distribution maps exhibiting grain growth after HT. The mean grain areas of 2.8 and 44.8 μ m² were estimated on *xy*/*yz* planes for AB specimens, respectively.

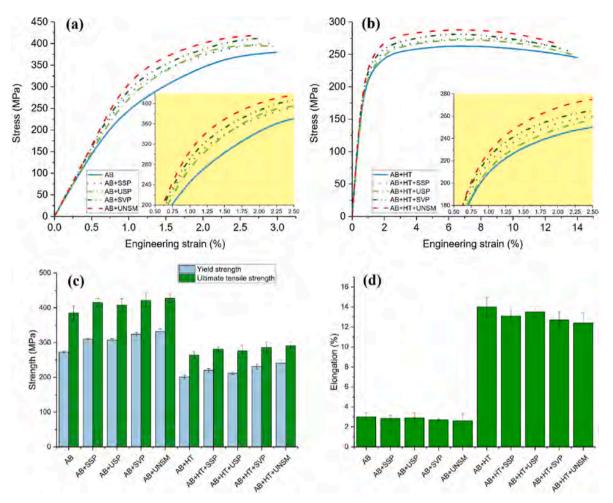


Fig. 3. Comparison of stress-strain curves of all sets of post-processed specimens in (a) as-built and (b) heat treated conditions. The average data of the three specimens tested for each set, in terms of (c) YS and UTS and (d) EL.

Table 7

Variation of tensile properties after surface post-processing in as-built and heat treated conditions.

	YS increase compared to the AB state (%)	UTS increase compared to the AB state (%)	EL reduction compared to the AB state (%)	YS increase compared to the HT state (%)	UTS increase compared to the HT state (%)	EL reduction compared to the HT state (%)
After SSP	13.55	10.98	-0.05	6.95	6.22	-0.32
After USP	12.45	8.42	-0.03	4.02	4.39	-0.18
After SVP	18.68	13.18	-0.10	10.98	8.05	-0.47
After UNSM	21.61	15.38	-0.13	14.65	9.89	-0.58

While, in the AB + HT specimens, elongated grains with larger size were measured with mean grain areas of 5.7 and 53.3 μ m² on *xy* and *yz* planes, respectively. The obtained values of mean grain sizes are compatible with the prior works [79–81].

Strain contouring maps that can be used as an index of plastic strain variations were obtained for AB/AB + HT specimens as revealed in Fig. 4e. Strain contouring reveals local plastic strain by interpretation of the inferred lattice distortion and dislocation density [82,83]. The results reveal that effective strains are located onto the melt pool regions where the density of equiaxed grains is higher mainly where the boundaries overlap. In addition, due to the significant homogenization of microstructure after HT as well strain relief, a lower maximum localized strain of 3 was obtained for AB + HT specimen compared to the AB specimen with a maximum strain of 3.5.

Surface layer of the specimens subjected to the peening-based treatments was assessed by EBSD analyses on the transverse cross-section concentrating on a scan area of 30 \times 150 μm^2 . Fig. 5 depicts IPF-Z, dynamic recrystallization (DR) intensity, kernel average misorientation (KAM) and strain contour (SC) maps all together. IPF-Z maps prove the coarsening of grains from nano-size in the outmost surface towards the core in pure mechanically operated specimens for both AB and HT conditions. It should be noted that the black fields in IPF-Z figures are related to the nanosized grains with an approximate size of 100 nm that were too small to be detected by using the allowable step size. Recrystallization maps indicate the level of mechanical deformation on the grains by using color mapping; severely deformed grains around the top surface are shown with concentrated red color. It is noticeable that the deformation density gradually decreases as they are

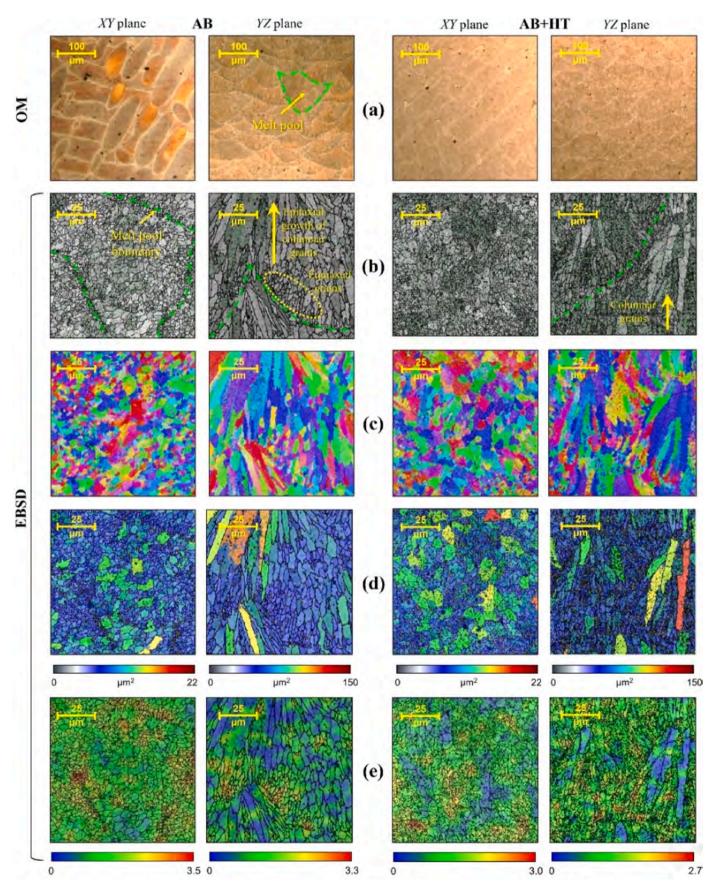


Fig. 4. Microstructural characterization results in both transversal and longitudinal cross-sections for as-built (2 left columns) and heat treated (2 right columns) states obtained by (a) OM investigations and EBSD analyses by means of (b) grain boundaries, (c) IPF, (d) grain size distributions and (e) strain contouring maps.

