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**MEASUREMENT OF FRACTURE TOUGHNESS OF METALLIC MATERIALS**  
**PRODUCED BY ADDITIVE MANUFACTURING**

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## ABSTRACT

This study focuses on the microstructure and mechanical properties of metallic materials produced by additive layer manufacturing (ALM), especially the Laser Beam Melting process. The influence of the specimen orientation during the ALM process and that of two post-build thermal treatments were investigated. The identified metal powder is Ti-6Al-4V (titanium base). Metallographic analysis show their effects on the microstructure of the metals. Mechanical experiments involving tensile tests as well as toughness tests were performed according to ASTM (American Society for Testing and Materials) norms. The results show that the main influence is that of the thermal treatments; however the manufacturing stacking direction may lead to some anisotropy in the mechanical properties.

**KEYWORDS:** *Additive Layer Manufacturing, Laser Beam Melting, Ti-6Al-4V, fracture toughness, mechanical properties*

## NOMENCLATURE

K <sub>IC</sub> – Mode I fracture toughness	ASTM - American Society for Testing and Materials
E - Young Modulus	EBM - Electron Beam Melting
YS - Yield Stress	HIP - Hot Isostatic Pressing
UTS - Ultimate Tensile Stress	LBM - Laser Beam Melting
A - Plastic Elongation	SEM - Scanning Electron Microscope
ALM - Additive Layer Manufacturing	SLM - Selective Laser Melting
AM - Additive Manufacturing	

## 1. INTRODUCTION

For a few decades, additive manufacturing (AM) appeared as on the way of revolutionizing both the design possibilities and the manufacturing standards. It is now used as a regular technique in space, aircraft and dental industry. Design limits were incredibly stretched thank to unprecedented freedom and part complexity. Moreover AM totally fits the use of CAD tools, thus matching topology (and

weight) optimization approach well. Global manufacturing time and cost were deeply changed due to the decrease in the number of processing operations and number of parts. Additionally, AM leads to an optimal use of raw material.

This paper focuses on the Laser Beam Melting AM technique dedicated to Ti-6Al-4V. AM also means a different manufacturing process resulting in a different microstructure due to rapid solidification rate. Consequently the mechanical properties have to be precisely determined with respect to lots of parameters available (powder properties, machine properties, batch design, post-processing, loading orientation, etc.). Much work has been done to investigate the process-microstructure-tensile property relations for LBM and shed light on the influence of the thermal treatment [1-3] or the building orientation [4-6]. Similar studies were done for EBM [7-10]. Carroll *et al.* [11] gave a summary of the tensile properties available in literature for Powder Bed Fusion, Direct Energy Deposition, and Electron Beam Melting techniques. Several papers dealt with the fatigue properties [12-23]. Li *et al.* [24] compared fatigue properties published in open literature for LBM and EBM. However to the author knowledge only few papers focused on the fracture toughness of AM Ti-6Al-4V [12, 17-19, 23, 25], although this is a key point in the damage tolerance approach. Table 1 gives an overview of the data available in open literature for Ti-6Al-4V toughness obtained with Laser Beam Melting (LBM) and Electron Beam Melting (EBM) techniques. There is huge scatter between the various results for a given technique; however the dependency on the heat treatment and on the layering direction was reported by most of the authors.

The present contribution focuses on the microstructure-tensile-toughness relations of Ti-6Al-4V with respect to sample orientation and heat treatment used. LBM technique was used to manufacture several specimens, including metallography cube, tensile and fracture toughness specimen. The various specimens were produced in the same batch to ensure unique manufacturing conditions. Each batch received, or not, a particular heat treatment. In the end, samples had to experience several stages: ALM process, thermal treatments, machining, fatigue pre-cracking, final mechanical tests. The statistical analysis required a fine tracking of each specimen to be sure the conclusions could be drawn concerning the influence of the parameters of interest.

The paper is organized as follows: after introducing the samples preparation and the tests set-up, results of metallographic analysis, tensile tests, and toughness tests are presented, followed by a discussion.

## **2. SAMPLES PREPARATION AND TESTING METHODS**

### **2.1 Materials, manufacturing parameters and thermal treatment**

Two batches of Ti-6Al-4V specimens were manufactured. Each batch was manufactured during a single job to ensure that the same manufacturing parameters were used for all the samples. For the same reasons the manufacturing parameters were also kept the same for the two batches, and the jobs were done in a row to avoid too much recycling of the powder between the two batches.

Batch #1 and Batch #2 were manufactured from Ti-6Al-4V powder from Advanced Powders and Coatings, with 15-45  $\mu\text{m}$  particle size. The powder had been recycled 14 times for batch #1 and 15 times for batch #2. The batches were manufactured in a EOS M280 machine using a 1060-1100  $\mu\text{m}$  wavelength laser. Layers were 60  $\mu\text{m}$  thick. Both batches followed a stress relieving treatment on the base platform (670°C / 5h) to release residual stresses. The as-obtained Batch #2 was submitted to an additional Hot Isostatic Pressing (HIP) process (920°C / 2h / 1020 bars).

Both batches were then ready to be machined to obtain the expected shape and surface quality ( $R_a = 0.8$ ), allowing mechanical testing.

