



# Review of defects in lattice structures manufactured by powder bed fusion

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## Abstract

Additively manufactured lattice structures are popular due to their desirable properties, such as high specific stiffness and high surface area, and are being explored for several applications including aerospace components, heat exchangers and biomedical implants. The complexity of lattices challenges the fabrication limits of additive manufacturing processes and thus, lattices are particularly prone to manufacturing defects. This paper presents a review of defects in lattice structures produced by powder bed fusion processes. The review focuses on the effects of lattice design on dimensional inaccuracies, surface texture and porosity. The design constraints on lattice structures are also reviewed, as these can help to discourage defect formation. Appropriate process parameters, post-processing techniques and measurement methods are also discussed. The information presented in this paper contributes towards a deeper understanding of defects in lattice structures, aiming to improve the quality and performance of future designs.

**Keywords** Lattice structures · Manufacturing defects · Additive manufacturing · Powder bed fusion

## 1 Introduction

Cellular solids are defined by Gibson and Ashby as ‘an interconnected network of solid struts or plates which form the edges and faces of cells’ [1]. Cellular solids can take either stochastic or non-stochastic forms, depending on the arrangement of their unit cells. A stochastic design, often called a foam, is controlled globally, often through the manufacturing process, for example, introduction of a gas into a molten material [2]. Stochastic designs, therefore, have high local variability, and the design is controlled only at the macroscale. Non-stochastic arrangements,

however, often called lattice structures, are highly ordered; full control over individual unit cells allows for greater control of the structure’s properties.

Cellular solids possess a range of desirable properties, such as high specific strength and stiffness, and high surface area to volume ratio [3], and exist in a large number of forms. Examples can often be observed in nature, for example the structure of wood (Fig. 1a), cork and trabecular bone. Non-stochastic examples include the highly ordered arrangements in the atomic structures of crystalline materials. Macroscale examples include truss structures in cranes, bridges (Fig. 1b) and roofing. Between these two scales, so-called mesoscale cellular solids have unit cell sizes ranging from approximately 0.1 mm to 10 mm [4, 5] (Fig. 1c, d).

Recent developments in additive manufacturing (AM) processes, particularly powder bed fusion (PBF), have seen the number of applications for cellular solids rise in a number of disciplines. Engineering applications include heat exchangers [6–9], which exploit the high surface area to volume ratio to maximise heat transfer, and sandwich structures for aerospace applications, which require lightweight, high-strength materials [10, 11]. There is also significant interest from the biomedical field, where biocompatible materials are used to develop cellular solids imitating bone tissue, thus enabling the development of bespoke biomedical implants [12–17].

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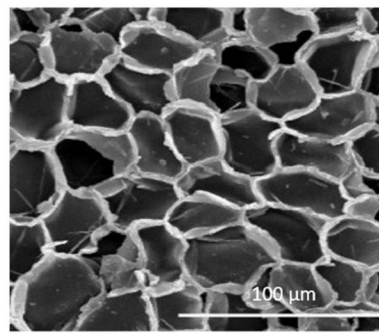
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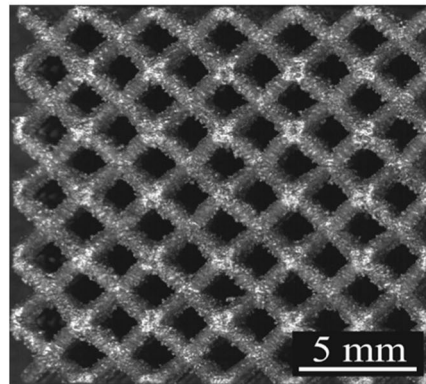
**Fig. 1** Examples of cellular solids: **a** scanning electron microscopy image of cork from the bark of *Quercus cerris* var. *cerris* [19], **b** a crane (upper) and a bridge (lower) showing examples of truss structures [20], **c**, **d** cellular solids produced by powder bed fusion processes: **c** non-stochastic AlSi10Mg body centred cubic structure [21], **d** stochastic Ti6Al4V structure [22]



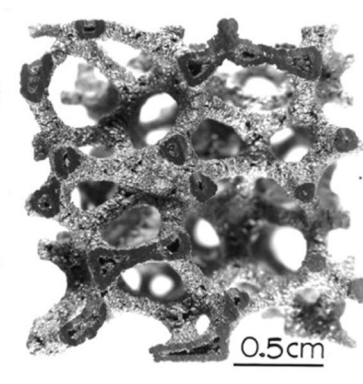
(a)



(b)



(c)



(d)

Cellular solids produced by PBF processes are prone to defects, due to their numerous overhanging, often millimetre-scale features, which challenge the performance limits of PBF systems. Support structures are required to dissipate heat and support overhanging features, however, in PBF, cellular solids prevent the use of support structures as post-manufacture removal is impractical. Therefore, metallic cellular solids produced by PBF are particularly plagued by defects due to higher processing temperatures. Manufacturing defects, that is any measurable deviations between an initial design and the manufactured product, can significantly hinder the desired operating performance of AM parts, for example through the introduction of stress concentrations causing a reduction in fatigue strength [18]. Therefore, particularly considering the load-bearing applications of AM cellular solids, a thorough understanding of manufacturing defects is required and is presented in this review.

### 1.1 Contents of the review

This review paper is focused on the literature regarding manufacturing defects in non-stochastic, metallic, cellular solids produced by PBF, which will hereafter only be referred to as lattice structures. Additionally, only mesoscale lattice structures will be reviewed. Such a scope is justified due to the prevalence of these structures in the literature related to

stochastic and/or polymeric forms and alternative manufacturing methods. Mesoscale lattice structures are well suited for the feature resolution of PBF systems and the control afforded to the designer yields a greater range of applications, as reflected in the introduction. Helou and Kara [23] showed laser melting to be the most common technique for lattice structure manufacture, reviewing literature between 1999 and 2016, with Ti6Al4V and 316L stainless steel the two most common materials. Furthermore, searching keywords on Scopus such as ‘lattice structure’, ‘stochastic’, ‘additive manufacturing’, ‘laser melting’ and taking from the most recent results in 2019 showed 85% (of 13 results) were related to non-stochastic lattice structures. Additionally, the most cited papers on lattice structures for additive manufacturing also consistently focus on non-stochastic metallic forms (for example, [24–26]).

There are existing reviews in the field of cellular solids which focus on reporting their design, analysis and manufacturing methods [2, 4, 23]. This paper considers defect formation from the perspective of lattice design, as opposed to process control, which is the focus of other AM reviews, such as [18, 27, 28]. To refine the discussion on lattice structure defects, several questions were posed, as shown below.

1. How do defects form in powder bed fusion processes?
2. What type of defects form in lattice structures?

3. How can defects be appropriately measured in lattice structures?
4. How can defect formation in lattice structures be minimised?

Firstly, an introduction to lattice structures is provided in Sect. 2, and the terminology used in this review is established. Subsequent sections aim to address the above questions, where Sect. 3 provides a description of the PBF process and a brief summary of some existing reviews on general defect formation mechanisms. Section 4 discusses the defects which have been observed specifically in lattice structures, followed by measurement methods for quality inspection and defect mitigation in Sects. 5 and 6, respectively. To close the review, a summary of the findings is provided along with suggestions on potential future directions. Addressing the above questions aims to offer a well-rounded literature search by providing not just a description of defects in lattice structures but also detailing how the quality of lattice structures has been improved and inspected more appropriately.

This literature review was performed using the main scientific databases (Scopus, Google Scholar) with publication dates ranging from 1994 to 2019. Papers were selected if the contents of the study contributed towards answering any of the above questions.

## 2 Lattice structures

Lattice structure designs are defined by their unit cell, which often falls into one of two categories: strut and surface based (Fig. 2). Strut-based unit cells consist of a network of often prismatic struts (i.e. constant cross section) connected at nodes, similar to truss structures. Surface-based unit cells are mathematically defined as the surface connecting set of points for which a given function has a constant value, that is, an isosurface.

Unit cell design holds large influence over the mechanical properties of a lattice structure. In strut-based unit cells, the connectivity of the struts, that is the number of struts connected at a given node, greatly controls the structure’s behaviour under compressive load. For low or high connectivity, unit

cells exhibit what is known as bending- or stretch-dominated behaviour, respectively. Bending-dominated structures are desired for applications where mechanical energy absorption is required. Stretch-dominated unit cells are preferred for structural applications, due to greater stiffness and yield strength [29]. Common examples of strut-based lattices are the body- (BCC) and face-centred cubic (FCC) structures (Fig. 2a, b) which are bending and stretch dominated, respectively.