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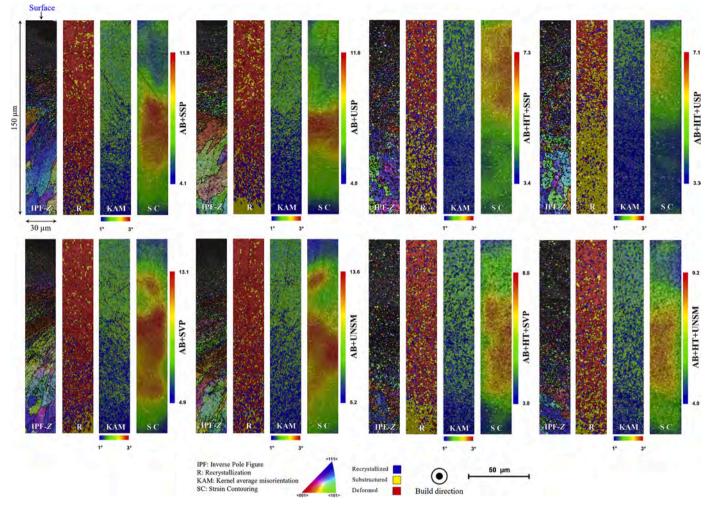


Fig. 5. EBSD results of IPF-Z, DR, KAM and SC maps in the single step treatments of AB + SSP, AB + USP, AB + SVP and AB + UNSM, as well as hybrid treatments of AB + HT + SSP, AB + HT + USP, AB + HT + SVP and AB + HT + USM.

replaced with sub-structured (yellow phase) and recrystallized (blue phase) grains towards the interior. The fraction of these three phases varies among the specimens due to the different impacting approach of the surface treatments as well as the initial differences of strength in the

as-built and HT conditions.

KAM maps that reflect the variations of applied plastic deformation and cold working [84–86], indicate the highest values in the surface outmost zone, which gradually decreases towards the center. The

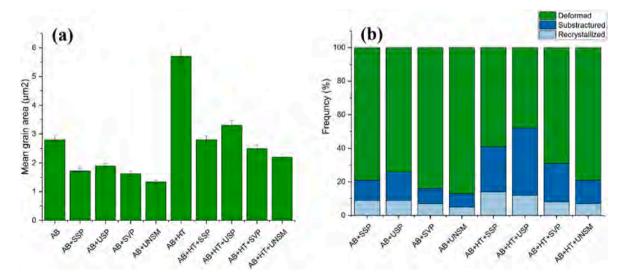


Fig. 6. Quantitative outputs of (a) mean grain area and (b) relative frequency of recrystallized, substructured and severely deformed grains detected by EBSD around the outmost layer.

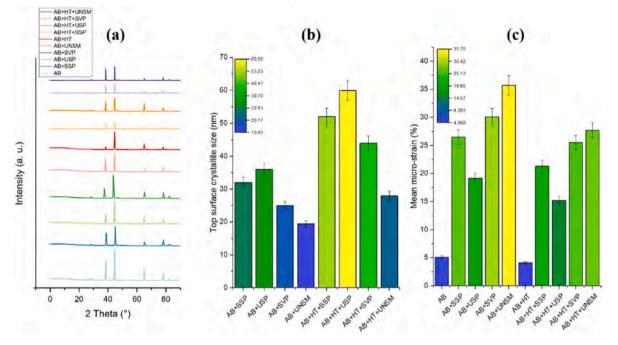


Fig. 7. (a) XRD patterns of the sets of specimens, (b) surface crystallites size and (c) micro-strains obtained by XRD analyses.

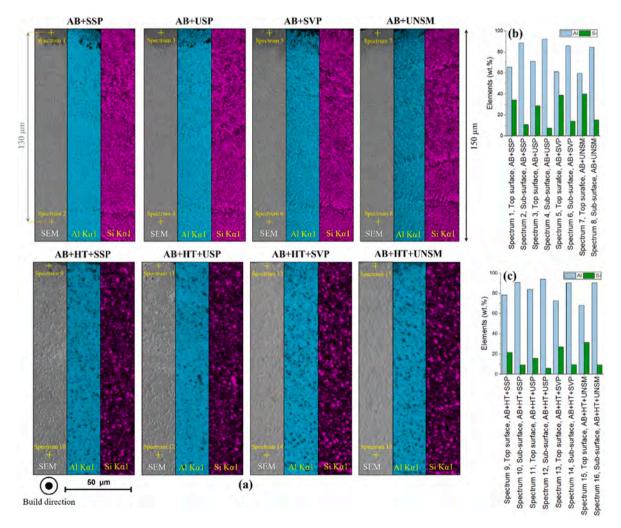


Fig. 8. (a) The obtained EDS maps and the spectrum points for treated specimens; EDS point analysis in the outmost layer for surface treated specimens (b) without and (c) with HT.