**Table 1: Results of LBM and EBM Ti-6Al-4V toughness reported in literature**

AM technique	Reference	Scanning strategy	Post processing	Sample orientation / load orientation	Surface finishing	$K_{IC}/K_Q$ (*)	
<b>LBM</b>							
	Greitemeier 2016 [23]	Rotation by 30° for the next layer	Annealing (710°C - 2h) followed by cooling under argon atmosphere	Vertical / Z	Milled	<b>49.5</b> (±1.5)	
			HIP (920°C - 1000 bars - 2h under argon)			93.5 (±0.5)	
	Edwards 2015 [18] (**)	Rotation by 67° for the next layer	As-built		XY / Y	Machined	66.9(**) (2.6)
					XZ / Z		64.8(**) (16.9)
					YZ / Z		41.8(**) (1.7)
	Cain 2015 [17]	Bi-directional: scan of the perimeter, then zigzag formation and rotation by 90° for the next layer	As-built		XY / X	Sliced from continuous block then milled	28 (±2)
					XZ / X		23 (±1)
					ZX / Z		16 (±1)
			Stress relief (650°C - 4h under argon)	Milled	28 (±2) 30 (±1) 31 (±2)		
			Annealing (890°C - 2h under argon)		XY / X XZ / X	Milled	41 (±2) 49 (±2)

				ZX / Z		49 ( $\pm 1$ )
	Van Hooreweder 2012 [12]	Bi-directional: scan of the perimeter, then zigzag formation and rotation by 90° for the next layer	As-built	XY	Machined by EDM	52.4 ( $\pm 3.48$ )
<b>EBM</b>						
	Greitemeier 2016 [23]	Alternates between 0° and 90°	Annealing (710°C - 2h under vacuum) followed by cooling under argon atmosphere	Vertical / Z	Milled	106.5 ( $\pm 3.5$ )
				Vertical / Z	Milled	119
	Edwards 2013 [25]	Rotates 90° after every layer	As-built	Horizontal Vertical / Z	Machined	110 (8.9) 102 (7.4)
	Seifi 2015 [19] (***)	?	As-built	XY / X XY / X XZ / X XY / Y XZ / Z	?	68 80 76 67 65

(\*) Toughness values that succeed to meet the ASTM E399 validity requirement are denoted by bold underline font.

(\*\*) Edwards *et al.* [18] used plain stress specimens instead of plain strain ones; this is likely to have given higher toughness values.

(\*\*\*) Seifi *et al.* [19] used bending toughness specimen whereas most authors used CT specimens.

## 2.2 Tensile and toughness specimens and specimen orientations

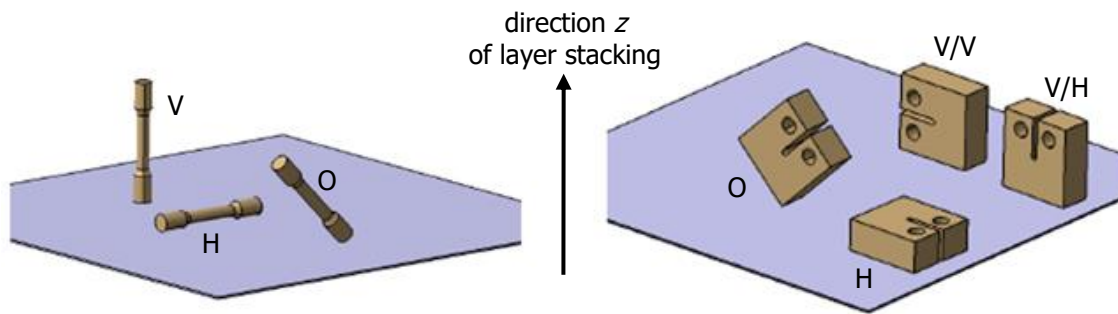
The aim of the project required to get all the samples of a given batch in one job. For the sake of a statistical analysis and due to the dimensions of the manufacturing zone, dimensions of the samples had to be sufficiently small. For tensile tests, standard round specimens were designed using the following dimensions specified by ASTM E8 norm [26]: diameter for the section: 6 mm, length of the reduced section: 27 mm, total length: 60 mm. Seven samples were manufactured for each orientation/material/thermal treatment. After the ALM process and the thermal treatments, specimens had to be machined (mainly by turning process) to achieve good dimensional quality and to avoid any dependency on the quality of the surface shape.

For toughness tests, compact (CT) specimens were designed according to ASTM E399 norm with 32 mm specimen width (W32-B16). This allowed to manufacture five samples for each combination of material/orientation/thermal treatment. After the ALM process and the thermal treatments, specimens were machined to get the corresponding dimensions and to create the notch. A fatigue crack was created and propagated according to ASTM E399 norm recommendations. Unfortunately, manufacturing difficulties appeared during Ti-6Al-4V process and lead to several cracks in the samples during or after the manufacturing stage. Consequently such samples were missing for the toughness tests, as will be detailed later. For each group of samples, the number of tested specimens will be given in the corresponding tables.

The ALM process consists in melting the part layer by layer, favouring the anisotropy of direction  $z$  of the "stacking". Consequently, three particular orientations of the samples were identified for the tensile specimens, as shown in Figure 1. Note that for tensile specimens, this direction corresponds to the loading axis: vertical (denoted by "V"), horizontal ("H"), slanted at  $45^\circ$  ("O"). Note that in-plane directions  $x$  and  $y$  were not distinguished since the laser advance direction is turning of  $30^\circ$  when changing for the next layer. For the toughness specimens however, one needs to report both the building plane and the loading axis due to multiple crack orientations. Consequently four different orientations were identified: horizontal (resp. oblique) specimens gave only horizontal (resp. oblique) loading direction denoted by "H" (resp. "O"), whereas vertical specimens could lead to both a vertical



notch and horizontal loading direction (denoted by "V/H"), and a horizontal notch and vertical direction ("V/V").



**Figure 1: Orientations of the samples for tensile and toughness tests**

### 2.3 Metallographic analysis

Preparation of the samples consisted in polishing, under water as lubricant, using a PRESI device (reference: MECAPOL P 220 U). Sequential grinding with silicon carbide (SiC) abrasive paper was performed for the preliminary steps, using water as lubricant. Four grades were used (P320, P800, P1200 and P4000), the corresponding abrasive grains median diameter of which are 46.2  $\mu\text{m}$ , 21.8  $\mu\text{m}$ , 15.3  $\mu\text{m}$  and 2.5  $\mu\text{m}$  respectively.