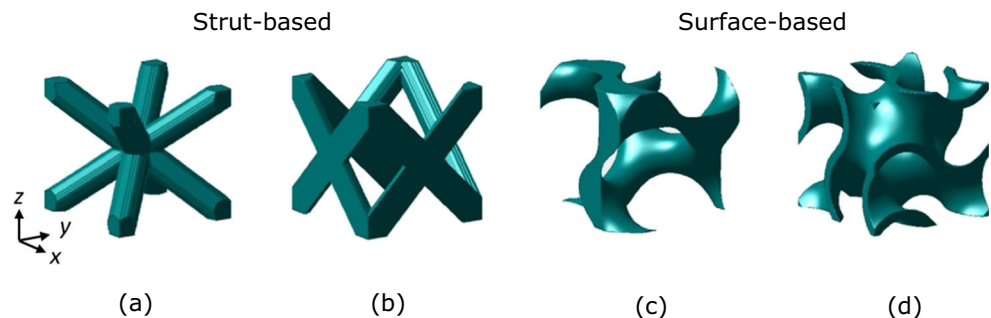
The most common types of surface-based unit cells are the triply periodic minimal surfaces (TPMS), which have a mean curvature of zero. An example of a surface-based unit cell is the gyroid, developed by Schoen [30] (Fig. 2c, d). The gyroid is defined by the isosurface  $\phi$  in three dimensions ( $x, y, z$ ):

$$\phi = \cos(x)\sin(y) + \cos(y)\sin(z) + \cos(z)\sin(x) = 0. \quad (1)$$

TPMS unit cells can be further distinguished between those with isosurfaces that have been thickened (sheet/matrix-TPMS), or their enclosed volume solidified (skeletal/network-TPMS), as shown in Fig. 2c, d. Some skeletal/network TPMS designs appear to possess strut-like features, as is the case for the skeletal gyroid (Fig. 2c). The user has less design control with TPMS unit cells, however they are currently studied for applications such as biomedical implants [12]. Additionally, there are custom designs outside of strut-based and surface-based forms, for example those produced using topology optimisation [31] and with internal resonators for vibration isolation [32].

A key property of a unit cell is its relative density, which is defined as the ratio of the density of the unit cell to that of the comprising material [1]. Therefore, the density of the unit cell, often called volume fraction (see Table 1), is a design feature of key importance, largely controlling the mechanical properties of the structure. Additional terms for lattice design features are listed in Table 1. Some definitions are described in the literature using different terms, for example volume fraction can be referred to as ‘porosity’, particularly in biomedical applications where the designed void space in a unit cell is required for bone ingrowth. Therefore, for consistency in this review, the terms in Table 1 will be adhered to. Note that Table 1 is not an exhaustive list, and the references serve only as examples of where such terms can be found within the papers in this review.

**Fig. 2** Types of unit cells: **a** body-centred cubic, **b** face-centred cubic, **c** gyroid (skeletal/network) and **d** double gyroid (sheet/matrix) [33]



**Table 1** Lattice terminology

Terminology in this paper	Definition	Alternative terms in literature
Lattice structure	Mesoscale, non-stochastic, cellular solid	Porous structure [34, 35], microlattice [5, 9], scaffold [16, 36]
Periodicity	Unit cell tessellations in $(x, y, z)$	Staggering [37]
Node	The intersection of multiple struts (or strut-like features, in the case of some surface-based cells)	Strut junction [38, 39], truss corners [40]
Volume fraction	Ratio between the volume of material in the unit cell and its bounding box (often expressed as a percentage)	Porosity [41, 42], solid fraction [43]
Pore/porosity	Unintentional void within the lattice structure/ratio of pore volume to material volume	n/a <sup>a</sup>
Designed pore/ designed porosity	Void space included in the unit cell design/ratio of designed pore volume to material volume (i.e. the complement of volume fraction)	Pore/porosity [36, 42]

<sup>a</sup>No alternative terms found from the papers in this review

### 3 Powder bed fusion

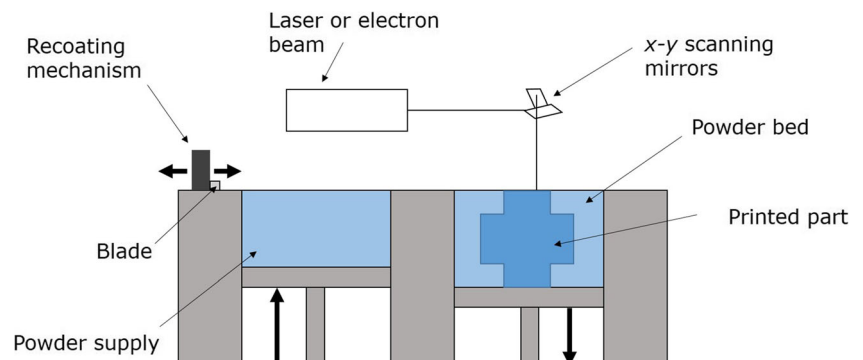
PBF operates by using focused thermal energy to selectively fuse regions of a powder bed [44], with potential to create highly intricate geometries, with millimetre scale features. Figure 3 is a schema of the PBF process, which can be summarised in the following steps. A layer of powder (typically of thickness 20  $\mu\text{m}$  to 150  $\mu\text{m}$ ) is spread across the build bed. A thermal energy source is focused onto the powder bed and fuses the predefined regions. The part is then lowered by one-layer thickness and the process repeats until completion. Within PBF, there are three main processes which share this operating procedure: laser sintering (LS), laser melting (LM) and electron beam melting (EBM). Laser sintering is used to process polymer powders while laser melting and electron beam melting process metal powder. The main difference between polymer and metal processing is in the input energy, where higher energies are required to produce full melting for metal powders. LM and EBM differ fundamentally in energy source, and this difference yields several effects, such as a reduced feature resolution in EBM due to defocusing of the beam to prevent a build-up of negative charge [45]. Within metal PBF processes, support structures are

often required to attach the part to the build bed in order to prevent distortion from residual stresses induced through high thermal gradients and to support overhanging features that would otherwise distort under gravity [45].

PBF process parameters can be grouped into four categories [45]: thermal source parameters, scan parameters, powder parameters and temperature-related parameters. The thermal source parameters, such as laser power, most directly control the fusing of powder material and, in the case of full melting, directly affect the melt pool characteristics. Scan parameters control the manner in which material is fused, for example scan speed and scanning patterns which can control surface finish. Powder parameters such as powder size and layer thickness can control the absorption characteristics of the powder bed and the thickness of each layer. Temperature related parameters, such as powder bed temperature, have a strong influence on the repeatability of the manufacturing process, where constant temperatures encourage uniformity throughout the build bed and between batches of production.

The term energy density  $ED$  defines the relationship between primary process parameters and provides a useful means for selecting optimal processing parameters.  $ED$  is defined as:

**Fig. 3** Schema of the powder bed fusion process





$$ED = \frac{P}{vht}, \quad (2)$$

where  $P$  is laser power,  $v$  is scan speed,  $h$  is hatch spacing (the spacing between scanning paths) and  $t$  is layer thickness [46]. These parameters are often used to investigate the effects of process parameters on print quality (for example, [47, 48]).

### 3.1 Defect formation mechanisms

This section provides a brief background on the defects commonly formed in PBF processes, providing useful context to the following sections which discuss defects found specifically in lattice structures. The heating, melting and cooling mechanisms within the powder bed are at the core of the final quality of the parts produced by PBF and are discussed within several review papers on defects in PBF processes. From [18, 27, 49], three main categories of defects were identified: porosity/incomplete fusion, residual stresses and surface defects.

Porosity is defined by Zhang et al. [27] as approximately spherical voids with diameter typically less than 100  $\mu\text{m}$  (Fig. 4a). Pores can form when gases within the melt pool are entrapped due to high cooling rates (up to  $10^8$  K/s). The entrapped gases initially form either from gaps within the powder bed or through evaporation of lower melting point constituents within the powder material. Pores can also form through the use of hollow powder particles and through ridges formed in previous layers which impede the flow of the melt pool. Incomplete fusion defects are explained by Zhang et al. [27] to occur mainly due to insufficient input energy and thus often form as irregular voids containing un-melted powders (Fig. 4b). Incomplete fusion defects can form through consecutive layers, again as melt pool flow is impeded. For easily oxidised materials, (for example AlSi10Mg) the formation of oxide layers reduces wettability, which also impedes melt pool flow and causes additional incomplete fusion.