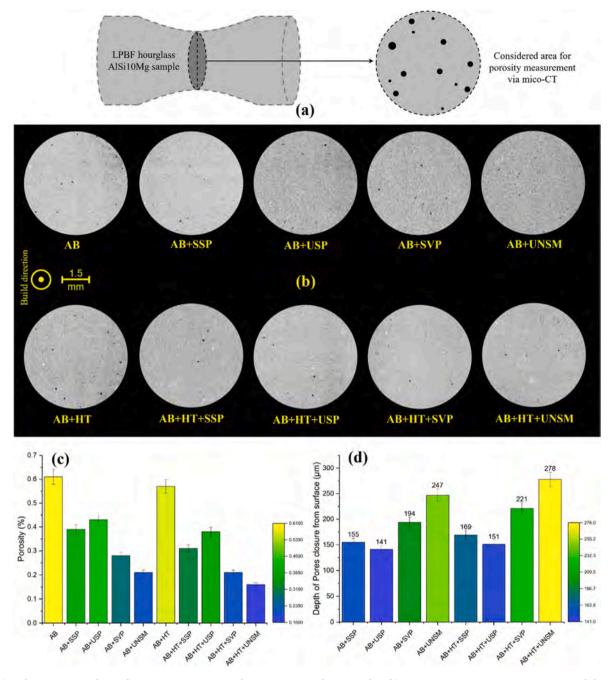


Fig. 9. (a) Mid-section scanned area for porosity measurement by X-ray computed tomography, (b) micro-CT images in transverse sections of the specimens, quantitative data of (b) mean porosities for all sets of specimens and (c) pore closure depth measured from the outmost zone in the mechanically surface treated specimens.

maximum KAM outputs were identified in certain regions with severe plastically deformed grain fields. The results reveal that specimens treated with UNSM had the highest average KAM values followed by SVP, SSP and USP, respectively, in both AB and HT conditions. In particular, mean KAM values of 1.77, 1.61, 1.87, 1.93, 1.26, 1.11, 1.52 and 1.67° were accomplished for AB + SSP, AB + USP, AB + SVP, AB + UNSM, AB + HT + SSP, AB + HT + USP, AB + HT + SVP and AB + HT + UNSM specimens, respectively. Furthermore, the high-KAM depths, corresponding to about 2° , compared to that of low-KAM values, corresponding to about 1° , indicates a good agreement with the depth of refined grains in the IPF-*Z* maps.

Strain contouring maps demonstrate that the highest plastic strains were induced in the specimens treated with UNSM followed by SVP, SSP and USP, respectively. The max. localized plastic strains of 11.9, 11.6, 13.1, 13.6, 7.3, 7.1, 8.8 and 9.2 were determined for AB + SSP, AB + USP, AB + SVP, AB + UNSM, AB + HT + SSP, AB + HT + USP, AB + HT + SVP and AB + HT + UNSM parts, respectively. It has to be mentioned that the max. plastic strains were located just below the surface as is common in the mechanically surface treated materials [87–89]. The results of KAM and strain contouring maps confirm that the applied surface treatments led to lower strain on the HT sets owing to better ductility compared with the sets with initial AB condition.

Quantitative grains size analysis based on EBSD data, shown in Fig. 6a, demonstrates mean grain size reduction in the considered scan area in the surface treated specimens. Mean grain areas of 1.7, 1.8, 1.6, 1.3, 2.8, 3.3, 2.5 and 2.2 μ m² were measured for AB + SSP, AB + USP, AB + SVP, AB + UNSM, AB + HT + SSP, AB + HT + USP, AB + HT + SVP and AB + HT + UNSM specimens, respectively. UNSM was shown to be

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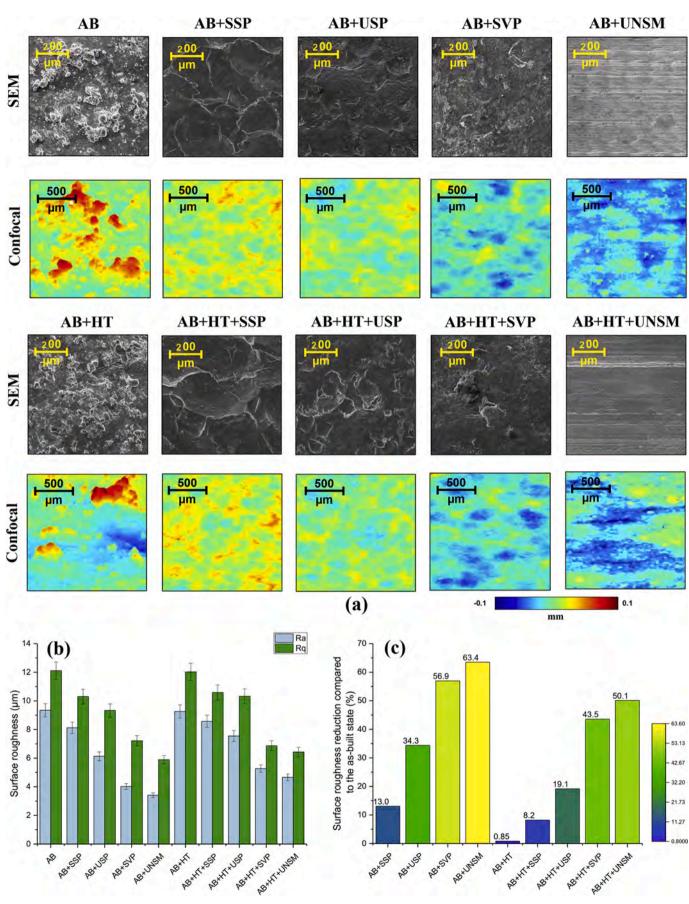


