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Letter

Compression and fatigue performance of additively manufactured NiTi architected shape memory alloys

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Abstract

Additive manufacturing of architected materials—particularly lattice or porous structures—has gained significant attention in recent years due to their enhanced strength-to-weight ratios, load-bearing capabilities, and energy absorption properties. The integration of these structures with shape memory alloys offers multifunctional performance for advanced engineering applications. This study investigates the compressive fatigue behavior of NiTi lattice structures fabricated by Laser powder bed fusion. Initial quasi-static compression tests, carried out to full structural collapse, were used to define load levels for subsequent fatigue experiments. Fatigue testing was then conducted at 40 °C to induce pseudoelastic behavior, and an S–N curve was generated to characterize fatigue performance. Results showed that the NiTi lattice could sustain cyclic loading at 8 kN for an average of approximately 86 000 cycles, and around 18 000 cycles at 11 kN. Post-mortem microstructural analyses revealed martensite accumulation near fracture regions, attributed to stress-induced phase transformation.

Keywords: additive manufacturing, LPBF, shape memory alloys, NiTi, architected structures, fatigue

1. Introduction

In recent years, considerable efforts have been directed toward enhancing structural functionality through the integration of the inherent capabilities of shape memory alloys (SMAs) [1, 2]. The remarkable flexibility and freeform capacity of additive manufacturing (AM) technologies have shown

significant promise in fabricating SMAs, thus opening up new avenues of opportunity by enabling the creation of intricate geometries, like complex shape lattice structures, previously unattainable with conventional manufacturing techniques [3, 4]. One of the most encouraging potential applications of AM SMAs is in the fabrication of lattice or porous structures with shape recovery performance, so that they can retrieve their original shape [5–7]. These structures have demonstrated remarkable thermomechanical performance in terms of light-weight properties [8–10], improved damping abilities [11], high energy absorption [12, 13], and high specific strength and stiffness [14–17].

Architected NiTi structures produced by AM are typically designed for conditions in which compressive forces predominate. One of the advantages of AM technology in

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fabricating these types of structures is the ability to optimize parameters such as strut diameter, strut length, wall thickness, or unit cell configuration [18–23]. This optimization is particularly beneficial for tuning the mechanical performance of such lattices as well as load bearing, Young's modulus of components, and energy absorption applications [24–28]. For example, there is a report on improvement in the design of NiTi-SMA gradient lattice structure by outstanding shape memory effect and functional stability with a larger displacement recovery interval of up to 99% for the unidirectional setup and an accumulative irrecoverable strain up to 0.8% which showed the excellent compression energy absorption [29]. In another study, enhancement in the architected design of Fe-based SMA through AM techniques demonstrated increased compressive strength. This improvement led to enhanced multifunctional behavior, energy absorption capacity, load-bearing ability, and geometric recoverability of the components [30].

Multiple studies have highlighted the complex fatigue behavior of superelastic Nitinol, emphasizing the influence of microstructure, loading conditions, and processing history [31–37]. For instance, Mahtabi *et al* reviewed the state-of-the-art and ongoing challenges, noting that fatigue in nitinol is more complex than in traditional alloys due to its unique superelastic and shape memory properties, with microstructural effects and testing conditions playing significant roles in fatigue life [38]. Malito *et al* established a methodology for determining small crack growth thresholds, finding that small cracks—comparable to native inclusions—are critical for predicting fatigue lifetime in medical devices [39]. Launey *et al* demonstrated that pre-strain, mean strain, temperature, and strain amplitude all significantly affect the fatigue response, and they proposed both empirical and energy-based models for predicting fatigue strain limits [40]. Pelton *et al* found that pre-strain and mean strain can either enhance or degrade fatigue performance, with cycling in the two-phase region generally leading to longer fatigue lives due to martensite stabilization and cyclic phase transformation [41]. Gugat *et al* reported excellent high-cycle fatigue resistance in sputtered nitinol thin films, with a fatigue safety limit of 1.75% strain amplitude for mean strains up to 2.5% [42]. Additionally, Shabani Nezhad *et al* showed that wire size and microstructural inhomogeneities introduced during processing significantly influence fatigue behavior, with smaller wires being more vulnerable to surface defects but exhibiting better functional performance [43]. These findings collectively underscore the importance of microstructural control, loading history, and device geometry in optimizing the fatigue performance of superelastic nitinol.

Alarcon *et al* reported a significant reduction in the fatigue performance of NiTi wires, particularly when loading approaches the stress-induced martensitic transformation (MT) regime, which results in large recoverable strains due to the superelastic effect [44]. Similarly, Sherif *et al* demonstrated successful fatigue cycling of NiTi wires and explored the influence of various parameters, such as maximum tensile stress and residual strain, on the material's cyclic performance. However, the fatigue behavior of NiTi SMA lattice

structures under varying stress conditions presents a distinct research direction. These investigations highlight the potential of tailored design strategies to achieve high-cycle functional performance, which is particularly critical for biomedical applications like stents [45]. It is important to note that the functional longevity of a printed component is not determined solely by structural failure; high functional fatigue life is equally important and can be heavily influenced by lattice geometry.