Residual stresses are formed through PBF processes due to the high thermal gradients and cooling rates within the powder bed subjecting the part to rapid expansion and contraction. Residual stresses can cause cracks to form [27], delamination between layers [18] (Fig. 4c) and dimensional inaccuracies through shrinkage or warping [18]. Cracking is more likely to occur in materials such as stainless steels and nickel-based superalloys, due to their low thermal conductivity and high coefficient of thermal expansion [27]. Mercelis and Kruth [50] define two mechanisms through which residual stresses are induced. Firstly, the temperature gradient mechanism, where upon heating the part, expansion is hindered by the solidified material beneath, creating compressive residual stresses. Secondly, upon cooling of molten layers, contraction is hindered, creating tensile residual stresses. Mercelis and

Kruth note that the temperature gradient mechanism does not require the material to be molten and can, therefore, occur in both LS and LM, whereas the cooling mechanism within the melt pool only occurs in LM.

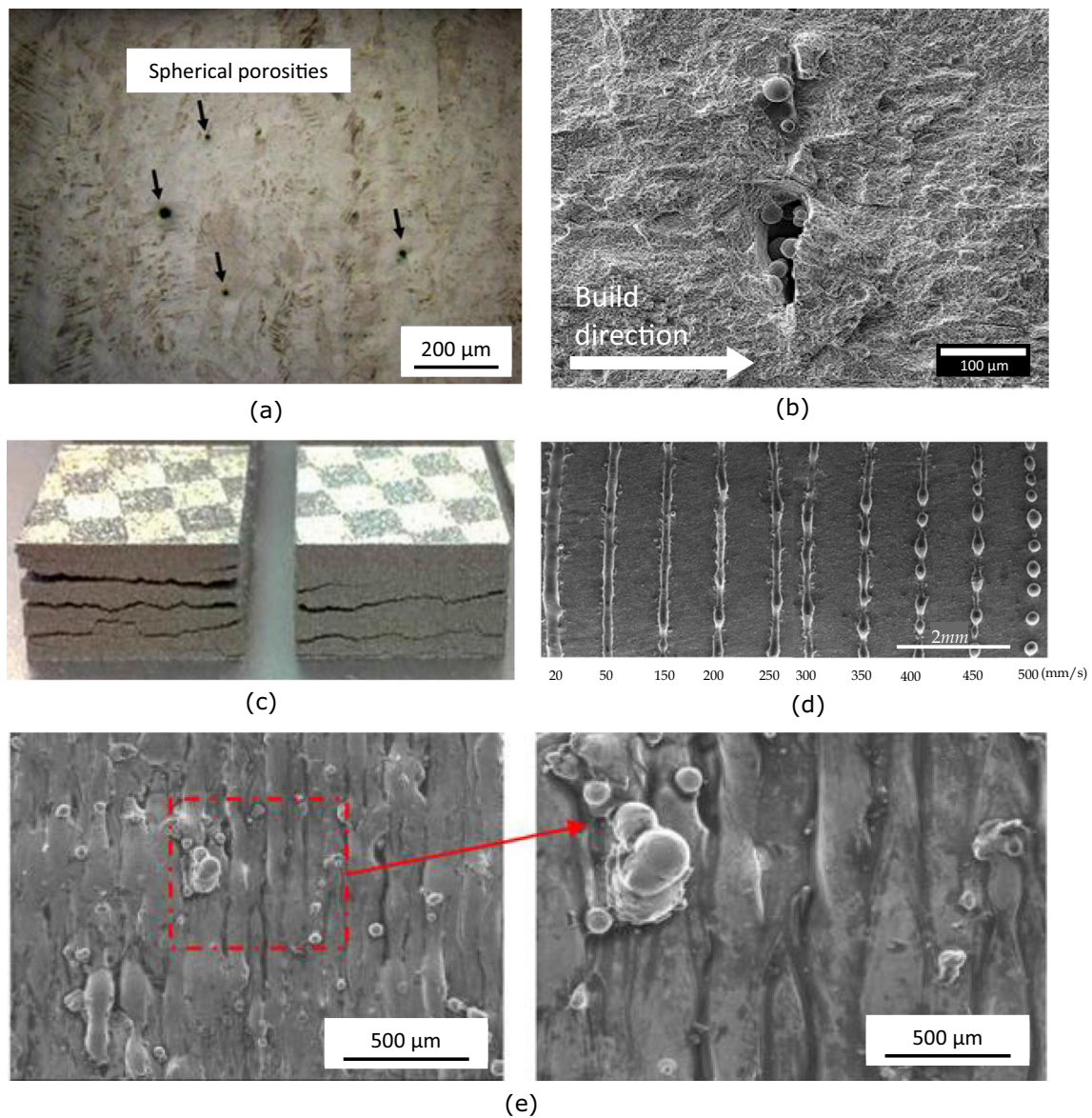
Highly complex surfaces are produced in PBF processes where the interaction between the powder material and energy source forms a number of surface irregularities. Unstable melt pools can cause balling and spatter. Balling occurs when the melt pool solidifies into a sphere instead of wetting the underlying substrate to form a layer—mainly controlled by the length-to-width ratio of the melt pool, where a ratio less than 2.1 is desirable [49] (Fig. 4d). Spatter occurs at higher energy densities, where molten material is expelled by the recoil pressure generated by evaporation within the melt pool (Fig. 4e). Inter-layer surface defects are also present, where discontinuities form between successive layers, i.e. stair stepping. The orientation of a surface further affects the quality. Surfaces whose normal is directed towards the build bed (called down-skin surfaces) produce significantly more irregular surfaces than those whose normal faces away (i.e. up-skin). This discrepancy is due to the support powder, which is a poor conductor of heat. Therefore, down-skin surfaces experience excess heating and poor cooling through support powder, whereas up-skin surfaces cool faster via conduction through the solidified part underneath [18].

## 4 Lattice structure defects

This section follows on from the previous discussion of defect formation and aims to establish the types of defects that form specifically in lattice structures, with a focus on how design features influence defect formation. In the literature, lattice structure defects are frequently categorised into the following groups: dimensional inaccuracies, surface texture and porosity. Dimensional inaccuracies occur on the same scale as the lattice features, for example a deviation in the length or overall volume. Surface defects refer to any deviation from the ideal surface and are often quantified using surface texture parameters, providing a quantitative description of surface texture. Porosity denotes the absence of material, where a pore describes a closed void within the part, or a surface void. When necessary, measurement methods will be mentioned to provide context, and will be covered in more detail in Sect. 5.1.

### 4.1 Dimensional inaccuracies

It is often helpful to first globally assess a lattice structure's general conformity to CAD data, as performed by [31, 36, 56–58], where the results generally show a reasonable adherence to CAD data. However, dimensional inaccuracies can be



**Fig. 4** Examples of the defects which form in PBF processes: **a** porosity [51] (annotations from [18]), **b** incomplete fusion [52], **c** severe cracking causing delamination [53], **d** balling occurring at higher scanning speeds [54] and **e** spatter [55]

better understood via a local analysis of specific lattice features, such as struts, nodes, designed pores and wall thicknesses. Struts have been observed to deviate from circular cross-sections to ellipsoidal in references [59–62]. Ataei et al. [61] particularly note the effect to be strongest at the ends of the struts. Arabnejad et al. [59] observe ellipsoidal deviations to a greater degree in horizontal struts, due to overmelting.

The average diameter of lattice struts has been observed to significantly deviate from the initial design, as shown in Table 2, where both over- and under-sizing have been

observed. Cuadrado et al. [63] and Arabnejad et al. [59] show strut diameter to be significantly affected by strut orientation, where vertically oriented struts were under-sized by up to 45%, and horizontal struts were over-sized by over 100%. Cuadrado et al. [63] linked deviations in strut diameter to the overall volume fraction of lattice structures, observing a volume fraction less than designed in lattice structures consisting of a significant portion of vertical struts, with the opposite occurring in designs with more horizontal struts. Zhang et al. observed similar orientation-dependent thickness variations in sheet TPMS unit cells (primitive, diamond, gyroid) [60].

**Table 2** Deviation in strut diameter

	Designed strut diameter (mm)	Strut deviation (mm)
Sing et al. [74]	0.6 to 0.8	− 0.5 to − 0.2; up to ~ 60%
Tancogne-Dejean et al. [71]	0.534	Average strut deviation, − 0.01 (~ 2%)
Choy et al. [75]	0.4 to 1.2	− 0.138 to + 0.156 (− 10% to + 40%)
Cuadrado et al. [63]	0.65	Vertical, 0.46 (− 30%) sloped, 0.36 (− 45%) horizontal, 1 (+ 54%)
Arabnejad et al. [59]	0.2	Vertical, 0.11 (− 45%) sloped, 0.26 (+ 30%) horizontal, 0.45 (+ 128%)

Melancon et al. [42] and Liu et al. [64] observed deviations in the position of struts' axes, defining this deviation as 'strut waviness' (Fig. 5a). As with variations in diameter, strut waviness was observed to occur to a greater degree in horizontal struts. Liu et al. calculated an increase in strut waviness from approximately 8% to 17% in diagonally oriented struts and horizontal struts, respectively. Melancon et al. also calculated an increase, to a lesser degree, of 7% to 9% in diagonal and horizontal struts, respectively. Similar observations on strut orientation dependency are also observed in references [60, 65–67].