Fig. 10. (a) Surface morphologies of specimens using SEM and confocal observations, (b) calculated surface roughness parameters R_a and R_q , and (c) comparison of the effects of all post-processing methods on roughness reduction compared to initial state (based on R_a values).

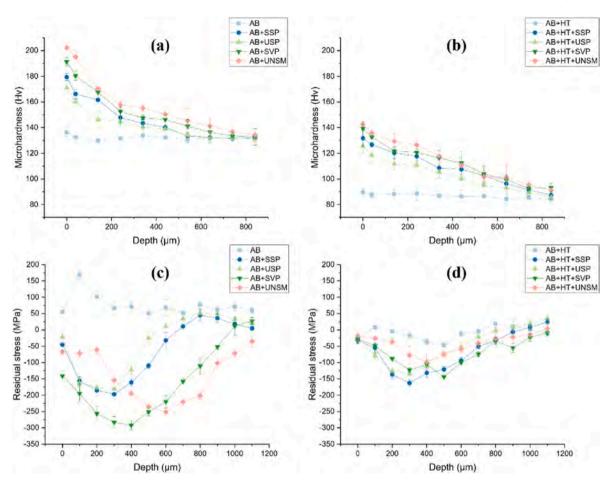


Fig. 11. Microhardness profiles for surface treated specimens with initial (a) AB and (b) AB + HT states, and distributions of residual stresses for surface treated specimens with initial (c) AB and (d) AB + HT states.

the most effective on refining the grains followed by SVP, SSP and USP. Moreover, as presented in Fig. 6b, specimens treated with UNSM achieved the highest fraction of plastically deformed grain intensity per unit area followed by SVP, SSP and USP.

XRD analyses were also used for the determination of top surface crystallite size employing Williamson-Hall methods. Fig. 7a presents the XRD patterns of all the conditions. The applied mechanical surface posttreatments diminished the peak intensity and caused peak-broadening and shift owing to the severe plastic deformation. Fig. 7b indicates that the calculated top surface crystallites size of the surface treated specimens is lower than 100 nm. In particular, surface crystallites size of 32, 36, 25, 19, 52, 60, 44 and 28 nm were measured after AB + SSP, AB + USP, AB + SVP, AB + UNSM, AB + HT + SSP, AB + HT + USP, AB + HT + SVP and AB + HT + UNSM operations, respectively. The UNSM treatment had the highest influence on top surface nanocrystallization thereafter SVP, SSP and USP, while all these treatments were implemented using the same intensity of 10-12 A [0.001 inch]. The microstrains for the specimens are demonstrated in Fig. 7c showing considerable plastic strain induced by the mechanical surface treatments. Moreover, it was found that trends of variations of grains size and plastic strains obtained by XRD agree with the ones obtained for grains size by KAM and strain contouring maps from EBSD analysis (see Figs. 5 and 6a).

In the previous works on the effects of SVP on microstructural evolution and fatigue performance of V-notched LPBF AlSi10Mg [37], we reported that besides the formation of gradient microstructure caused by grain refinement, a chemical gradient of Si phase was also observed. Herein, to investigate this phenomenon, EDS analysis was performed on the surface as depicted in Fig. 8a. Two spectrum points were considered in each specimen. EDS maps were obtained in the same area used for EBSD analyses (30 \times 150 μm^2), adding one spectrum point on the outmost deformed surface and another one in the sub-surface area, i.e. 130 μm below the surface. The results reveal that the fraction of Al phase was reduced after surface treatments while Si phase fraction increased on the outmost zone. In addition, the fibrous Si networks observed in the AB state were transformed to homogeneously dispersed spherical Si particles after HT. This considerable increase of Si fraction on the outmost surface is quantitatively demonstrated in Fig. 8b and c.

In the top surfaces, Si fractions of 34.2, 28.6, 38.7, 40.1, 21.4, 15.7, 27.1 and 31.7% resulted from AB + SSP, AB + USP, AB + SVP, AB + UNSM, AB + HT + SSP, AB + HT + USP, AB + HT + SVP and AB + HT + UNSM operations, respectively. However, in the sub-surface regions, the Si fractions were only 7.3-15.1 and 5.7-9.4% for plastically deformed specimens with initial AB and HT conditions, respectively. The results indicate the occurrence of Si phase chemical gradient near the surface zone. The amount and distribution of Si content can considerably impact the tensile strength and tribological features of Al-Si alloys [90,91]. Increasing the content of Si in Al alloy can enhance the hardenability, which typically leads to reduced EL [92]. It has been shown that by increasing the content of Si in LPBF Al-Si alloys, the solidification cracking risk can be decreased [93] and tensile strength can be improved [94]. Therefore, application of these surface treatments can enhance performance and local strengthening through simultaneous induction of microstructural (confirmed by EBSD analyses) and chemical (confirmed by EDS analyses) gradients.