Despite this, most existing studies have focused on the limited cycling behavior of SMA lattices, underscoring the need for further research involving extended fatigue cycles [46–49]. The interplay between geometric design and scanning strategy plays a vital role in fatigue performance, as it governs the initiation and growth of defects or inclusions that can severely degrade the material's endurance [50]. For instance, Chen *et al* demonstrated that careful component design can lead to enhanced fatigue life in NiTi lattice structures [29]. Biffi *et al* investigated the cyclic compression and shape recovery of NiTi architected meta-structures produced via the laser powder bed fusion (LPBF) process, employing three different compression loads to assess a broad range of shape recovery behavior and energy dissipation characteristics [51, 52]. Similarly, Speirs *et al* reported that the fatigue performance of NiTi structures is strongly influenced by design features that generate stress concentrations and facilitate crack initiation [53], a finding further validated by Ren *et al* in a separate study [54].

This study examines fatigue behavior over an extended range of cycles. Fatigue tests are conducted under varying load levels to enable the development of an S–N curve tailored specifically to the NiTi lattice structure. Additionally, the relationship between transformation temperatures and the evolving microstructure in different regions of the tested samples is analyzed to provide insights into how the material's properties change throughout cyclic loading. This approach enhances the understanding of the material's behavior and phase transformations during the experimental process.

2. Experimental procedure

Spherical Ni-rich NiTi powder (Ni56Ti44 wt.%) with a particle size of 16–63 μm (Nanoval, Germany) was processed using an LPBF system (Renishaw AM400, UK). The process parameters were optimized to achieve a 99.5% relative density of [51]. Table 1 reports the adopted process parameters.

The architected structures investigated in this study exhibit auxetic behavior, characterized by a negative Poisson's ratio, as reported in our previous work [51, 52] and illustrated in figure 1. A total of ten NiTi samples were fabricated in a horizontal orientation on the build platform and subsequently heat-treated at 500 °C for 5 min, followed by water quenching. Samples 1 and 2 were subjected to monotonic compression tests to determine the maximum compressive force leading to failure, while Samples 3 through 10 were allocated for fatigue testing.

Compression tests were performed at room temperature with a 100 kN load cell, using a crosshead speed of

Table 1. List of process parameters used for the printing of the architected NiTi samples.

Laser power	Exposure time	Oxygen level	Atmosphere	Layer thickness	Hatch distance	Laser spot size	Point distance	Scanning strategy
150 W	75 μ s	<20 ppm	Argon	30 μ m	50 μ m	65 μ m	120 μ m	Meander

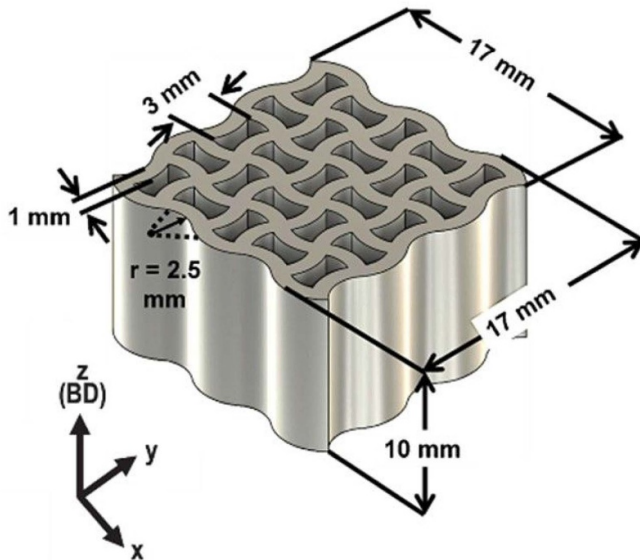


Figure 1. CAD model, highlighting the principal dimensions of the investigated architected structure [55]. Reproduced from [55]. CC BY 4.0.

1 mm min⁻¹ (Instron 5989). Once the failure force was identified, fatigue tests were conducted at 5 Hz with a stress ratio of $R = 0.1$, varying the applied load at three levels (MTS 810). Before testing, samples were heated to 40 °C to maintain the austenite phase. Specifically, three samples (5–7) were tested at a maximum load of 8 kN, two (3–4) at 9 kN, and three (8–10) at 11 kN. Given the stress ratio of 0.1, the corresponding minimum loads were 0.8, 0.9, and 1.1 kN, respectively. Based on the effective cross-sectional area of the samples (60 mm²), this corresponds to maximum stress amplitudes of approximately 133, 150, and 183 MPa. The lattice specimen has a nominal (projected) cross-sectional area of 10 mm \times 15 mm (150 mm²). However, due to its geometry, the structure consists of 6 individual struts, each approximately 1 mm wide and spanning the full 10 mm depth. Therefore, the effective cross-sectional area is estimated to be 6 \times 1 mm \times 10 mm (60 mm²).