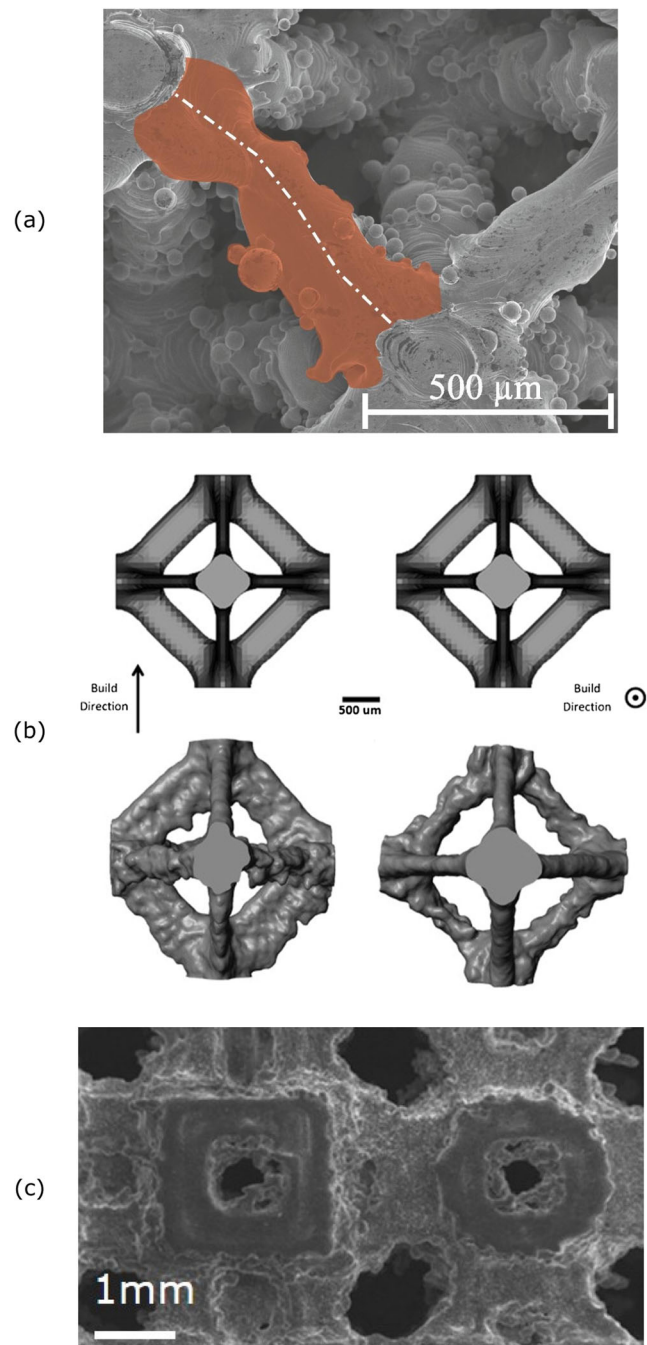
The effects of strut orientation dependency are highlighted by Sercombe et al. [43], who found horizontal struts to cause lattice structures to fail under compression, as the horizontal struts carried tensile loads (Fig. 5b).

Several defects have been observed around lattice nodes. Excess material accumulation at nodes has been observed by Gümriük et al. [68]. Similar results were seen by Li et al. [69], where strut diameters increased by approximately 30% around the nodes (from 0.3 mm to a maximum of 2 mm). Al-Ketan et al. [70] compared several unit cell designs and found nodes to be smoother in surface-based unit cells. Additional comments on nodal distortions are made in references [71, 72].

Regarding designed pores, Hollander et al. [73] and Lin et al. [31] both observed material accumulation called 'surface protrusions' (Fig. 5c)—the most severe of these extended across 0.3 mm (30%) of the designed pore diameter [73]. Obstructions within designed pores can impede bone ingrowth in lattice structures for biomedical applications.

## 4.2 Surface defects

As previously discussed, highly complex surfaces are produced in PBF processes and surface defects are often formed due to melt pool instabilities. The most prevalent surface defects in lattice structures can be observed when comparing down-skin and up-skin surfaces. Local over-heating in down-skin surfaces is often observed due to the many overhanging features in lattice structures [16, 43, 56, 69, 70, 76, 77]. Surface texture parameters, as defined by ISO 4287 [78],



**Fig. 5** Examples of dimensional inaccuracies in lattice structures. **a** An example of strut waviness. SEM micrograph provided by ref. [42]. Varying strut diameter can also be observed. **b** Example of strut orientation-dependency. Top, CAD representation of unit cell. Bottom, 3D rendering of an X-ray computed tomography model. Horizontal struts are shown to be highly irregular [43]. **c** Example of surface protrusions in designed pores [31]

have been used by Pyka et al. [79] and Leary et al. [72] to quantitatively analyse up-skin and down-skin lattice structure surfaces, as shown in Table 3. Pyka et al. found the  $R_a$  value of down-skin surfaces to be approximately twice that of the up-skin surface. Leary et al. calculated  $R_a$  and  $R_z$  values for down-skin and up-skin surfaces in BCC and FCC struts, which have



**Table 3** *Ra* values for up-skin and down-skin lattice surfaces

	Surface roughness parameter	
	Up-skin ( $\mu\text{m}$ )	Down-skin <i>Ra</i> ( $\mu\text{m}$ )
Pyka et al. [79]	<i>Ra</i> = 8	<i>Ra</i> = 15
Leary et al. [72]	45°: <i>Ra</i> = 8.9, <i>Rz</i> = 60.5; 35°: <i>Ra</i> = 8.8, <i>Rz</i> = 53	45°: <i>Ra</i> = 43.5 <i>Rz</i> = 235.5; 35°: <i>Ra</i> = 120.2, <i>Rz</i> = 456.5

inclination angles to the build bed of approximately 35° and 45°, respectively. Leary et al. calculated an *Ra* value of down-skin BCC struts to be almost three times that of FCC. The *Rz* values for BCC down-skin surfaces was also significantly larger than for BCC. The *Ra* values for up-skin and side surfaces showed little variation upon changing orientation.

Al-Ketan [70] compared the surfaces of strut- and surface-based unit cells, and observed the stair-stepping effect to be less pronounced in surface-based unit cell—an example is shown in Fig. 6. A reduced stair-stepping effect has been attributed to the continuous change in inclination angle in surface-based designs (further discussed in Sect. 6.1). Other supporting comments on the texture of lattice surfaces can be found elsewhere [11, 36, 37, 41, 57, 58, 68, 71, 73, 80–87].

### 4.3 Porosity

Pores are typically referred to as enclosed voids beneath the part's surface, however, more regular, hemispherical voids can be identified as surface pores. Under load, pores form stress concentrations which can have a significant effect on fatigue properties.

Low levels of porosity (approximately 1%) are achievable in AM processes, as in [88, 89], and as expected, similar results are found among lattice structures [24, 56, 57, 71, 75, 87, 90–92]. Process parameters are often the focus of the cause of porosity, as in references [89, 93, 94]. In lattice design, however, Yan et al. [24] observed porosity to increase up to approximately 10% upon increasing the size of gyroid unit cells (2 mm to 8 mm). This was attributed to the longer scanning

paths required for larger unit cells which allows a greater period of time for pores to form in between adjacent scanning tracks—as constant process parameters were used for the different cell sizes. Yan et al. highlight that further optimisation of the parameters depending on unit cell size should alleviate any increase in porosity. Amani et al. [95] also attribute scan strategy to an increase in porosity, where pore size was observed to increase at the nodes of lattices (Fig. 7).

Dong et al. [65] investigated the influence of strut orientation on porosity for AlSi10Mg lattices. Struts were printed at varying inclination angles from the build bed and each strut was partitioned into an upper and lower region (A and B as shown in Fig. 8). The results (Fig. 8) showed overall porosity to increase as inclination angle decreased. Furthermore, pore distribution was significantly skewed as the majority of the pores were located in the lower regions of the struts, with this effect increasing upon decreasing inclination angle. These observations were attributed to the cooling mechanisms, where inclined struts are subject to lower cooling rates due to direct contact with support powder, which yields higher temperatures and thus increases the formation of hydrogen pores. The differences in porosity were reflected in the tensile mechanical properties of the struts, for example a 22% increase in ultimate tensile strength was observed between struts printed at 35.5° and 90°. Similar observations were made by Delroisse et al. [96], also for AlSi10Mg struts.