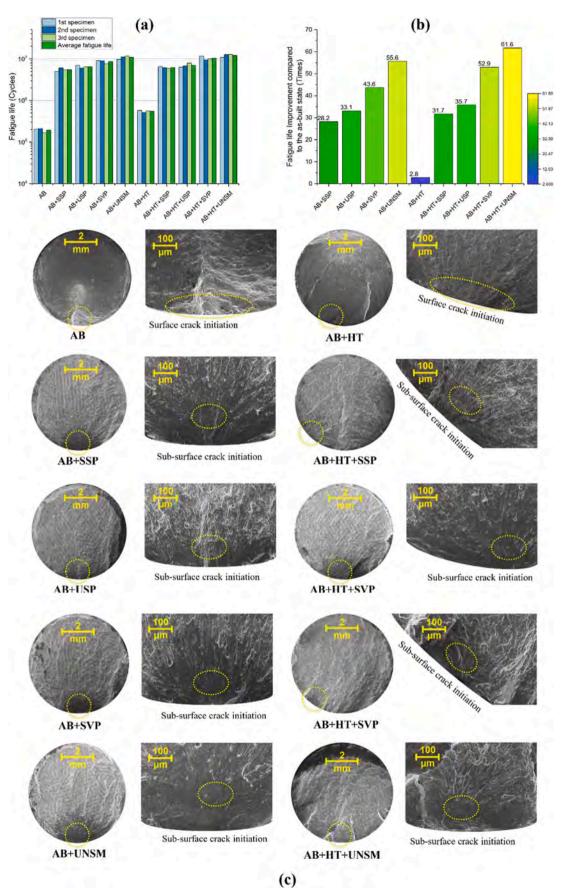


Fig. 12. (a) Fatigue test results at a stress of 110 MPa, (b) fatigue life alteration of post-processed specimens compared to the AB ones and (c) fracture surfaces specifying crack nucleation site and crack propagation direction.

3.3. Porosity analysis

Fig. 9a indicates the scanned area using X-ray computed tomography for porosity measurement on the mid-part of the specimens. The narrowest section of specimens was selected as it theoretically should be the most critical section prone to fatigue failure. Three different specimens were employed for porosity analysis of each condition. Fig. 9b shows the micro-CT images of the specimens in which the heterogeneously distributed pores can be distinguished with black color. The micro-CT images reveal that porosity was reduced by the mechanical surface treatments. Considering all the scanning area, mean porosities of 0.61, 0.39, 0.43, 0.28, 0.21, 0.57, 0.31, 0.38, 0.21 and 0.16% were measured for AB, AB + SSP, AB + USP, AB + SVP, AB + UNSM, AB + HT, AB + HT + SSP, AB + HT + USP, AB + HT + SVP and AB + HT + UNSM specimens, respectively (see Fig. 9c).

In addition, most of the sub-surface porosities were closed in the mechanically treated specimens; however, within different depths. As shown in Fig. 9d, depths of pore closure from surface were measured to be 155, 141, 194, 247, 169, 151, 221 and 278 μ m after AB + SSP, AB + USP, AB + SVP, AB + UNSM, AB + HT + SSP, AB + HT + USP, AB + HT + SVP and AB + HT + UNSM operations, respectively. UNSM exhibited the highest efficacy for pore closure, followed by SVP, SSP and USP, respectively. Overall, the depth of pores closure was higher in the HT series due to their lower strength, resulting in deeper effect of mechanical surface treatment, compared to the specimens in AB initial condition.

3.4. Surface roughness and morphology analysis

The SEM and confocal surface morphological observations of all specimens are presented in Fig. 10a. Due to the attached unmelted/ partially melted powders, the surface of both AB/AB + HT specimens were considerably rough. However, these powder particles were removed after the application of peening-based treatments leading to more uniform surface morphologies. Due to the physics of the surface operations and the specifics of the corresponding apparatus, each method created a unique surface morphology. In the case of SSP, USP and SVP treatments, the impact of shots created dimples and overlapping craters, with different depth and shapes due to the differences in the used media for each treatment. Nevertheless, the UNSM treated specimens exhibited a totally different surface morphology with semi-parallel tracks indicating the scanning directions.

The roughness parameters of R_a and R_g are presented in Fig. 10b for each surface treatment condition. The results reveal that Ra values of both AB/AB + HT specimens with 9.34 and 9.26 μ m, respectively, were decreased after mechanical surface treatments down to 8.12, 6.13, 4.02, 3.41, 9.26, 8.57, 7.55, 5.27 and 4.66 µm for AB + SSP, AB + USP, AB + SVP, AB + UNSME, AB + HT + SSP, AB + HT + USP, AB + HT + SVP and AB + HT + UNSM specimens, respectively. Roughness of the surface treated specimens in the non-HT state were generally lower than the ones in heat treated condition. The phenomena could be clarified via higher deformability of HT specimens allowing the mechanical treatments to induce deeper surface deformation. Rq values exhibited a similar trend with Ra. Fig. 10c shows the effectiveness of the postprocessing methods on surface roughness reduction by comparing their resulting R_a to the initial untreated specimens. UNSM was the most influential operation on surface texture followed by SVP, USP and SSP. Although SSP induced higher plastic strains compared to USP (according to the obtained results of microstructural characterization in section 3.2 and porosity analyses in section 3.3), surface morphology images reveal a more uniform surface after USP. In addition, it was found that USP can be more effective than SSP in surface roughness reduction. Similar results were reported comparing the application of USP and SP processes with almost same intensities on LPBF Ti-6Al-4V [95]. The phenomenon is due to the intrinsic differences between the two treatments.