After the fatigue testing, the fractured regions of the samples were observed with Optical Microscopy (Keyence, Model VHX-1000) at low magnification and scanning electron microscopy (Nova, Model NanoSEM450) at high magnification. Furthermore, the results obtained from electron back-scattered diffraction (EDAX Trident system) were correlated to study the microstructural evolution of the tested samples. The MT temperatures were measured using differential scanning calorimetry (DSC, Q25 model, TA Instruments, USA). The analysis was conducted with a heating/cooling rate of 10 °C min⁻¹ over a temperature range of -50 °C to 100 °C. DSC was performed on the as-built samples and two samples after fatigue analysis.

3. Results and discussion

The mechanical behavior of the architected structures was evaluated through compression tests on two samples (Samples 1 and 2) until fracture. The corresponding force-displacement curves (figure 2(a)) show an initial linear increase in load up to approximately 0.35 mm of displacement, followed by a second slope leading to structural collapse at around 20 kN. This second, shallower slope likely reflects a transition from purely elastic deformation to localized structural rearrangement or the onset of plastic deformation within the architecture, such as buckling of struts or progressive failure at critical nodes. This behavior stabilizes until reaching a maximum displacement between 1.0 and 1.25 mm. Notably, this transition occurs at approximately 10 kN, indicating that this load range may be of interest for further study.

Fatigue tests in compression were conducted at 40 °C to assess the functional performance of NiTi architected structures under high-cycle conditions. At this temperature, the material is fully austenitic, as the austenite finish temperatures (A_f) of all samples, determined via DSC, are well below 40 °C (see figure 5). While fatigue testing occurred in the austenitic state, it is possible that MT occurred during post-test SEM analysis at room temperature. Loads of 8, 9, and 11 kN were applied to determine the S-N curves, with figures 2(b)–(d) illustrating the deformation evolution per cycle under different load conditions. Higher loads resulted in greater deformation and a shorter lifespan, highlighting the load-dependent nature of the fatigue response. Notably, among the samples tested at 11 kN (Samples 8, 9, and 10), Sample 9 failed significantly earlier than the others, 13 841 cycles vs. 19 898 and 20 960 cycles for Samples 8 and 10, respectively. This discrepancy may be attributed to variations in the LPBF process affecting the internal structure, such as porosity or local residual stresses. Additionally, the initial deformation at the start of each test was approximately 0.2 mm, except for the 11 kN tests, where it was noticeably higher.

Figures 3(a)–(c) presents the stress–position hysteresis loops for three additively manufactured NiTi lattice samples (Samples 4, 6, and 10), tested under cyclic compressive fatigue loading at maximum force levels of 9 kN, 8 kN, and 11 kN, respectively. Selected cycles are shown to illustrate the evolution of mechanical behavior throughout fatigue life. The hysteresis loops are plotted using stress versus crosshead displacement, rather than true strain, as direct strain measurements were not available. In this study, strain was approximated from global displacement. Whereas this approach introduces some limitations due to machine compliance and boundary effects, the stress–displacement curves still provide meaningful and consistent insights into energy dissipation and deformation trends across samples.