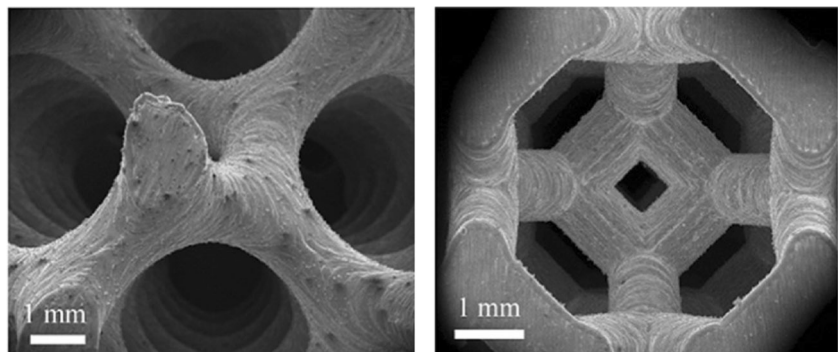
## 5 Quality inspection of lattice structures

This section will cover the methods used to inspect the quality of lattice structures. This will be followed by a discussion on the limitations of these methods and additional considerations to enable more reliable results.

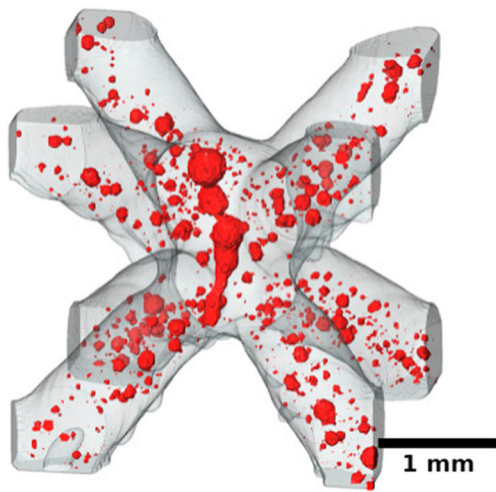
### 5.1 Measurement methods

Table 4 summarises the methods that have been used in the studies discussed in this review to inspect defects in lattice structures. Only quantitative methods will be discussed in

**Fig. 6** Left to right, SEM micrograph of two unit cells, skeletal gyroid (surface-based) and Kelvin structure (strut-based). A more pronounced stair-stepping effect can be seen in the Kelvin structure [70]







**Fig. 7** XCT results from [95] showing increased porosity in the nodes of a lattice

the following section. For additional measurement techniques, a thorough summary is provided by Banhart et al. [2].

### 5.1.1 Dimensional inaccuracies

X-ray computed tomography (XCT) data can be aligned to the original CAD model for comparison against the nominal values. This alignment can be used to calculate the maximum or average deviation of the XCT data points [36, 56, 57]. Other methods using XCT include measuring the volume or surface area of the lattice structure, as demonstrated by Van Bael et al. [97, 98] and Parthasarathy et al. [58]. For local strut measurements, Qiu et al. [87] and Melancon et al. [42] calculated strut diameter as the largest circle that could be inscribed, using the two-dimensional data from individual XCT slice images. Pyka et al. [79] and Heintl et al. [82] expand this into three dimensions by using sphere-fitting algorithms to measure the diameters of struts and designed pores. Hrabe et al. [83] suggest that using the median value of

inscribed sphere diameter is a more appropriate representation of the strut diameter. Hildebrand and Rüeggsegger [99] highlight additional parameters useful for characterising lattice structures with XCT, for example the ratio of surface area to volume which could allow for a comparison of attached particles in multiple samples.

Dimensional measurements have been performed using XCT [98, 100], SEM [11, 13, 68, 69, 73, 85–87, 101–103] and optical microscopy [25, 36, 63, 75, 76, 104–106]. It is also relatively common to use Vernier callipers for the measurement of outer dimensions (for example, lattice diameter, length, width) [74, 103, 104, 107]. The lattice’s mass has been used to calculate volume fraction, as performed by Al-Ketan et al. [70] and Wieding et al. [108].

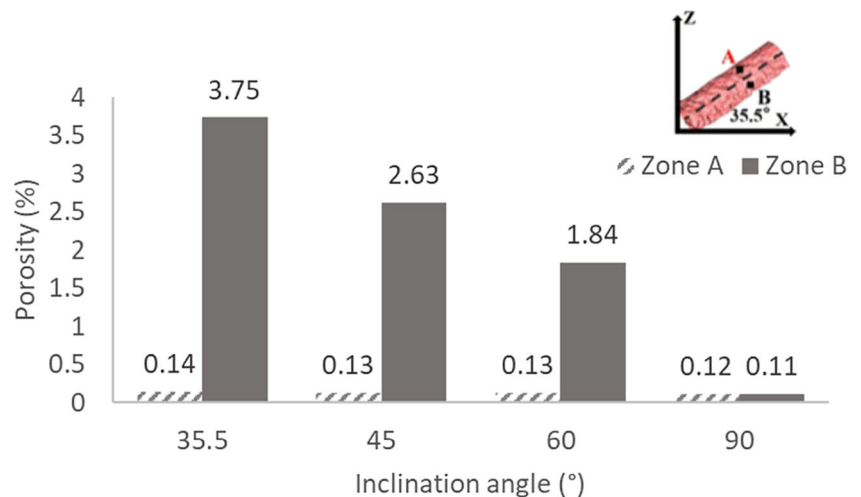
### 5.1.2 Surface texture

*Ra* values from profiles of strut surfaces have been calculated using XCT, by Kerckhofs et al. [77], Pyka et al. [79], and de Formanoir et al. [109], and using optical microscopy by Leary et al. [72]. Similarly, *Ra* was determined using the variation in strut cross-sectional area using XCT by Tancogne-Dejean et al. [71].

### 5.1.3 Porosity

Archimedes’ principle is often used to calculate porosity by comparing the mass of a lattice in two different fluids [24, 57, 75, 92]. XCT has been used to calculate porosity [56, 65, 95, 96]. Amani et al. [95] employ a stitching method with XCT, whereby small portions of the lattice are imaged at higher magnification and are stitched together. This enabled smaller voxel sizes to be achieved and a greater proportion of pores were detected. Optical microscopy was used by Qiu et al. [87] and Köhnen et al. [90] to calculate an average porosity from a

**Fig. 8** Results from Dong et al. [65] showing the effects of strut inclination on porosity in AISi10Mg lattice struts. (This graph was developed from the tabulated results and is not from the original paper. Inset illustration of strut partitioning (top right) was taken directly from the paper)



**Table 4** Defect measurement methods

Defect	Method	Reference	Quantitative characterising techniques
Dimensional inaccuracies	XCT	[31, 36, 42, 43, 56–59, 62, 64, 71, 72, 79, 82, 83, 87, 95, 97, 98, 100, 110, 111]	<ul style="list-style-type: none"> <li>• Comparison with CAD. Maximum or average deviation</li> <li>• Calculate total volume</li> <li>• Measure strut diameter (largest circle/sphere inscribed)</li> <li>• Elliptical strut cross-section</li> </ul>
	SEM	[11, 13, 68, 69, 73, 85–87, 101–103]	Dimensional measurements (length, diameter etc.)
	Optical microscopy	[25, 36, 63, 75, 76, 104–106]	
	Vernier callipers	[74, 103, 104, 107]	Infer volume fraction
	Weighing	[70, 108]	
Surface texture	XCT	[31, 56, 71, 77, 79, 87, 109]	Calculate $Ra$ value from strut profile or cross-sectional area
	Optical microscopy	[58, 72, 81, 105]	$Ra$ value from strut profile
	SEM	[11, 25, 41, 57, 68, 69, 79, 80, 82–85, 106]	Qualitative analysis
Porosity	Archimedes' method	[24, 57, 75, 92]	Infer porosity from weight in two fluids
	XCT	[56, 65, 95, 96]	Calculate total porosity
	Optical microscopy	[87, 90]	Average porosity determined from select cross-sections
	SEM	[71]	Surface porosity

select number of cross-sections. SEM has also been used by Tancogne-Dejean et al. [71] to view surface porosity.

## 5.2 Discussion of measurement methods

Optical microscopy and SEM offer established, high resolution imaging methods, however, using these imaging methods to measure structures is challenging, as highlighted by Van Bael et al. [98] who found large differences between optical microscopy and XCT results for the diameter of designed pores, particularly for non-spherical designed pores. Additionally, in order to work within the smaller measurement volume, sample preparation will often require destroying the sample. Furthermore, to obtain porosity measurements, the sample is destroyed upon sectioning to view the pores and pore morphology may be modified. Pores can easily be undetected as only those exposed upon sectioning contribute towards the calculation.