A specific Silica suspension (particle size: 0.04  $\mu\text{m}$ , PRESI reference: SPM) was used for the finishing. No additional lubricant was necessary in this case. The best results were obtained by controlling the parameters (rotation speed  $\nu$ , load  $F$  and duration  $t$ ) as follows:

- for P320:  $\nu=300$  rpm,  $F=25$  N,  $t=1$  min;
- for P800:  $\nu=300$  rpm,  $F=25$  N,  $t=0.5$  min;
- for P1200:  $\nu=300$  rpm,  $F=25$  N,  $t=0.5$  min;
- for P4000:  $\nu=300$  rpm,  $F=25$  N,  $t=0.5$  min;
- for silica suspension,  $\nu=150$  rpm,  $F=15$  N,  $t=2$  to 4 min.

To reveal the microstructure of the specimen, *Kroll's* reagent (6 mL of  $\text{HNO}_3$ , 3 mL of HF and 91 mL of  $\text{H}_2\text{O}$ ) was used: 2 min 30 sec at room temperature.

Optical observations were performed with a *ZEISS Stemi 2000* binocular (for magnifications ranging between x0.65 to x5) and a *ZEISS Axiotech* microscope (for magnifications ranging between x5 to

x100). In both cases, a *PRESVIEW* software allowed acquiring images. For larger magnification (up to x10 000), a *HITACHI TM1000* scanning electron microscope (SEM) was available.

One must note that observations were performed along two orientations:

- perpendicularly to the plane of manufacturing. The corresponding observation (horizontal cross section) can then reveal the microstructure of a layer;
- perpendicularly to the layers of the material. The corresponding observation (vertical cross section) reveals the homogeneity along the height of the sample, from the first to the last fused layer.

For practical reasons, we will call these two orientations as horizontal and vertical respectively in the following.

## **2.4 Tensile and toughness tests set-up**

Tensile tests were performed on a Instron electromechanical testing machine of type UTM-EM. Machine driving was performed by Bluehill 2.21 software. A 2525-801 type 100 kN Instron load cell was used to record the tensile force. A 12.5mm 2620-601 Instron clip-on extensometer was used to measure the axial strain of the specimen. Tests were performed according to recommendation of norms ASTM E8 [26] and NF EN 10002-1 [27]. Load rate of 2 MPa/s was used till reaching 70 MPa. Then strain rate of  $0.00025 \text{ s}^{-1}$  was prescribed. At about 2% strain, the clip-on extensometer was removed manually, and strains were measured directly by the displacement of the load frame.

Toughness tests were performed on the same Instron testing machine using Bluehill software 2.21. Apparatus involved the same 100 kN load cell and a 2670-130 Crack Opening Displacement Gauge from Instron. Tests were performed following the ASTM E399 recommendation [28]. Load rate of 0.33 kN/s was used up to global failure of the specimen.

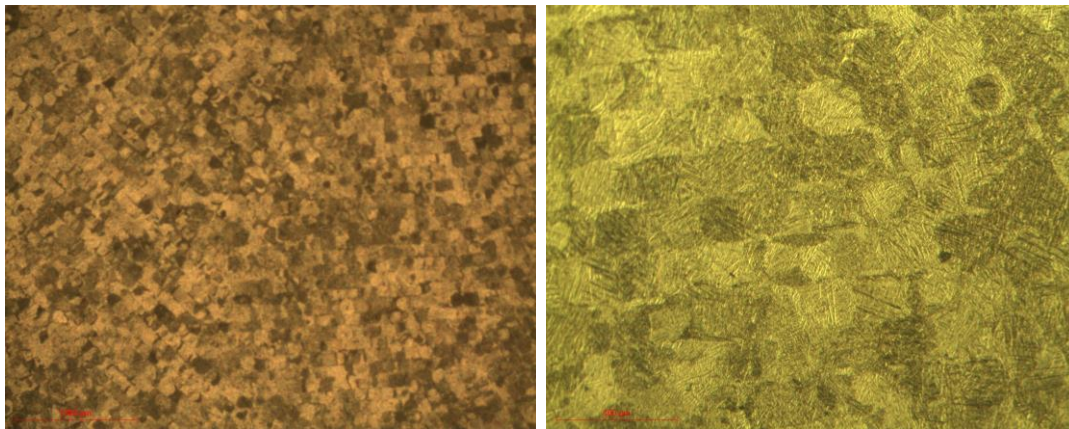
## **3. RESULTS FOR TI-6AL-4V**

### **3.1 Results of metallographic analysis for Ti-6Al-4V alloy**

#### **Batch #1: Stress-relieved Ti-6Al-4V**

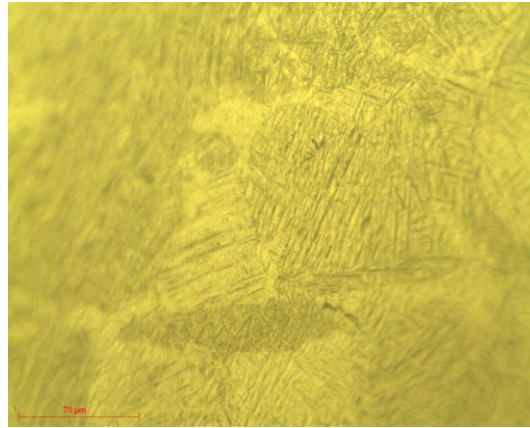
Observations at low magnification revealed a good microstructural homogeneity of the sample, whatever the orientation. This result could prove that the cooling rate was similar during the manufacturing, whatever the number of layers. The mass of obtained sample has no (or not optically detectable) influence on the resulting microstructure.

**Horizontal:** At low magnification, each layer is composed of lines parallel one to each other, each of them showing a succession of more or less regular squares (see Figure 2). This typical microstructure reflects the successive scanning orientations of the laser beam during the manufacturing. Indeed, for each layer, the laser beam produced a melted bead and, after recrystallization, a line of solid material (the width of which is ranging between 100 and 120  $\mu\text{m}$ ) aligned to the direction of the laser beam displacement. The direction of the beam changing from layer to layer, and the induced fusion affecting not only the concerned layer but also the 2-3 lower layers, the recrystallization produced square grains detectable along each line. Observing several orientations of the lines indicates that the polishing affected several successive layers.