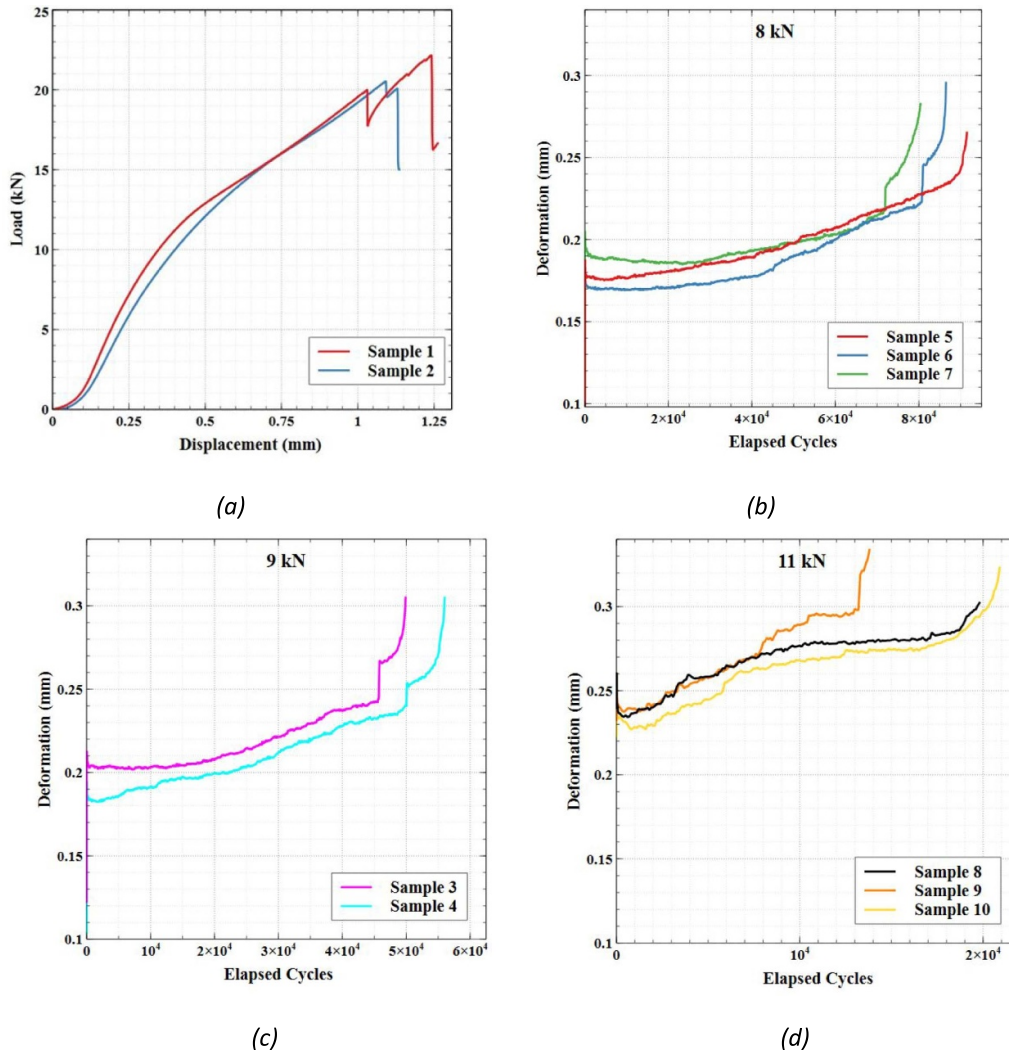


Figure 2. (a) Compression test results for Samples 1 and 2; (b)–(d) deformation per cycle until failure for samples tested under different loads.

In all cases, the applied force range was held constant during testing. Since the cross-sectional area of the samples remained unchanged, the peak compressive stress values remained stable over the fatigue life. This indicates that the samples maintained their load-bearing capacity and exhibited a stable superelastic response during the initial and intermediate stages of cycling.

Despite the differences in load levels, all samples demonstrated similar cyclic behavior. A gradual accumulation of residual deformation was observed over the course of the tests, accompanied by a progressive reduction in the slopes of both the loading and unloading curves—indicative of stiffness degradation. These effects became more pronounced in the final cycles before failure, where the hysteresis loops widened and the slopes decreased significantly, reflecting increased energy dissipation and a marked loss of structural integrity. Among the three, Sample 10 (tested at the highest load of 11 kN) consistently showed greater deformation throughout the test, consistent with the higher strain response expected under elevated loading conditions.

The energy dissipated per cycle was estimated by calculating the area enclosed by the stress–position hysteresis loop, with strain approximated by dividing the crosshead displacement by the specimen’s initial height. Figure 3(d) shows the evolution of energy dissipation over the fatigue life for each sample. As expected, the sample tested at the highest load (11 kN) exhibited significantly greater energy dissipation, reflecting increased internal deformation and energy loss mechanisms. In contrast, samples subjected to lower loads (8–9 kN) displayed lower and more stable dissipation trends, consistent with fatigue behavior dominated by elastic deformation. Some scatter and early increases in dissipation, particularly in Sample 9, may be attributed to microstructural settling or the onset of early damage accumulation. Overall, the consistency across samples supports the validity of the measurement approach for comparative analysis.

Figure 4(a) shows the average number of cycles to failure and the corresponding average displacement at failure for each load. The data indicates that as the maximum applied load decreases, the average number of cycles to failure increases