One of the main strengths of XCT is its ability to image complex AM parts and assess dimensional inaccuracies; surface texture and porosity can all be assessed from the same measurement data. In this regard, XCT is superior to optical instruments which rely on line of sight and consistent reflective properties—both of which are hindered by AM surfaces through features such as high slopes and variable aspect-ratios [112]. A drawback of XCT measurements can be found in its long acquisition times and the current limited understanding of its range of influence factors (i.e. sources of variation within the measurement output [113]). Depending on the measurement task, influence factors can significantly alter results. For example, surface determination, which defines the material boundary, has been shown to cause

greater errors in edge-to-edge length measurements over sphere-centre distances [113]. These errors may cause strut diameter measurement to have less accuracy than when measuring the position of strut centres and node centres. Additionally, the alignment of XCT and CAD data is effective for analysing the overall geometry of a lattice. However, segmentation of individual features is a more informative approach for analysing local deviations. Furthermore, large-scale deviations, such as shrinkage and warping, present a challenge during alignment of significantly dissimilar geometries. For porosity measurement, employing a stitching method [95] enables detection of smaller pores through the smaller voxel size, however, errors may form at the boundaries of reconstructed volumes [113]. Lastly, an additional discussion on XCT is provided by Ho et al. [114], focusing on its suitability for the characterisation of lattices for biomedical applications.

The remaining approaches i.e. Vernier callipers, weighing and Archimedes' method all provide useful initial assessments of lattice quality but present repeatability challenges. For example, it may be unreliable to determine porosity with the Archimedes' method as its accuracy can be reduced due to surface pores and cracks which can cause infiltration of the soaking fluid [115].

Sample preparation can further effect measurement results, through procedures such as the removal of parts from the build bed. Sing et al. [74] found lattice height was reduced upon removal via electrical discharge machining. Data processing also has a strong influence over measurement results, for example, the selection of surface texture parameters. The parameter  $Ra$  is often used to describe surface texture, and has been useful for assessing lattice surfaces, however, it may inadequately capture complex

information on AM surfaces [116]. There are investigations into employing areal parameters, which aim to more adequately capture AM surface information, such as  $Sa$ , the areal counterpart of  $Ra$  [117]. Regarding instrumentation, XCT has been used to extract areal parameters from AM surfaces, however, only from planar surfaces, as in references [118–120].

## 6 Methods for minimising defects

This section details the methods used in the literature for improving the quality of lattice structures produced by PBF. The methods are categorised in chronological order, firstly design constraints, followed by process parameter optimisation and post-processing methods, thus considering all the stages at which the quality of the lattice can be influenced.

### 6.1 Design constraints

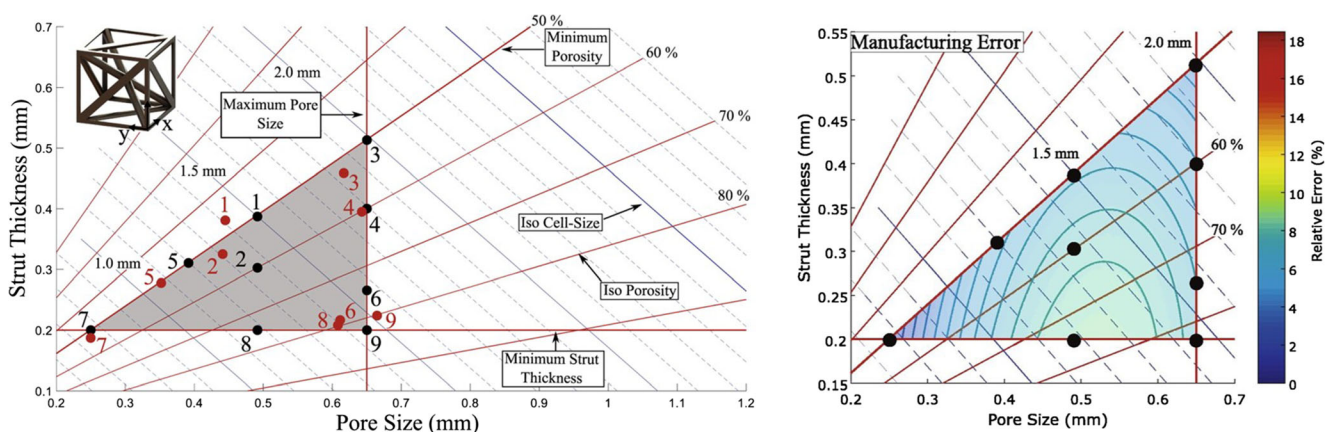
As previously discussed, the design of strut-based unit cells can be accurately controlled through the dimensions of individual members. Therefore, there is more investigation in the literature into the design constraints on strut-based unit cells as opposed to surface-based, which are mathematically defined. Several studies in the literature have investigated lattice design constraints by considering strut diameter, strut overhang angle, designed pore size and volume fraction.

Regarding strut diameter, designed pore size and volume fraction, Melancon et al. [42] provide a thorough consideration of design constraints for Ti6Al4V tetrahedron and octet-truss unit cells produced by LM. The authors define an admissible design space by selecting a range of designed pore sizes and a maximum volume fraction required for bone ingrowth, that is 50  $\mu\text{m}$  to 650  $\mu\text{m}$  and 50%, respectively. A minimum strut diameter of 0.2 mm is also defined, yielding a triangular design space as shown in Fig. 9. Arabnejad et al. [59] also use a similar design

space for tetrahedron unit cells, however, with designed pore size ranging from 50  $\mu\text{m}$  to 800  $\mu\text{m}$ . Melancon et al. assess the manufacturability of tetrahedron and octet-truss unit cells by producing several samples within this design space. For each unit cell, the strut diameter, designed pore diameter and volume fraction were each measured, calculating the relative error from nominal for strut diameter, volume fraction and designed pore size. The arithmetic mean of the relative errors was used to obtain an overall manufacturing error for each design. One of several results are shown in Fig. 9 which displays the overall manufacturing error for tetrahedral unit cells. The contour plot shows manufacturing error to increase to approximately 10% at the minimum strut diameter. Other observations included a maximum designed pore error of 18% at lowest strut diameter and a maximum error of 15% strut diameter upon decreasing volume fractions.

The minimum strut diameter constraint of 0.2 mm used by Melancon et al. is further supported by other studies. Further reduction in strut diameter is generally hindered by process limitations, particularly melt pool size. Van Bael et al. [97] found LM 0.1 mm diameter Ti6Al4V struts unachievable due to a melt pool size of approximately 0.18 mm, yielding a strut diameter of approximately twice the designed value. Strut oversizing due to the minimum melt pool size was also observed by Pyka et al. [79] and Sing et al. [104] who attempted strut diameters of 0.1 mm and 0.08 mm, respectively.

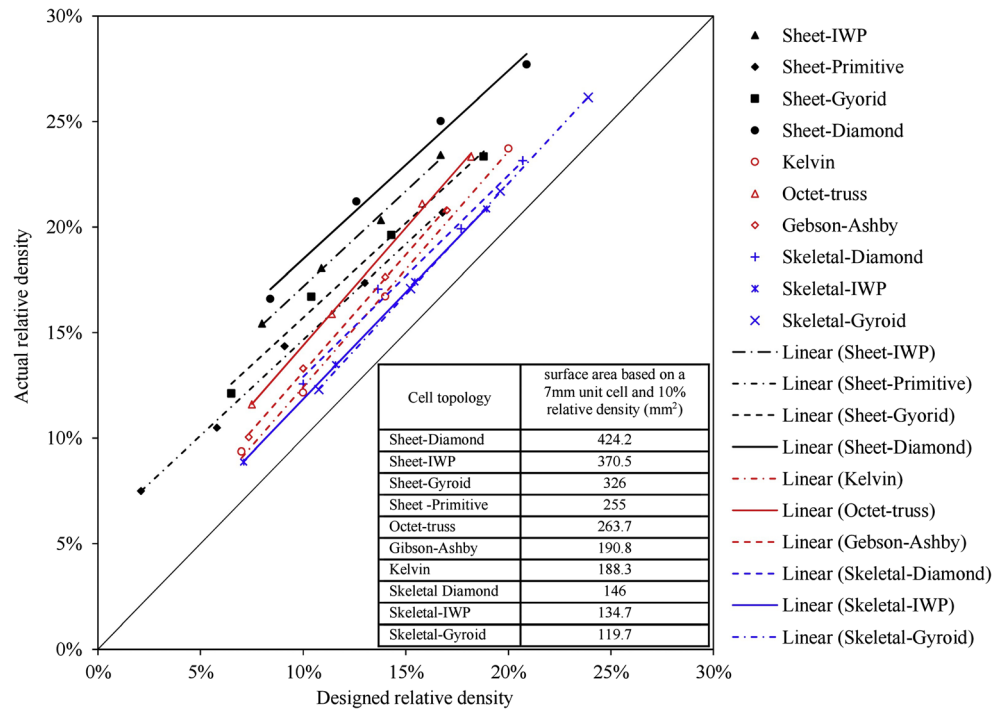
Volume fraction has also been assessed by Al-Ketan et al. [70] ('relative density', in their paper), who highlight surface area as a useful indicator for determining deviation in volume fraction for several lattice structure designs. The error between designed and manufactured relative densities increased in unit cells with higher surface area, as a greater amount of powder can remain attached to the part. Deviations of 1% to 10% in volume fraction were found in the unit cells of lowest and highest surface area, respectively—the skeletal gyroid and sheet diamond (Fig. 10).