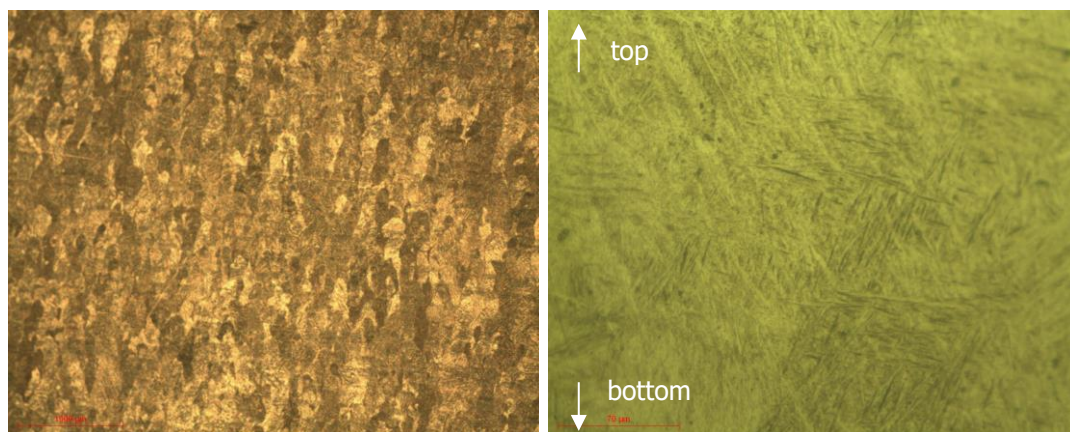
**Figure 2: Microstructure observed for the Ti-6Al-4V sample at low magnifications**

Higher magnification allows observing the martensitic structure (see Figure 3) of each square of the Ti-6Al-4V sample. This microstructure indicates that the cooling rate during the manufacturing process was fast enough to lead to the  $\alpha'$  metastable phase.



**Figure 3: Martensitic microstructure of the Ti-6Al-4V sample**

**Vertical:** Observations of the Ti-6Al-4V sample along its height revealed that, if the microstructure is composed of layers, the obtained grains after recrystallization have a columnar structure (see Figure 4), the orientation of which is perpendicular to the layers. This microstructure is only possible by fusion of the lower layers, leading to a melted bead with a “keyhole” section when scanning the laser beam [29, 30]. Figure 4 confirmed that the grains are made of  $\alpha'$  martensitic phase.

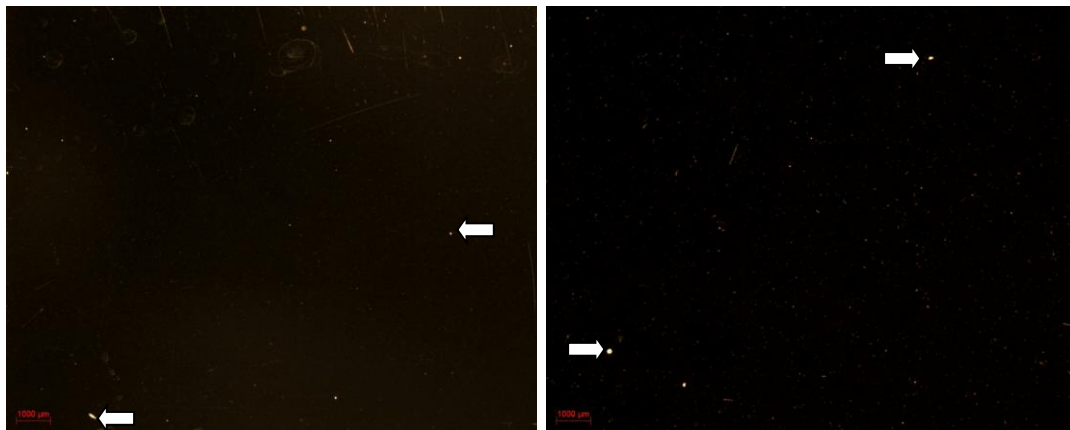


**Figure 4: Microstructure along the height of the Ti-6Al-4V sample at low (left) and high (right) magnification**

As a result, it appears that the microstructure of the Ti-6Al-4V samples obtained by LBM is quite homogeneous, even if it does not seem to be isotropic, nor in the layer plane, nor in the height plane.

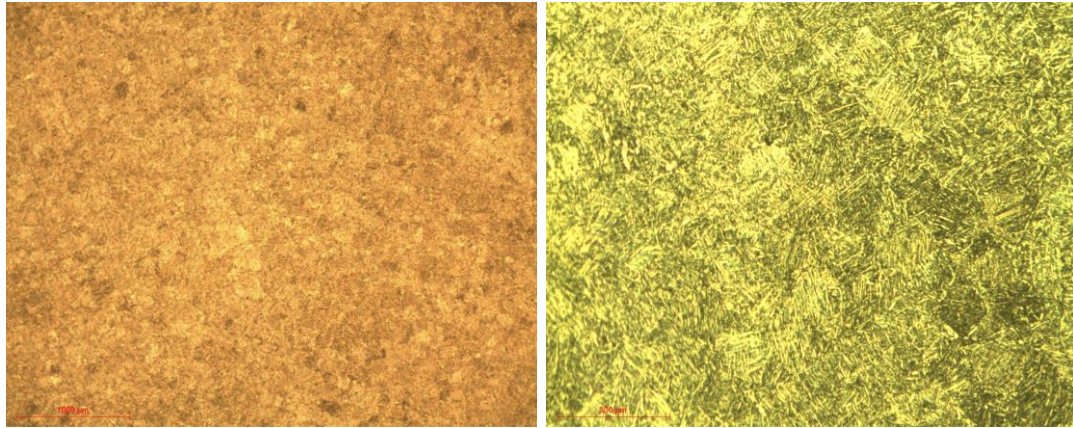
## Batch #2: Ti-6Al-4V after stress-relief and HIP

The HIP treatment was expected to eliminate, or at least reduce, the porosity rate in the obtained samples. Among all the potential techniques, we estimated the porosity rate by observing the polished surface of the sample and detect the porosity by difference of colour. The surface of the sample appears dark while porosity appears as white dots (see Figure 5). As a first result, we can say that the HIP finally did not reduce the porosity rate, maybe due to the very low proportion of porosity (lower than 0.2%) in the initial material.