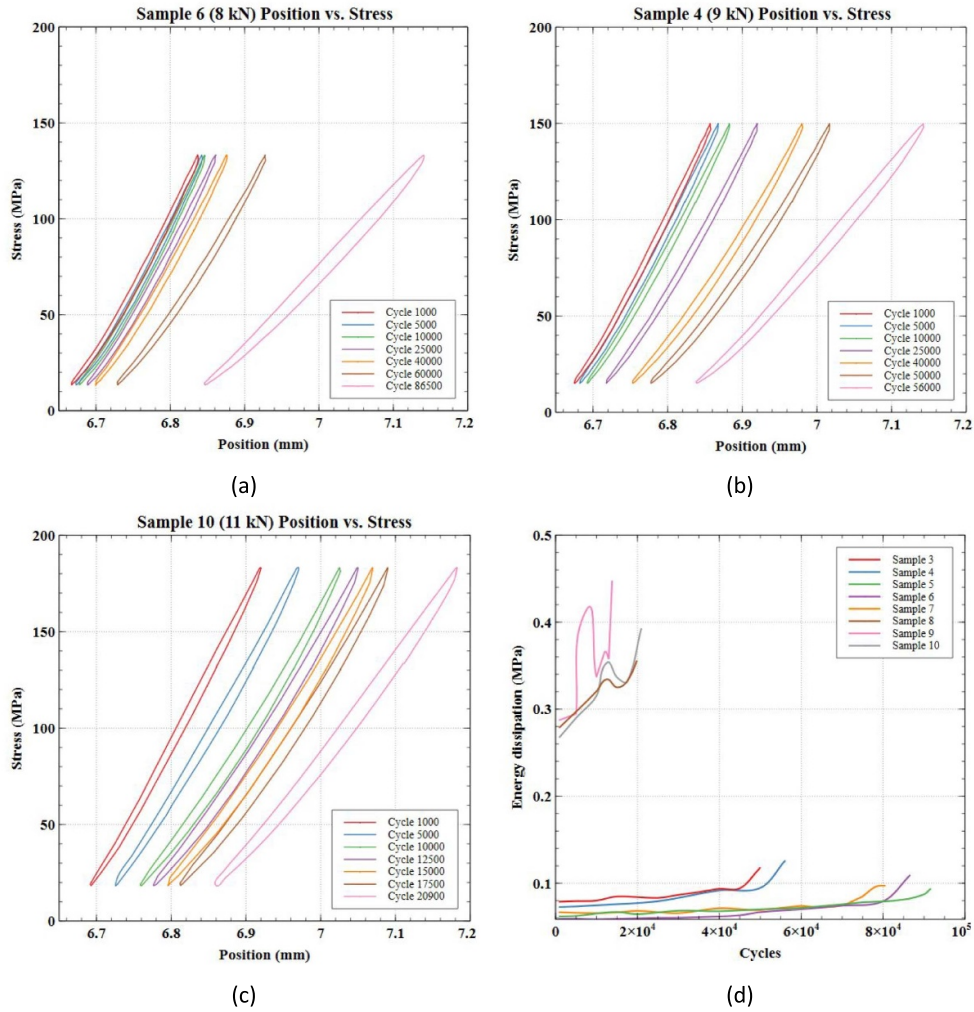


Figure 3. Stress—position hysteresis loops for the NiTi samples under different loading conditions: (a) 8 kN; (b) 9 kN; (c) 11 kN; and (d) energy dissipation over fatigue cycles for each sample.

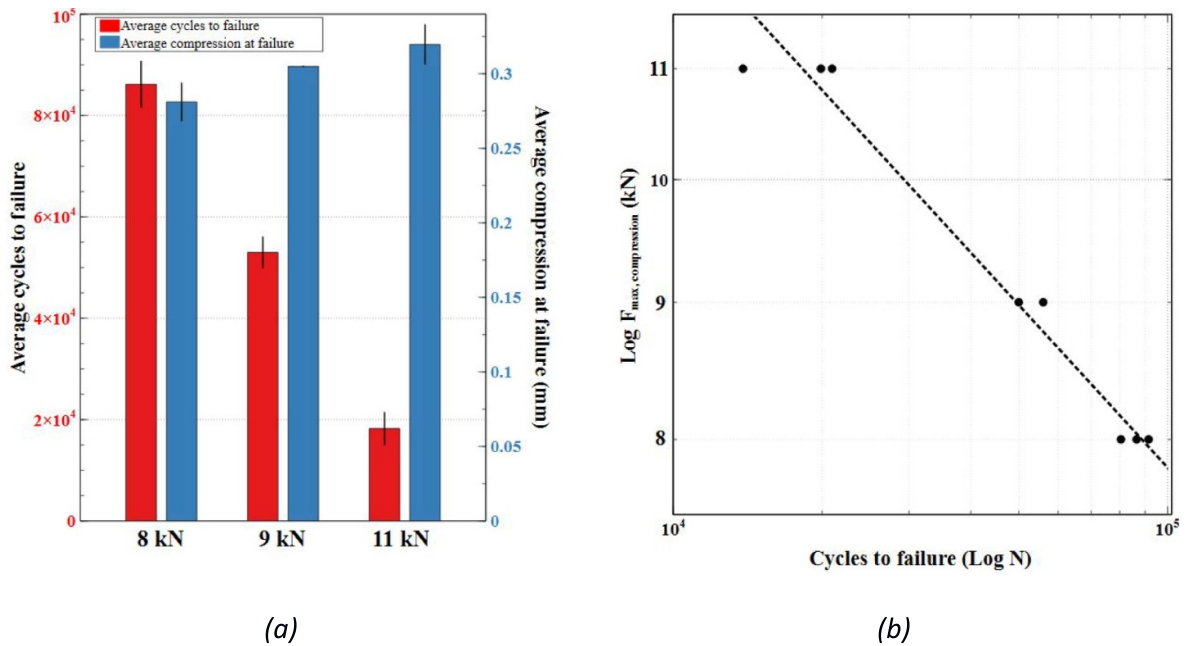


Figure 4. (a) Average cycles to failure and average compression at failure for each load; (b) S–N curve for fatigue tests on lattice samples under 8, 9, and 11 kN loads.