**Fig. 9** Left, admissible design space for a tetrahedron-based unit cell. Here, 'pore size' refers to the designed pore size and 'porosity' to volume fraction. Right, the results contour map showing the overall manufacturing error for each sample [42]



**Fig. 10** Plot showing the deviation in volume fraction (here, termed 'relative density') for several unit cells. Inset table illustrates the relationship between surface area and deviation in volume fraction [70]



The overhang angle of struts (the angle between the struts and the build bed) is a key feature of lattice design, as support structures cannot be incorporated into designs. As mentioned in Sect. 4.2, horizontal struts have been successfully printed, albeit with significant deviations from their designed cylindrical shape. Yan et al. [24] comment on the importance of small overhang lengths for successfully achieving horizontal struts, as demonstrated with unit cells approximately  $3 \times 3 \times 3$  mm in size. Yan et al. [25] used the skeletal gyroid unit cell to demonstrate that manufacturing horizontal struts is more achievable when non-prismatic struts are used. The struts in the skeletal gyroid unit cell exhibit a continuous change in inclination angle, allowing for horizontal struts to be successfully printed. These observations from Yan et al. may also explain the previously discussed observations by Al-Ketan [70] in Sect. 4.2, regarding a less pronounced stair stepping effect in surface-based unit cells in comparison with strut based. Similarly, Li et al. [69] manufactured horizontal struts by linearly increasing strut diameter.

Regarding prismatic struts, Calignano defined the minimum angle for overhanging struts to be  $30^\circ$  from the build bed, for LM of AlSi10Mg and Ti6Al4V [76], which prevents aggravation of the stair stepping effect. Mazur et al. [121, 122] found a minimum strut angle and diameter of  $20^\circ$  and 0.3 mm respectively for LM of Ti6Al4V and H13 tool steel struts. Wang et al. [81] provides a comprehensive study on the manufacturability of the octahedral unit cell using 316 L stainless steel. A 'critical fabrication zone' of  $35^\circ$  to  $40^\circ$  was found for the strut overhang angle. Angles above this range were

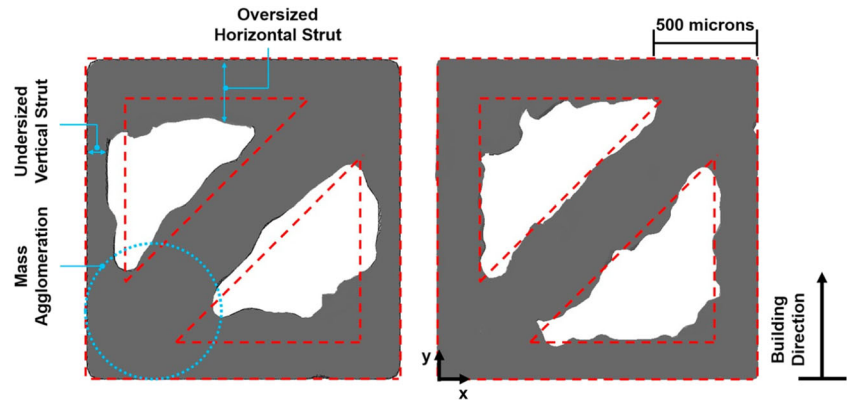
regarded as safe to print and those below were prone to failure. For EBM of Ti6Al4V, Cansizoglu et al. [123] states a minimum strut diameter of 0.7 mm and overhang angle of  $20^\circ$ .

There have been several studies optimising lattice designs by considering the inherent defects which form. Simple approaches to design optimisation have proven effective, for example Bagheri et al. [124] altered strut diameters to compensate for orientation-dependent over- or under-sizing (Fig. 11). The results included a reduction in strut diameter deviation from 60% to 3% and additional compression tests on compensated lattices showed better agreement with simulations. Li et al. [107] increased the diameter of designed pores to accommodate for later shrinkage due to excess consolidation of material. Similarly, irregular, material accumulation was combated by Li et al. [69], by increasing strut diameters near the nodes.

## 6.2 Processing parameters

Recalling Eq. (2) from Sect. 3, the process parameters governing energy density are often investigated to optimise part quality. Laser power has been investigated in several studies. Sing et al. [104] compared the effects of laser power, layer thickness and scanning speed on strut diameter and concluded that struts were most sensitive to laser power. An increase of laser power from 120 W to 240 W caused strut diameters to increase more than twofold. Qiu et al. [87] found strut diameters to increase by over 60% upon increasing laser power from 150 W to 400 W, with the amount of attached particles

**Fig. 11** XCT image of uncompensated unit cell (left) and the compensated unit cell (right). Red-dashed lines show the initial unit cell design [124]



also increasing (Fig. 12). Shen et al. [11] also observed, over a smaller range (80 W to 120 W) increasing diameter at higher laser powers. Using SEM, Hrabec et al. [83] observed smoother surfaces at higher energy input. Additional tests by Qiu et al. [87], across the same power range (150 W to 400 W), saw a peak porosity content of 2% at a laser power of 200 W.

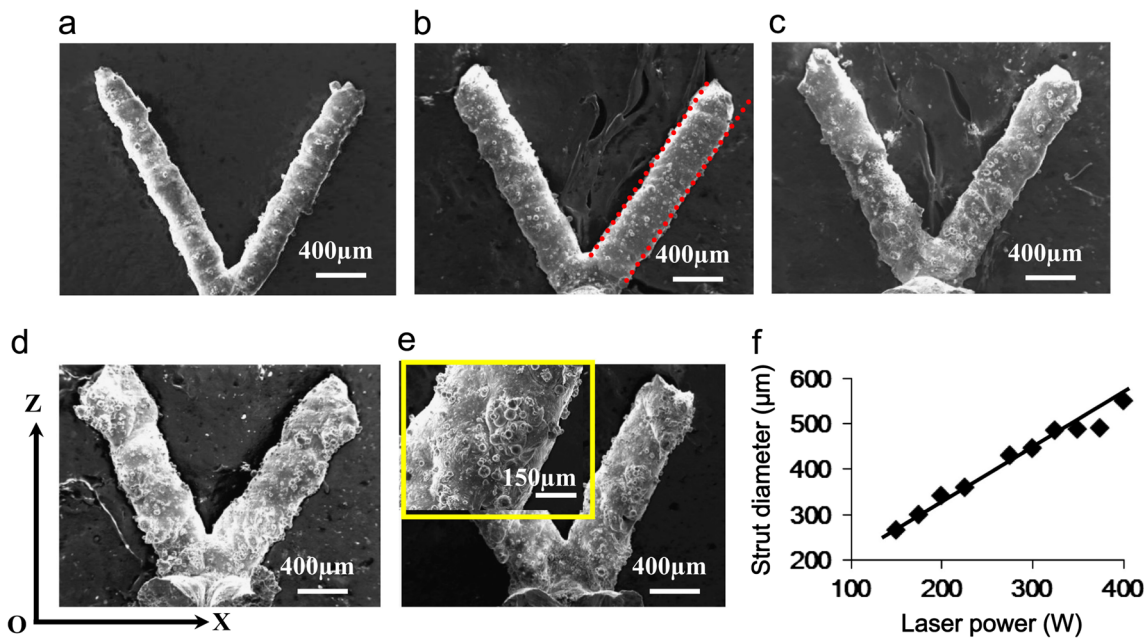
Scan speeds have also been investigated. The length-to-width ratio of the melt pool is greatly controlled by scanning speed, where at higher speeds the ratio increases and in severe cases the melt pool can become unstable and separate [54]. Qiu et al. [87] observed that increasing the scanning speed from 3000 mm/s to 7000 mm/s caused an initial decrease in strut diameter, followed by no effect at speeds above 4000 mm/s. Sing et al. [74] also found increasing scanning speed to cause a reduction in strut diameter. Han et al. highlighted effects of scan strategy on strut size, citing reference [101], explaining that continuous scan paths that make a u-turn at the ends of profiles will cause strut oversizing [36].