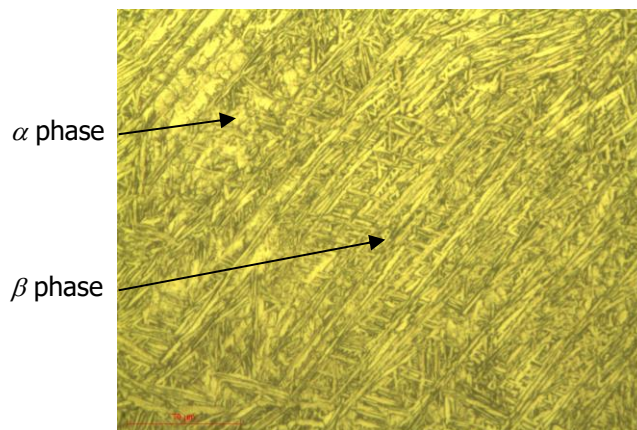
**Figure 5. Porosity (white dots) detected in the Ti-6Al-4V sample before (left) and after (right) the HIP treatment**

**Horizontal:** The HIP treatment clearly blurred the anisotropic microstructure of the Ti-6Al-4V sample (see Figure 6), even if some residual anisotropy can be detected. The lines and squares largely vanished.



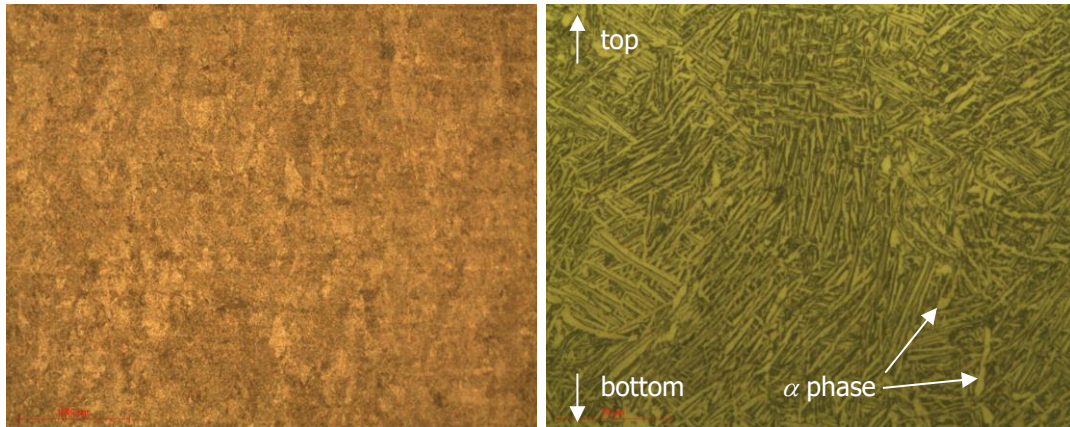
**Figure 6. Microstructure observed for the Ti-6Al-4V sample after HIP at low magnifications**

At higher magnification (see Figure 7), observations showed that the martensitic structure disappeared; the sample is now constituted by a  $\alpha/\beta$  structure. The proportion of obtained  $\alpha$  phase is low due to a thermal treatment either at too low temperature or with a too short duration.



**Figure 7. Microstructure observed for the Ti-6Al-4V sample after HIP at high magnification**

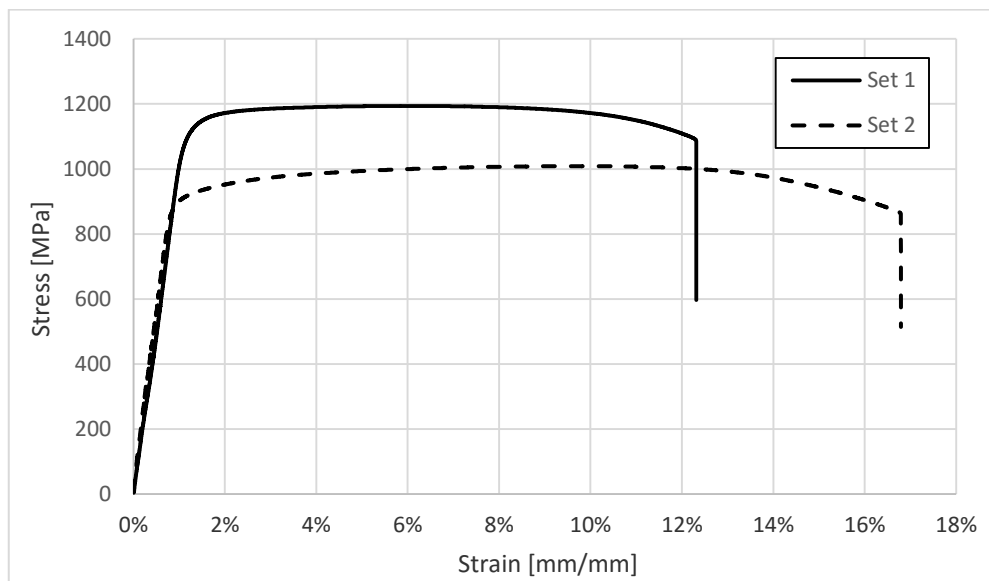
**Vertical:** One can note at low magnification (Figure 8, left) that the anisotropy is not fully blurred, as layers are still slightly visible, as well as some residual columnar grains. We could then estimate that the anisotropy is more strongly present in the vertical plane than in the horizontal one. Higher magnification (Figure 8, right) allows to conclude that the  $\beta$  phase appears along the grain boundaries between the columnar grains, as expected.