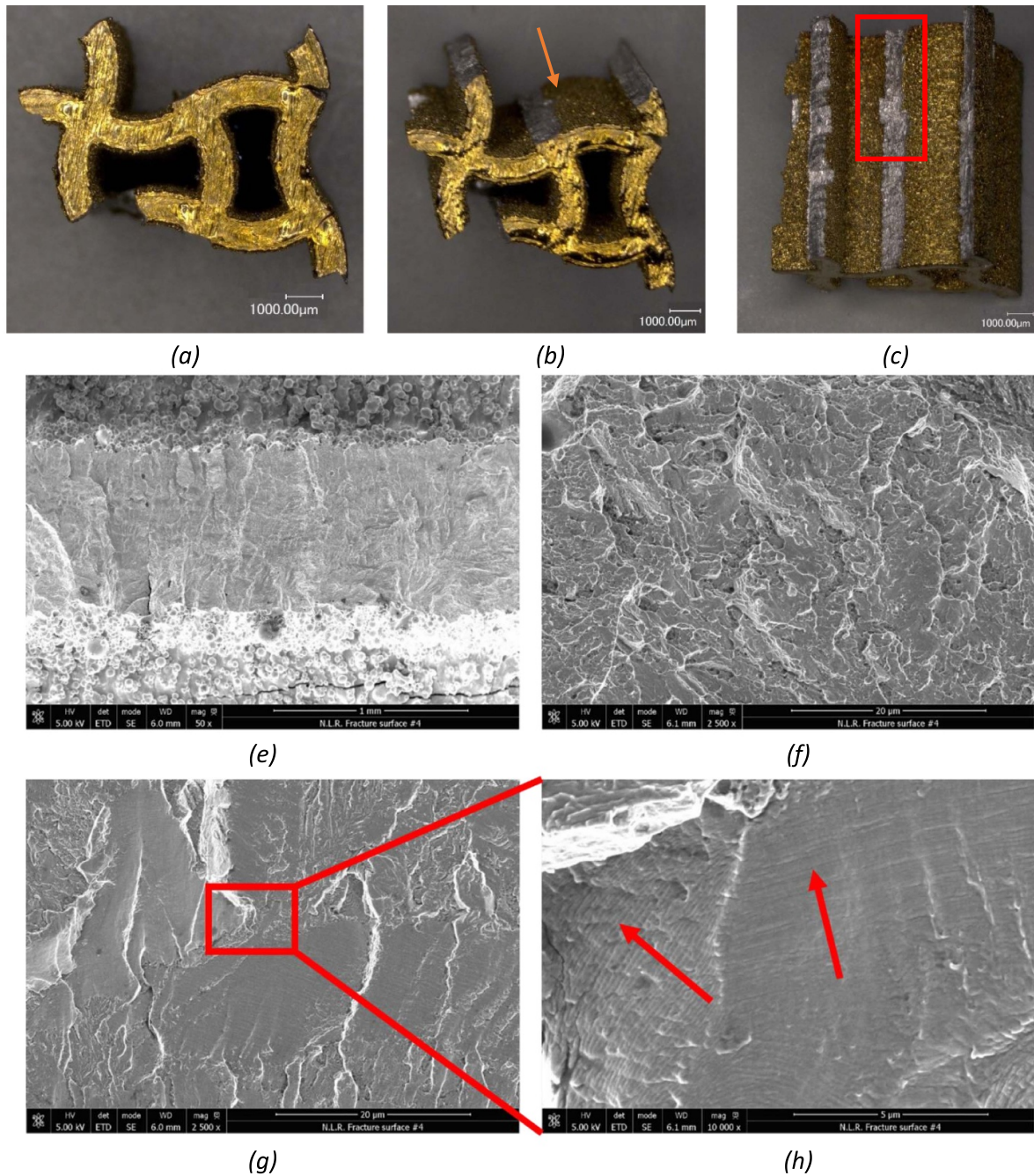


Figure 5. OM and SE-SEM images of the NiTi architected sample (Sample 4): (a) top view, (b) isometric view with an arrow indicating the fractured region for further investigation, and (c) lateral view highlighting the fractured area, with a red box marking the region for high-magnification observation. SE-SEM images of the fracture surface at increasing magnifications: (e) overview of the fractured region, (f) close-up of selected areas showing dimples and smooth subsurfaces, and (g) magnified view of the red box in (h), highlighting the striations.

from 18 000 to 86 000, while the average displacement at failure slightly decreases from 0.32 mm to 0.28 mm. The figure also illustrates the deformation progression per cycle until failure, with each graph corresponding to a specific applied load. Deformation gradually increases with each cycle, followed by a sudden spike, indicating structural failure.