3.5. Microhardness and residual stress measurement

The near surface microhardness profiles of specimens in non-HT and HT conditions are presented in Fig. 11a and b, respectively. AB + HT specimens, with higher ductility and lower strength, exhibit lower microhardness compared to the AB ones [58]. The surface treatments led to considerable surface layer hardening with the highest microhardness on the outmost surface that steadily decreases towards the center of the specimen. Grain size reduction is well known to induce strengthening [96]. Considering grain refinement data obtained from the EBSD analysis, as expected UNSM exhibited the highest surface hardness followed by SVP, SSP and USP. Surface microhardness improvements of 31, 25, 40, 48, 47, 40, 56 and 59% were accomplished for AB + SSP, AB + USP, AB + SVP, AB + UNSME, AB + HT + SSP, AB + HT + USP, AB + HT + SVP and AB + HT + UNSM specimens, respectively. As results indicate, higher hardness improvement was obtianed in HT specimens owing to their higher deformability.

The residual stress distributions from the outmost layer towards the inside of the specimens for all conditions are presented in Fig. 11c and d. TRS in the AB specimens were released and transformed to CRS in the AB + HT specimens. Mechanical surface post-treatments remarkably induced CRS in the specimens in both AB and HT initial condition. The trend of the induced CRS varied based on the applied surface treatment. Surface residual stresses of 55, -45, -21, -141, -67, -25, -32, -24, -28 and -18 were measured for AB, AB + SSP, AB + USP, AB + SVP, AB + UNSME, AB + HT, AB + HT + SSP, AB + HT + USP, AB + HT + SVP and AB + HT + UNSM conditions, respectively. Among surface treated specimens that were initially non-HT, the highest maximum CRS was obtained for the AB + SVP specimens (-292 MPa) and the highest depth of CRS was achieved in AB + UNSM specimens (1100 μ m).

3.6. Fatigue behavior

Rotating bending fatigue experiments at stress level of 110 MPa were conducted using 3 specimens per set to assess the fatigue behavior of LPBF AlSi10Mg as affected by post-processing. Fig. 12a presents the results of fatigue tests confirming that all the individual and hybrid posttreatments positively impact the fatigue performance of the hourglass LPBF AlSi10Mg specimens. Fatigue lives improved after applying HT because of microstructural homogenization, enhanced ductility, and relaxation of the TRS. From another perspective, mechanical surface treatments enhanced the fatigue performance resulting from the combined effects of considerable pore closure just below the surface, surface grain refinement and hardening, induced CRS and surface texture reduction. The hybrid post-processing methods (HT + surface treatments) take advantage of all the above-mentioned effects in a synergistic manner.

Mean fatigue lives of 1.95×10^5 , 5.51×10^6 , 6.47×10^6 , 8.53×10^6 , 1.08×10^7 , 5.5×10^5 , 6.19×10^6 , 6.98×10^6 , 1.03×10^7 and 1.20×10^7 cycles were obtained for AB, AB + SSP, AB + USP, AB + SVP, AB + UNSM, AB + HT, AB + HT + SSP, AB + HT + USP, AB + HT + SVP and AB + HT + UNSM conditions, respectively. The results exhibited note-worthy fatigue life improvement particularly in HT + UNSM, HT + SVP and UNSM processes with over 10^7 cycles. Fig. 12b represents the corresponding fatigue life enhancement by post-processing in comparison with the AB condition. UNSM demonstrated higher fatigue resistance than SVP, USP and SSP operations for both AB and AB + HT conditions. The results indicate that while SSP treated specimens had higher hardness and CRS as well as more porosity reduction compared to the SVP treated ones, the SVP sets exhibited higher fatigue lives highlighting substantial effect of lower surface roughness and morphology regularity.

Fatigue fracture surfaces are shown in Fig. 12c. The crack initiated from surface and propagated in the bulk in both AB and AB + HT specimens. Nevertheless, in the specimens subjected to mechanical surface treatments (in both initially AB and AB + HT states), sub-surface crack initiations were observed resulting from surface roughness

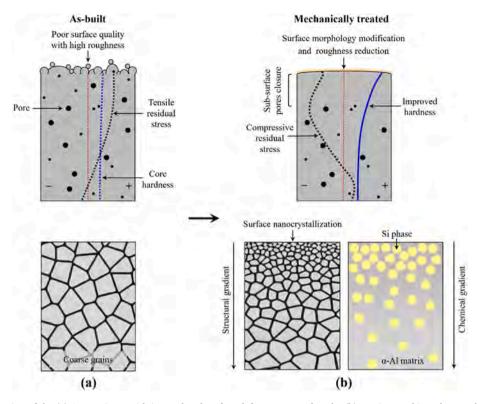


Fig. 13. Schematic illustration of the (a) AB specimen with internal and surface defects compared to the (b) specimen subjected to mechanical surface treatments with modified properties.

reduction, as well as surface layer hardening and CRS.