**Figure 8. Microstructure along the height of the Ti-6Al-4V sample after HIP at low (left) and high (right) magnification**

### 3.2 Results of tensile tests

Two typical stress-strain curves obtained for samples of batches #1 and #2 are shown in Figure 9. Detailed results including Young's modulus (E), tensile yield stress (YS), ultimate tensile stress (UTS) and elongation (A%) for batches #1 and #2 are reported in Table 2 and Table 3 respectively. Unlike toughness specimen, tensile specimens did not experience cracking during the ALM process thank to more slender shape, and all specimens manufactured could be tested.



**Figure 9 - Typical stress-strain results for titanium specimen**

Except for elongation, results for batch #1 were rather repeatable. For Young’s modulus, yield stress and ultimate stress, the slight differences between the various orientations are hardly significant and the material can be considered as quasi-isotropic for those properties. However a moderate 16% increase in the elongation tends to appear for the vertical direction (building direction), although low repeatability maintains doubt about this.

Samples of batch #2 were found to provide very repeatable results including the elongation. Again the material looks almost isotropic including the elongation. This tends to confirm that the HIP treatment removed the slight z-anisotropy noted in the elongation in batch #1.

Comparisons between batch #1 and batch #2 lead to the most interesting conclusions. Young’s modulus seemed to have slightly decreased (by 4.6%) after the HIP treatment, but the difference is not high enough to be sure that it is relevant. Important drops were noticed concerning the yield stress (-19.2%) and the ultimate stress (-14.7%). On the opposite elongation increased by 59%. As a conclusion, the HIP treatment did not lead to any improvement in the tensile characteristics, except the large 59% increase in the inelastic deformation of the material.

**Table 2: Results of tensile tests for batch #1**

Specimen orientation		H	O	V
Number of tested samples		7	7	7
E [GPa]	<b>Aver.</b> (std dev %)	<b>118.2</b> (7.4%)	<b>118.0</b> (3.5%)	<b>119.2</b> (2.5%)
YS [MPa]		<b>1085</b> (1.7%)	<b>1098</b> (2.0%)	<b>1116</b> (1.9%)
UTS [MPa]		<b>1166</b> (1.5%)	<b>1166</b> (1.9%)	<b>1155</b> (6.2%)
A%		<b>9.9%</b> (5.3%)	<b>9.9%</b> (18.4%)	<b>11.5%</b> (10.7%)



**Table 3 - Results of tensile tests for batch #2**

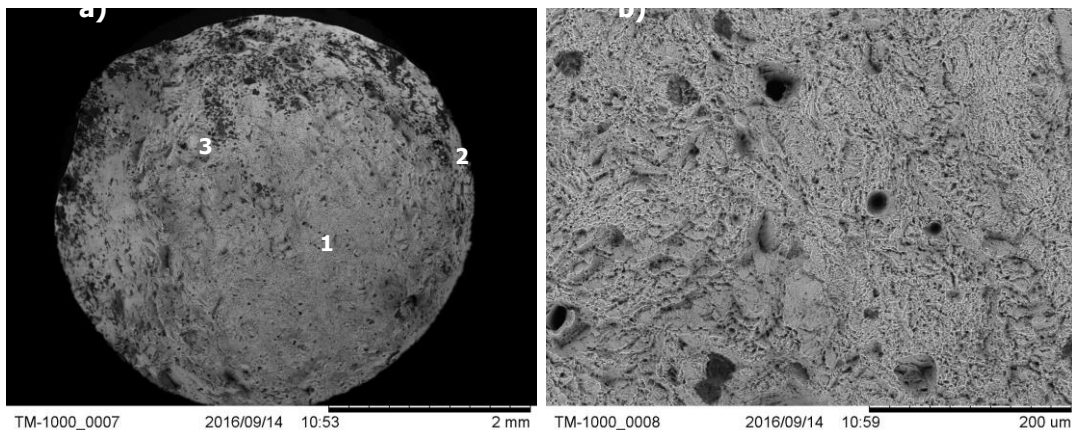
Specimen orientation		H	O	V
Number of tested samples		7	7	7
E [GPa]	<b>Aver. (std dev %)</b>	<b>114.6</b> (1.6%)	<b>111.6</b> (3.2%)	<b>112.8</b> (0.9%)
YS [MPa]		<b>896</b> (0.6%)	<b>909</b> (0.8%)	<b>861</b> (4.3%)
UTS [MPa]		<b>995</b> (0.5%)	<b>1013</b> (0.6%)	<b>967</b> (2.9%)
A%		<b>16.2%</b> (2.1%)	<b>16.6%</b> (3.6%)	<b>17.1%</b> (6.4%)

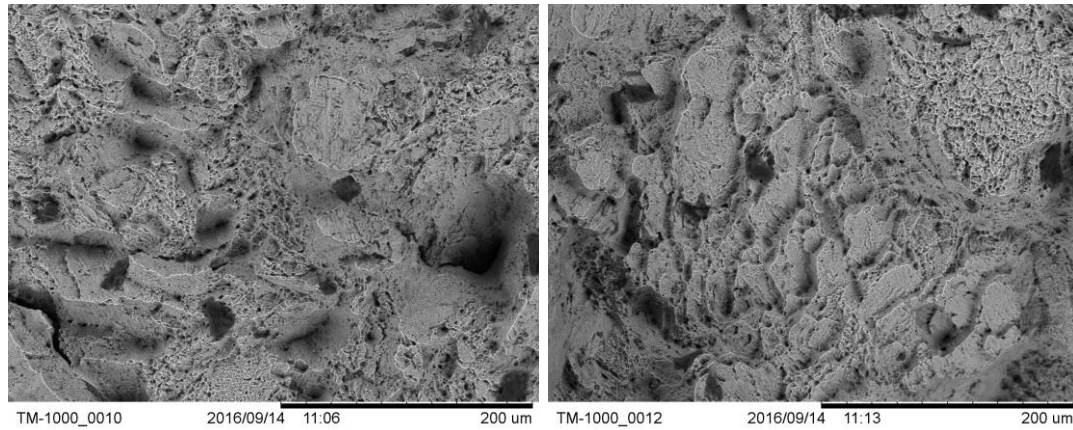
### 3.3 Fracture surface analysis of tensile tests

#### Batch #1

The overall view of the tensile fracture surface of the vertical as built samples is presented in Figure 10a. It appears that failure mechanism was mostly ductile as a transgranular fracture surface with a network of dimples can be largely observed (see Figure 10b).