Figure 4(b) presents the fatigue test results in an S–N curve, where the y-axis represents the logarithm of the maximum compressive force, and the x-axis represents the logarithm of

cycles to failure. The data exhibits a strong correlation, fitting well to a power-law trend on a log–log scale.

Since the MT in NiTi SMAs can be triggered by stress, the evolution of the microstructure near the selected fracture area was examined to better understand the damage mechanisms in LPBF-fabricated NiTi lattice structures.

Figures 5(a)–(c) display OM images of a lattice sample at the end of the fatigue test. These images show a portion of the fractured sample, where a ligament detached due to exceeding

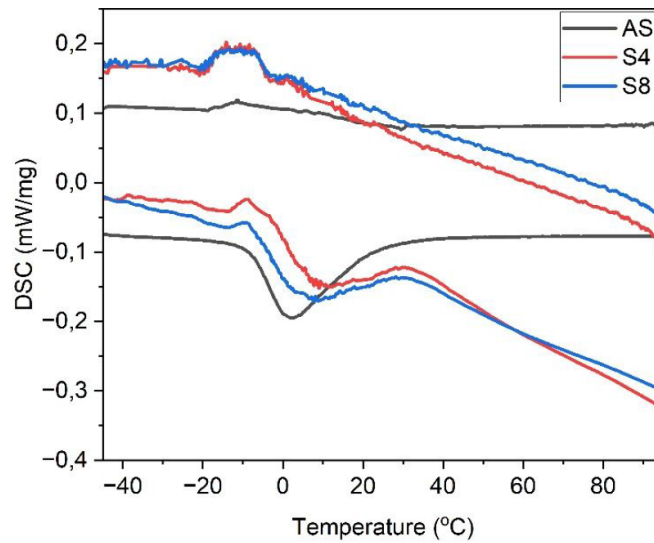


Figure 6. DSC scans of NiTi architected structures in three conditions: as-built (A_S) sample before fatigue testing, and fractured regions after testing for Samples 4 and 8.

the material's maximum load. A high-magnification SEM analysis was performed on the fracture region highlighted by the red box in figure 5(c), using the secondary electron (SE) signal.

Figure 5(e) presents an overview of the fracture surface from figure 5(c), showing remnants of unmelted powder on the outer surface. The fracture surface exhibits two main regions: an irregular surface and a smooth one, both typical of AM parts subjected to fatigue. The irregular surface (figure 5(f)) contains dimples and is associated with the early failure of adjacent material, leading to an overloaded area. In contrast, the smoother surface (figure 5(g)) displays ductile failure behavior. Continuous load cycling led to the formation of fatigue striations, visible in the high magnification shown in figure 5(h). These striations follow inclined directions, as indicated by the red arrows in figure 5(h).

Figure 6 presents the DSC results for the fractured regions of the NiTi lattice samples, specifically Samples 4 and 8, along with the as-built (A_S) sample, which was tested before fatigue. These samples were chosen for comparison due to their different load conditions and fatigue responses: Sample 4 underwent a lower load with a higher number of cycles to failure, while Sample 8 was subjected to a higher load, leading to fewer cycles to failure.

The evolution of the transformation temperatures for the three samples is summarized in table 2. The austenite finish temperature (A_f) for both fractured samples was found to be higher than that of the base material, while the other transformation temperatures were not significantly influenced by the sample condition. In contrast, the applied load during cyclic compression testing did not cause any notable changes in the transformation temperatures.

The transformation temperatures of NiTi alloys can shift due to microstructural changes such as dislocation accumulation and phase transformation fatigue. The thermal hysteresis, defined as the difference between the austenite finish (A_f) and martensite finish (M_f) temperatures,

Table 2. Transformation temperatures of the investigated samples.

	M_f	M_s	A_s	A_f
Heat-treated sample	-18.2	—	-8.8	23.9
Sample 4	-19.3	-4.2	-3.6	32.5
Sample 8	-20.2	-3.8	-6.1	30.5

increased after fatigue testing, primarily due to crack formation within the samples. This observation is supported by EBSD results (figure 7), which show that regions where cracks propagated, ultimately leading to failure of the lattice structure, also exhibited alterations in the MT. In particular, high-stress concentrations in these areas led to significant plastic deformation and the formation of stress-induced martensite.