### 6.3 Post-processing

Post-processing techniques have proven effective for reducing the remaining defects. This section discusses post-processing methods for improving surface texture and porosity.

To reduce the surface texture of AM parts, mechanical surface treatment methods, such as sandblasting, are commonly used [57, 73], however, limited access to the internal features in lattice structures may prove difficult to treat, as highlighted by Hollander et al. [73]. Xuanyong et al. [125] provide a thorough review of procedures for surface modification of titanium and its alloys, focusing on chemical and physical methods, such as chemical etching and thermal spraying, respectively.

Chemical etching has been used in several studies to investigate its effect on Ti6Al4V lattice structures [77, 79, 109, 126]. Taking an illustrative example, de Formanoir [109] investigated the effect of hydrofluoric acid and nitric acid solutions on the



**Fig. 12** An example of the effects of laser power on strut diameter, provided by [87]

surface texture of Ti6Al4V octet lattice structures. A clear decrease in attached particles was observed, as shown in Fig. 13, with supporting  $R_a$  and  $R_t$  values, determined from XCT data. The etching process also caused the volume fraction to decrease by approximately 7%. Compression tests revealed a reduction in stiffness upon etching, with this reduction being proportional to the change in volume fraction. Chemical etching was also observed to reduce the discrepancies between experimentally determined and simulated stiffness values. This better agreement with simulation data was attributed to the removal of attached particles which allowed for a better representation of ideal, cylindrical struts.

To reduce the porosity in AM parts, hot isostatic pressing is commonly used, where porous parts are subjected to high pressure and temperature [127–129]. Wu et al. [130] observed a significant improvement in fatigue properties upon hot isostatic pressing of Ti6Al4V lattice structures, quantified by an increased endurance ratio of approximately 80%. The improved fatigue properties were attributed to a phase change from brittle  $\alpha'$ -martensite to tough  $\alpha + \beta$  mixed phases causing an increase in toughness—the key material property for resisting crack propagation. A 20% and 30% reduction in hardness and yield strength respectively was also recorded. Hot isostatic pressing was also performed by Dutton et al. [131] on EBM and LM parts. It was observed that for LM parts, the high pressure caused the gas filled pores to compress into sharp edges. However, EBM parts were unaffected due to the evacuated chamber preventing gas filled pores from forming.

Van Hooreweder et al. [132] compared the effects of stress relieving, hot isostatic pressing and chemical etching on the fatigue properties of Ti6Al4V lattice structures. The results showed chemical etching as significantly superior, due to the removal of attached powder yielding smoother surfaces.

## 7 Summary and conclusions

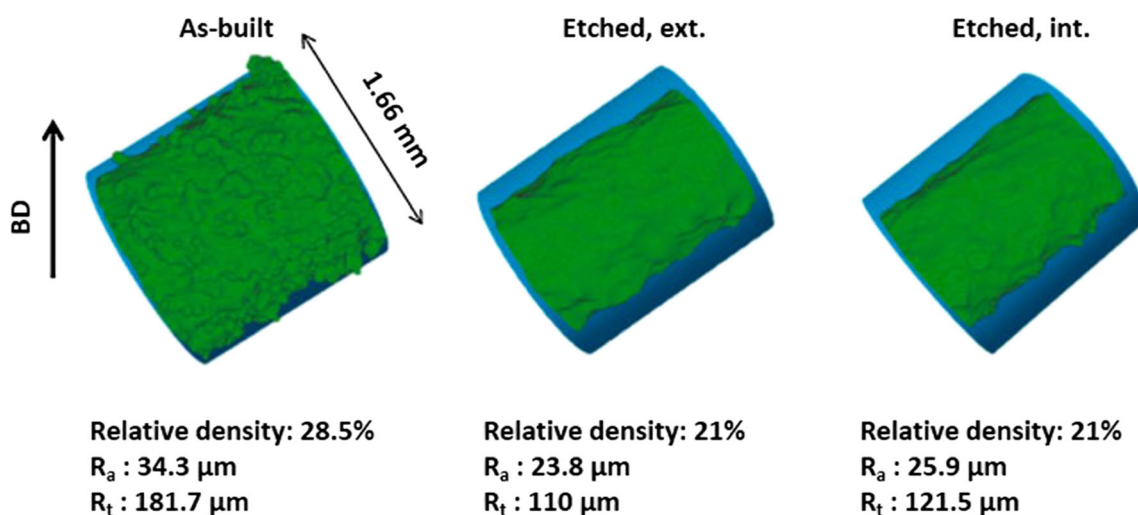
This review aims to serve as a resource on manufacturing defects in lattice structures produced by PBF. A review of defects in lattice structures manufactured by powder bed fusion processes has been presented, discussing how dimensional inaccuracies, surface defects and porosity have been observed in different designs. Additional topics of general defect formation mechanisms in PBF processes, quality inspection and defect minimisation methods have been included as supporting sections. To summarise, the questions from Sect. 1.1 are readdressed.

1. How do defects form in powder bed fusion processes?

There are numerous studies investigating defect formation mechanisms in PBF processes. It is often documented that melt pool instabilities and the heating/cooling mechanisms give rise to pores, dimensional inaccuracies and surface defects, where pores are often formed through entrapped gases and unmelted powders, dimensional inaccuracies through residual stresses and surface defects through melt pool instabilities.

2. What type of defects form in lattice structures?

Defects in strut-based lattices were found to be considerably more documented in the literature in comparison with those found in surface-based lattices. Dimensional inaccuracies are often documented for strut-based designs, with defects such as varying diameter, strut waviness, material accumulation and severe orientation dependency in the struts.



**Fig. 13** Example of the effects of chemical etching. XCT reconstruction of Ti6Al4V struts before and after chemical etching, with accompanying volume fraction (here, termed 'relative density'),  $R_a$  and  $R_t$  values. BD,

build direction; etched, ext./int, chemically etched struts in exterior/interior locations within the lattice [109]



Surface defects have been observed to be highly dependent on orientation, as is the case in general for PBF parts, with down-skin surfaces yielding rougher surfaces, supported by extracted surface texture parameters. The stair stepping effect was seen to be less severe in surface-based designs, due to the continuously varying inclination angles on the surfaces.

Porosity has been particularly documented in several strut-based designs, where larger pores have been observed in nodal areas. Strut orientation has also been documented to influence porosity, where lower inclination angles yield higher porosity, with the pore distribution grouping towards the lower region of the struts.

### 3. How can defects be appropriately measured in lattice structures?

From the literature in this review, XCT is often used to inspect the quality of lattice structures, which enables dimensional inaccuracies, surface texture and porosity to be assessed from the same data set. Limitations include long acquisition times and a limited understanding of the effect on measurement uncertainty of various influence factors. It has been shown that XCT voxel size limits the detectable porosity—stitching methods can improve this but introduce new errors when combining multiple scanning data sets. Additional established methods such as optical microscopy and SEM remain effective for inspecting defects within significantly smaller measurement volumes. Inexpensive techniques and apparatus such as Archimedes' method, and Vernier callipers remain as useful tools for porosity and dimensional measurements, respectively.

### 4. How can defect formation be minimised?

There are several methods employed to minimise defect formation in lattice structures; this has been attempted through development of design maps, design constraints, process parameters and post-processing. Dimensional inaccuracies can also be significantly reduced by accounting for orientation dependent errors measured from test samples.

Energy density calculations continue as an appropriate method for assessing primary process parameters. Expected results have been observed: increased laser/electron beam power increases strut size, increase scanning strategy reduces strut size. Scanning profiles can further influence strut size, particularly at the ends of scanning paths.

Post-processing methods have been documented as particularly useful for reducing surface irregularities, however, limited access to the lattice geometry inhibits the performance of mechanical processes such as sandblasting. Chemical etching, which circumvents problems due to limited access, provides no control over specific regions of interest and alters mechanical properties of the lattice, for example causing a reduction

in stiffness. Chemical etching has been observed as significantly superior for improving fatigue properties than stress relieving and hot isostatic pressing.

Looking forward, further research opportunities include more in-depth characterisation of manufacturing defects in surface-based lattice structures, as the literature found for the review was heavily skewed towards strut based. Continued development of design maps for individual unit cells will help increase understanding of their manufacturability. Additionally, incorporating defects into existing lattice structure models will be useful for quantifying the influence of defects on the desired function of the lattice [133]. From a metrology perspective, process monitoring methods will aid in increasing understanding of defect formation, particularly regarding the influence of process parameters. Furthermore, the popularity of XCT must be met with establishing traceability, providing confidence to the measurement of defects. Overall, this content, and subsequent work in the field, will contribute towards a better understanding of defects in PBF lattice structures, accelerating the application of functional lattice structures into various industries.

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