Some defects can also be observed, such as porosity (resulting from microvoid coalescence during the manufacturing process), mostly located in the center part of the section of the sample (see Figure 10b), even if this porosity does not seem to have a strong influence on the mechanical properties, at least for tensile tests. One could assume that the porosity could be the main cause of the fracture [13].





**Figure 10. Fracture tensile surface of the vertical stress-relieved sample. (a): overall view of the fracture surface. (b): defects detected in location 1 in (a). (c): microcracks detected in location 2 in (a). (d): brittle fracture surface in location 3 in (a).**

Nevertheless, another defect seems to have much more influence on the mechanical behaviour of the sample: the micro-cracks. They were mostly detected in the external part of the section of the sample (see Figure 10c), even if few of them can be observed in the centre of the section. These micro-cracks were attributed in our case to the residual stress in the sample after the laser manufacturing, which was not erased after the stress relieving treatment. The large number of such micro-cracks at the fracture surface should underline their strong influence on the mechanical performance of the stress-relieved Ti-6Al-4V sample. This conclusion was confirmed by the brittle surfaces detected around most of these micro-cracks, leading to cleavage facets and a layered profile (see Figure 10c-d). These brittle surfaces were attributed to cleavage within layers of the material, joining two ductile surfaces, which were the interface between two successive layers. Nevertheless, in such a mixed ductile-brittle behaviour, the ductility was largely dominating.

Similar results were obtained by observing samples in other directions (horizontal and sidelong), confirming the quasi-isotropy of the obtained LBM samples.

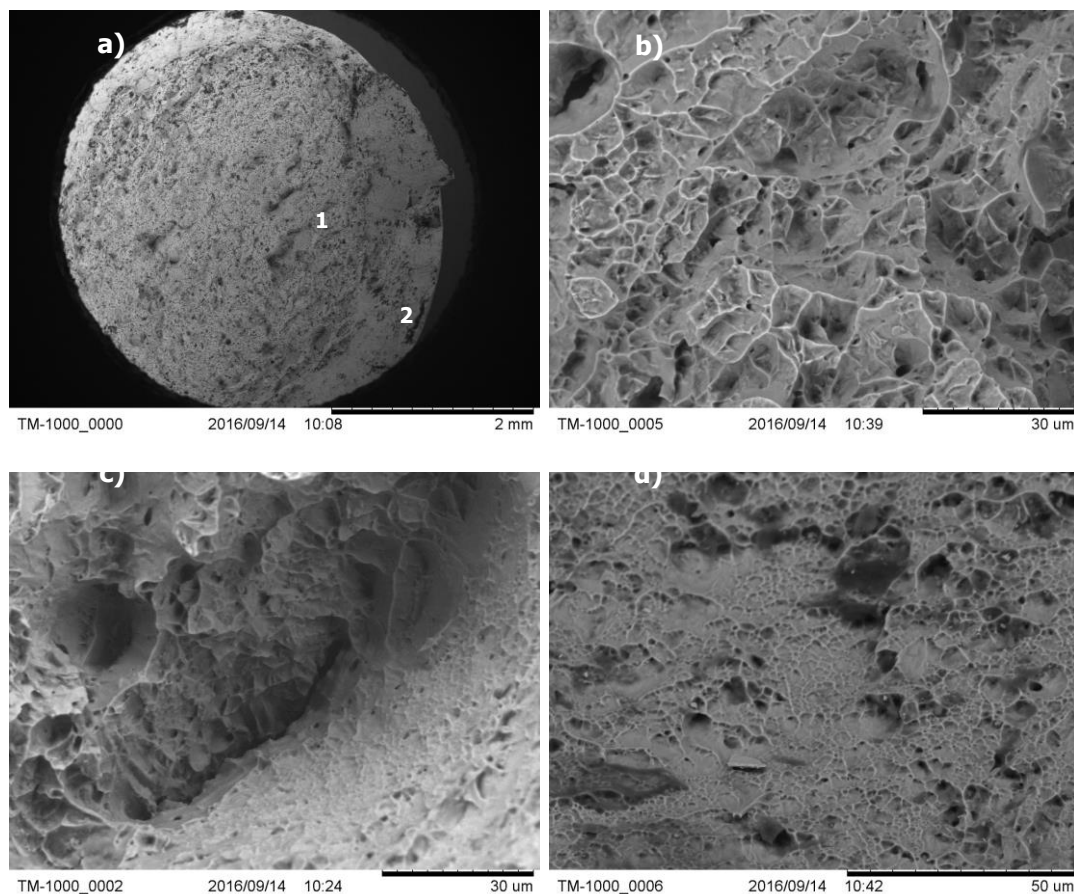
## **Batch #2**

As a first result, one must note that at the lowest magnification (x40), the whole fracture surface of the vertical sample after HIP could be observed (see Figure 11a), what was not possible for the

stress-relieved sample. This remark means that the HIP treatment lead to a higher ductility, and so to a larger necking, confirming the larger elongation measured for such samples during the tensile tests.

Then, it appeared that there was no porosity detected on the fracture surface, proving that if there was residual porosity in the whole sample (as noted in Figure 5), it had no influence on the mechanical performances of the HIPed sample. The thermal treatment seems to be very efficient to increase the plastic deformation of LBM manufactured Ti-6Al-4V.

Increasing magnification during the observation allowed revealing a pure ductile behaviour of the sample, characterized by the dimples network (see Figure 11b). Such a behaviour could be confirmed by both a larger necking of the sample and a "cup and cone" aspect (typical of ductile material).