Figure 7(a) presents an SEM image of the region of interest in Sample 4. In the central zone (figure 7(b)), no cracks were observed, and EBSD analysis confirmed that austenite was the dominant phase at room temperature. In contrast, figure 5(d) shows a magnified view of a crack propagating from the outer to the inner part of the NiTi lattice. This region exhibits a clear microstructural transition from austenite to martensite, attributed to localized stress intensification. The dark contrast in this region, typically indicative of martensite, further supports this interpretation, in agreement with previous studies [17].

Thus, elevated stress levels not only triggered microstructural changes but also facilitated crack initiation and propagation, contributing to structural degradation. Although the applied loading was uniaxial and compressive, local tensile stresses can develop due to geometric features or residual stresses, enabling crack formation. This is especially relevant in auxetic lattices, where nonuniform deformation patterns can amplify localized stress states.

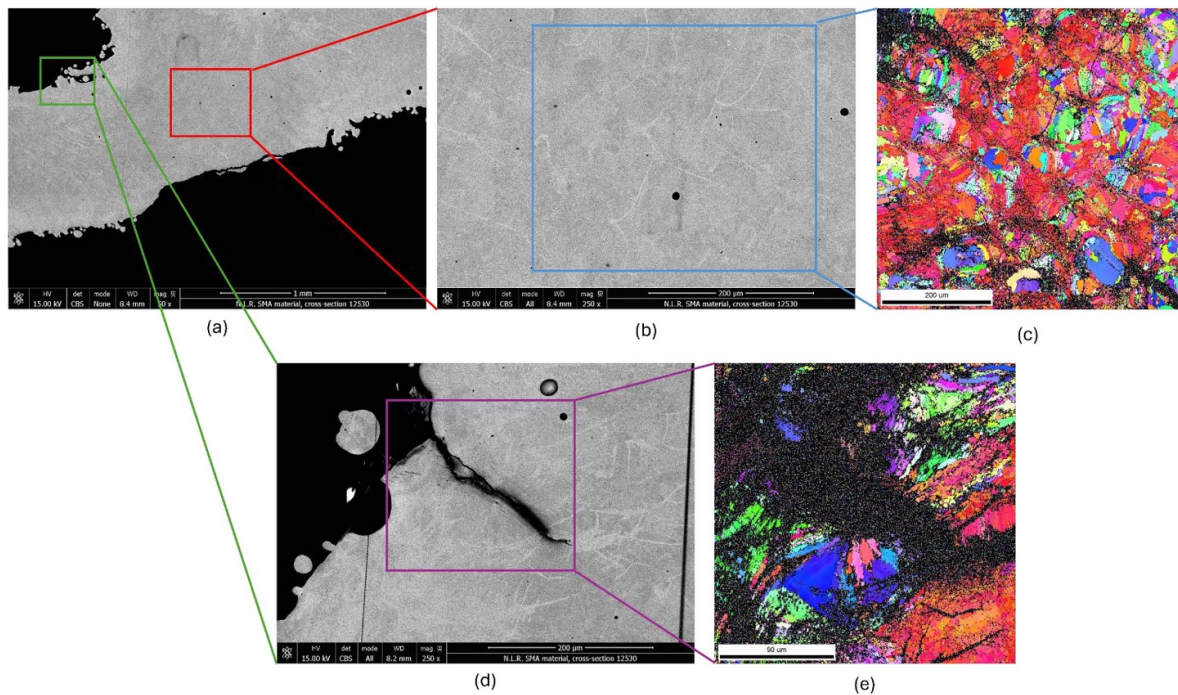


Figure 7. SEM-SE overview of the region near the fracture in Sample 4: (a) overview of the region, (b) magnification of the central part, with (c) its corresponding EBSD map; (d) magnification of a detected crack, with (e) its corresponding EBSD map.

4. Conclusion

This study investigated the fatigue behavior of LPBFed NiTi lattice structures across a broad range of cycles. By performing fatigue tests under varying load levels, an S–N curve following a power-law relationship was developed, revealing a consistent and predictable correlation between applied stress and fatigue life. This reliable fit enables confident extrapolation to different loading conditions, supporting the use of these lattices in cyclic applications.

The findings highlight how fatigue performance is governed not only by macroscopic stress-life trends but also by the complex interactions between phase transformations, lattice architecture, and manufacturing defects. Under higher loads, localized stresses can surpass the transformation threshold, triggering phase changes that affect energy dissipation and damage evolution within the material. The auxetic lattice design, with its negative Poisson's ratio, produces unconventional mechanical responses that cause uneven stress distribution and promote localized failures. Furthermore, manufacturing defects inherent to the LPBF process, especially at lattice corners and junctions, act as critical crack initiation sites, explaining instances of premature fatigue failure.

Together, these insights emphasize the importance of optimizing both design and manufacturing parameters to enhance the fatigue life of NiTi lattice structures for demanding cyclic applications.

Data availability statement

All data that support the findings of this study are included within the article (and any supplementary files).

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