4. Discussions

According to the obtained results it was found that applying postprocessing methods can highly modify the internal and surface anomalies of LPBF AlSi10Mg specimen compared to the as-built state. Fig. 13 schematically illustrates the near-surface state of AB specimens with the intrinsic internal and surface anomalies compared to the specimens subjected to mechanical surface treatments. These surface postprocessing methods induced gradients in microstructure and chemical composition on the surface, resulting from the formation of nanosized grains and promoting the density of Si particles. UNSM showed the best performance on surface nanocrystallization, formation of structural and chemical gradients, inducing plastic strains and surface layer hardening followed by SVP, SSP and USSP. Formation of gradients in surface layer led to improved hardness and strength, induce CRS and sub-surface pore closure, while at the same time enhance the surface morphology and reduce surface texture. In addition, mechanical surface treatments were effective in inducing sub-surface pore closure and compressive residual stresses. A maximum depth of pore closure of 278 μm was obtained through the hybrid post-treatment of HT + UNSM. Also, the deepest compressive residual stresses were achieved with UNSM (up to the depth 1100 μm).

UNSM demonstrated highest efficiency for improved fatigue behavior followed by SVP, USP and SSP operations, respectively, for both as-built and heat treated conditions. In addition, the results revealed that the effective parameters of relative density, depth of pore closure, surface layer grain refinement and hardening, residual stress distributions and surface texture as well tensile properties, all together simultaneously affect the fatigue behavior but with different impacts. For instance, the results indicate that while SSP treated specimens had higher hardness and CRS as well as more porosity reduction compared to the USP treated series, the USP sets exhibited higher fatigue lives

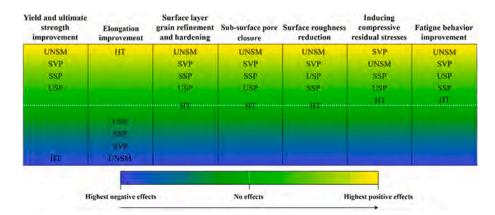


Fig. 14. Comparison of the effect of post-processing methods on YS and UTS, ductility, grain refinement and hardening, roughness, CRS, and fatigue performance.

		_	(;	a)						
	Т	ime	Money :	nalys	is matrix	4				
	Very High	5	5	10	LS.	20	25	1 TOT	Very	•
	Iligh	sb 4		*	12	12 16 9 12	20		Elig Modes	
Time	Moderate	3	.3	6	4		15	Cust		
	Low	2	2	4	6	8	10		L	
	Very Law	1	1	2	3	4	5		Very	
Idea	1		1	1	X	4	5	Ide	al	
Mode	_		Very Low	Low	Moderate	High	Very High	Mode	-	
Not-ic	teal		11		Money		1	Not-it	deal	

	Cost	-Per	formanc	e ana	lysis mat	rix		
107	Very High	5	5	10	15	20 16 12	-25	
	tligh	4	4	8	12		20 15	
Cost	Moderate	ate 3		-6				
	Low	1	2	+	6	8	10	
	Very Low	ù	10	ż	3	4	5	
Idea	al	٩.	1	2	3	*	5	
Mode	-		Very High	High	Moderate	Low	Very Low	
Noten	deal			Performance				

(h)

(c)

	1.	Cost (Tim	e-Money)		Cost-Performance				
Post-treatment	Time	Money	Final Score	Rank	Cost	Performanc e	Final Score	Rank	
НТ	Moderate	Low	6	3	Low	Very low	10	4	
SSP	Low	Very low	2	11	Very low	Moderate	3	1	
USP	Very low	Moderate	3	2	Very low	Moderate	3	<u>.</u>	
SVP	Moderate	Lów	6	3	Low	High	4	2	
UNSM	Low	High	8	4	Moderate.	Very high	3	1	
HT+SSP	Very high	Moderate	15	5	High	Moderate	12	5	
HT+USP	High	High	16	6	Very high	Moderate	15	6	
HT+SVP	Very high	Moderate	15	5	High	Very high	4	2	
HT+UNSM	High	Very high	20	7	Very high	Very high	5	3	

(d)

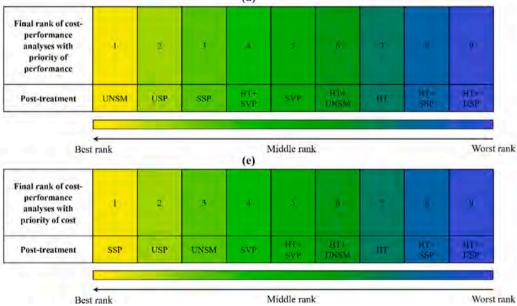


Fig. 15. Rating variables using five levels and associated scores for (a) Time-Money, (b) Cost-Performance outputs and (c) obtained scores of considered post-processing methods by means of cost/performance analysis, as well as overall ranks based on the results of Cost-Performance analyses by considering priority for (d) performance and (e) cost.

highlighting substantial effect of lower surface roughness and regular morphology. The influence of each post-processing method individually on YS and UTS, ductility, grain refinement and hardening, pore closure, surface texture modification and CRS and finally fatigue performance of the post-treated specimens are summarized in Fig. 14.