**Figure 11. Fracture tensile surface of the vertical sample after HIP treatment. (a): overall view of the fracture surface. (b): observed ductile behaviour. (c): microcrack detected in location 1 in (a). (d): magnification in location 2 in (a).**

Some few micro-cracks were still remaining (see Figure 11c), but they did not give rise to cleavage facets attributed to brittle behaviour. The final overloaded surface (see Figure 11d) also presented a ductile aspect, but the size of the dimples is much smaller. This result can be explained as the duration of final fracture was too short to allow large plastic deformation of the corresponding area, and then to lead to larger dimples.

Similar results were obtained for other directions (horizontal and sidelong).

### **3.4 Results of toughness tests**

Results of toughness tests for Ti-6Al-4V batches #1 and #2 are reported in Table 4 and Table 5 respectively. This includes the number of tested samples, the number of pre-cracking fatigue cycles and the results of plane strain fracture toughness  $K_{IC}$ .

Unlike tensile samples, let us first remark that several CT samples experienced cracking during or immediately after the ALM process. This was due to residual stresses in such compact parts, although samples were "only" 19 mm thick (aiming at 16 mm after machining).

**Batch #1** showed very repeatable values. The number of fatigue cycles was similar, except for the "V/V" orientation that was 20% lower. Fracture toughness was almost equal for "H", "O" orientation. On the contrary, the "V/H" and "V/V" samples were found to show a 17% increase and a -4% decrease, respectively.

Results of **batch #2** tests were also found to be very repeatable. Fatigue cycling was similar for all orientations, and was 19% lower than that of batch #1. Fracture toughness was equal for "H" and "O" orientations. Although "V/H" and "V/V" very low differences (+5% and -7.5% respectively) could be attributed to scatter, it is interesting to note that they have the same tendency than in batch #1, that is: slightly higher toughness for "V/H" samples, and lower but similar values for "V/V", "H" and "O" samples.

The most important point is the 57% average increase (+48% to +63% according to the orientations) in the average fracture toughness of batches #1 and #2. Unlike the tensile characteristics, the HIP treatment greatly improved the fracture strength of the samples.

**Table 4: Results of toughness tests for batch #1**

Specimen orientation		H	O	V/H	V/V
Nb of samples (*) <i>(valid) (***)</i>		4 (4)	4 (4)	4 (4)	3 (3)
N <sub>cycles</sub>	Aver.	46275	48765	48639	40732
K <sub>IC</sub> /K <sub>Q</sub> (**) [MPa.√m]	<b>Aver. (Std dev %)</b>	<b>40.3 (3.4%)</b>	<b>38.9 (6.3%)</b>	<b>45.6 (4.3%)</b>	<b>38.1 (1.6%)</b>

(\*) Due to manufacturing hazards, several samples of each orientation in batch #1 were cracked and could not be tested.

(\*\*) The results for K<sub>IC</sub> for batch #1 were all valid according to ASTM E399.

(\*\*\*) According to the P<sub>max</sub>/P<sub>Q</sub> condition in ASTM E399.

**Table 5 - Results of toughness tests for batch #2**

Specimen orientation		H	O	V/H	V/V
Nb of samples (*) <i>(valid) (***)</i>		5 (4)	2 (2)	4 (3)	2 (2)
N <sub>cycles</sub>	Aver.	38471	39998	38549	38050
K <sub>IC</sub> /K <sub>Q</sub> (**) [MPa.√m]	<b>Aver. (Std dev %)</b>	<b>64.9 (1.6%)</b>	<b>63.6 (0.5%)</b>	<b>67.3 (5.3%)</b>	<b>59.4 (0.2%)</b>

(\*) Due to manufacturing hazards, several samples of each orientation in batch #2 were cracked and could not be tested.

(\*\*) The results for K<sub>IC</sub> for batch #2 were not all considered as valid according to norm ASTM E399, due to the P<sub>max</sub>/P<sub>Q</sub> condition. Specimens that failed to meet the condition were very close to reach it though.

(\*\*\*) According to the P<sub>max</sub>/P<sub>Q</sub> condition.

#### 4. DISCUSSION

One must first note that if the LBM process lead to metallographic anisotropy of the material, such anisotropy was not significantly observed in the measured tensile properties. Nevertheless, the layered microstructure has a notable influence on the toughness values, as the "V/H" samples

exhibited higher characteristics than the other three sample orientations ("O", "H", "V/V"). These later presented similar toughness values. This could be easily explained by the fact that the interface between layers acted as grain boundaries, limiting the crack propagation and so leading to a higher toughness. This explanation could also explain why the "H" and "V/V" samples showed similar toughness (no additional layer interface to overcome). The "O" values were surprisingly similar to the "H" and "V/V" ones, although it could be expected to exhibit an intermediate behaviour between the "H" and "V/H" ones.

The HIP treatment applied almost restored the complete isotropy to the samples from both a metallographic and tensile properties points of view. The anisotropic behaviour was still remaining in the toughness test for the "V/H" orientation. As presented in other works, the HIP treatment (the role of which is to reduce or erase any porosity) mostly allowed larger plastic deformation (+59%) by strongly reducing the residual stresses induced by the LBM manufacturing. This higher plasticity was twined to a notable decrease in the yield and ultimate stresses. Unlike in [4, 31], observations did not reveal any notable decrease in the porosity rate, but this may be attributed to a very low initial rate: 0.2%. Nevertheless, the potential residual porosity was not detected on the fracture surface. This indicates that the plasticity was sufficient to strongly reduce the influence of the porosity, which was not anymore a crack initiation location. This could explain the resulting 57% increase in the toughness of the HIPed samples. The HIP heat treatment is then recommended for LBM manufactured Ti-6Al-4V samples submitted to fatigue solicitations.

## **5. CONCLUSIONS**

The obtained results seem to be promising for the Ti-6Al-4V material even if some other characterization are still missing. Further studied are under investigation concerning the toughness fracture surface and the influence of potential defects within the raw material.

Similar investigations are also in progress for AlSi7Mg0.6 material. The influence of building orientation and heat treatments after LBM manufacturing, as well as internal defects, will be soon presented in another paper.

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