Moreover, in order to compare the performance of applied posttreatments in economical fashion, Cost analysis and Cost-Performance analysis were carried out to determine the efficiency of the postoperations, particularly in terms of fatigue performance improvement. Following the methods used in risk analysis [97,98], each input was assessed as five levels of quite (very) low, low, moderate, high and quite (very) high with progressive scores of 1-5. Subsequently, a matrix was used for each set of data (Time-Money or Cost-Performance) to calculate scores for the post-operations [99]. The developed matrices of duration (Time)-Money and Cost-Performance relations are shown in Fig. 15a and b, respectively. The scores and ranks of cost (Time-Money) analysis were used in the Cost-Performance analysis. The determined scores for post-processing methods and the evaluated final ranks are shown in Fig. 15c. The criteria of assigning levels to time (considering required time for treatment of 15 specimens) and performance parameters are presented in Appendix A. The comparative cost and time levels related to each treatment were determined based on the authors' experience. The analysis results indicate similar rankings for some of the treatments regarding both cost and performance related factors. Considering the priorities of performance and cost, the applied post-treatments can be ranked as depicted in Fig. 15d and e, respectively. For instance, by considering the priority of performance in the Cost-Performance analyses, UNSM was found as the first rank treatment followed by USP, SSP, HT + SVP, SVP, HT + UNSM, HT, HT + SSP and HT + USP.

5. Conclusions

In this study, the efficacy of various mechanical surface posttreatments, involving SSP, USSP, SVP and UNSM, independent of and combined with T6 type heat treatment, was investigated on reducing the hazardous influence of internal and external (surface) imperfections on fatigue performance of LPBF AlSi10Mg specimens. Based on the experimental observations and analysis of the results, the following conclusions can be drawn:

- The mechanical surface treatments induced maximum of 0.5% reduction of elongation, while improving the yield and ultimate tensile strengths up by 21 and 15%, respectively, owing to grain size reduction and compressive residual stresses.
- While each mechanical surface treatment generated a distinct surface morphology, all of them eliminated the surface imperfections and reduced the surface roughness leading to a more uniform surface morphology. Contrary to all surface treatments that generated

Appendix A

Table A1

Time intervals considered for assigning time levels

overlapping surface dimples due to impacting of shots with different depths and shapes of valleys, UNSM induced semi-parallel lines along scanning directions. UNSM was found to be the most influential technique in surface roughness reduction, followed by SVP, USP and SSP, in order.

- In general, all the employed mechanical surface treatments improved the fatigue performance due to their remarkable synergistic effect of increasing tensile strength, inducing pore closure, reducing the surface grain size, increasing the surface hardening, and inducing compressive residual stresses in addition to surface texture reduction. The best fatigue life enhancement was obtained for the hybrid treatments; heat treatment + UNSM (× 61) followed by UNSM (× 55) and heat treatment + SVP (× 52) compared to the AB condition.
- Obtained results of Cost-Performance analyses by considering parameters of time, cost and performance with a special focus on fatigue behavior identified UNSM as the first-rank treatment followed by USP, SVP, HT + SVP, SVP, HT + UNSM, HT, HT + SSP and HT + USP. However, it should be noted that the selection of a proper shotprocessing treatment for additively manufactured parts requires a case-by-case study in which the effect of geometry and material properties should be also considered.

CRediT authorship contribution statement

Erfan Maleki: Conceptualization, Investigation, Formal analysis, Visualization, Writing – original draft. Sara Bagherifard: Conceptualization, Methodology, Validation, Writing – review & editing, Supervision. Okan Unal: Methodology, Investigation, Formal analysis, Writing – review & editing. Shuai Shao: Formal analysis, Writing – review & editing. Nima Shamsaei: Resources, Writing – review & editing. Mario Guagliano: Conceptualization, Resources, Writing – review & editing, Supervision.

Declaration of competing interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

Data availability

Data will be made available on request.

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Time level	Very low	Low	Moderate	High	Very high
Time interval (min)	0–199	200–399	400–599	600–799	800-1000

Table A2

Fatigue life improvement intervals considered for assigning performance levels

Performance level	Very low	Low	Moderate	High	Very high
Fatigue life improvement (times)	0–12.3	12.4–24.7	24.8–37.1	37.2–49.5	49.6–62

Table A3

Assigned levels of time and performance for all the applied post-treatments

Post- treatment	Required time for treatment of 1 sample (min)	Required time for treatment of 15 samples (min)	Time level	Fatigue life improvement (times)	Performance level
HT	550	550	Moderate	2.8	Very low
SSP	25	$25 \times 15 = 375$	Low	28.2	Moderate
USP	5	$5 \times 15 = 75$	Very low	33.1	Moderate
SVP	30	30 imes 15 = 450	Moderate	43.6	High
UNSM	15	15 imes15=225	Low	55.6	Very high
HT + SSP	550 + 25	$550+(25 \times 15) = 925$	Very high	31.7	Moderate
HT + USP	550 + 5	$550+(5 \times 15) = 625$	High	35.7	Moderate
HT + SVP	550 + 30	$550+(30 \times 15) = 1000$	Very high	52.9	Very high
HT + UNSM	550 + 15	$550+(15 \times 15) = 775$	High	61.6	Very high

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