

L-PBF processing and characterization of a Ti 35 Nb 30 Zr 29 Mo 3 Ta 3 multiprincipal element alloy for medical implants

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L-PBF processing and characterization of a Ti₃₅Nb₃₀Zr₂₉Mo₃Ta₃ multiprincipal element alloy for medical implants 2

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ABSTRACT 16

17 This letter reports on the processing, mechanical properties, electrochemical performance and toxicity 18 behavior of a multiprincipal element $Ti_{35}Nb_{30}Zr_{29}Mo_3Ta_3$ alloy. The recovered centimeter-sized cubes 19 or parallelepipeds were polished before analysis. It was found that the homogeneity and yield stress of 20 the alloys get increased when higher volumetric energy densities (VEDs) are applied during 21 processing. The synthesized alloys also exhibited a Young modulus closer to those of bones than those 22 usually reported on the conventional orthopedic metals (60-80 GPa for this alloy against 110 and 4-30 23 GPa for the Ti-6Al-4V and bone, respectively). The electrochemical corrosion assays performed at 25°C and 37°C, in sodium chloride (NaCl, 0.1 M), deoxygenated high glucose Dulbecco's Modified 24 Eagle Medium (DMEM) and deoxygenated DMEM implemented with Fetal Bovine Serum (FBS) 25 showed corrosion potentials (E_{corr}) ranged in the order NaCl_(25°C) \approx DMEM + PBS_(37°C) > DMEM_(37°C), 26 27 and impedances much larger in the DMEM and DMEM + PBS media than in NaCl. Finally, preosteoblasts cells cultured in the medium conditioned by the alloy did not evidence cytotoxicity effect, 28 because no effects on the cell proliferation and morphology were observed. These data suggest the 29 biocompatibility of this alloy, indicating no acute toxicity, which is a prerequisite for the integration of 30 an alloy as a bioimplant. 31

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33 Faced with the aging population in our modern societies and the increased risk of bone diseases and 34 accidents in these populations, the development of long-lasting implant materials is a significant

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public health issue. It is, nowadays, essential to adapt advanced structural materials to extend the net duration of the revision surgery beyond the current limit of ten years.¹ According to the recent United Nations reports, the over 60s population of the world is growing, and the proportion of the old persons is expected to increase by 45% by 2050, with a life expectancy that could gain another five additional years in the developed countries,² making this period of ten years before the surgical revision extremely short. The problem of the longer-lasting implants also concerns people with acute and chronic bone diseases while maintaining an acceptable perspective and quality of life.

Multiprincipal element alloys (MPEAs) are a new class of materials that offer a range of unique 42 properties, such as high strength and corrosion resistance.^{3,4} These properties make them ideal 43 candidates for designing medical devices. Moreover, by adjusting the composition of the MPEAs, it is 44 45 possible to create materials that are optimized for specific applications. For instance, their Young's modulus can be reduced to values close to those of bones (20 to 30 GPa), without losing their yield 46 strength, thereby reducing the probability of the occurrence of osteoporosis near the implant. 47 Consequently, the MPEAs are increasingly being used in the medical industry for various 48 applications.⁵ 49

50 In parallel, the emergence of the additive manufacturing (AM) has revolutionized the production of 51 complex and intricate structures, previously impossible to be manufactured⁶; opening thus real 52 opportunities for medical implant processing, meaning that AM allows the production of medical 53 implants tailored to the patient's needs, matching correctly the targeted function.

The technology behind the AM is based on the principles of the layer-by-layer deposition of materials. This means that the implant is built up from a series of thin layers, and each layer is deposited on top of the previous one. In this way, the AM eliminates the need for the expensive and time-consuming tooling. It also enables the production of porous structures, which can serve as channels for drug storage and then controlled delivery,⁷ as it allows surface texturing, which can be useful for tissue integration and rejection risk reduction.⁸

Altogether, the AM designed MPEAs-based biomedical implants have the potential to revolutionize
healthcare.⁹ Combining these two innovative concepts, the implants composition and morphology

would be easily tailored to meet patients' anatomy, age, and gender, laying the ground for personalizedtreatment.

In this study, a series of the MPEA with the nominal composition of $Ti_{39}Nb_{28}Zr_{28}Mo_{2.5}Ta_{2.5}$ (at. %) were elaborated by the laser powder bed fusion (L-PBF), using the selected laser melting (SLM) technology (see Table 1S for the actual achieved composition). By adjusting the fabrication parameters, especially the volumetric energy density (VED) during the SLM processing, the Young's modulus of the alloy was significantly reduced, without losing its yield strength, reducing thereby the probability of osteoporosis occurrence near the implant.

70 Through the processing optimization, several types of microstructural defects were identified
71 (experimental details in the supplementary data). These were mainly unmelted powders, cracks, and
72 pits.

Melting elements like Ta, Mo and Nb with high melting points (3020, 2623 and 2477°C, respectively)
is obviously arduous. The low VED resulted in the accumulation of unmelted Nb/Mo/Ta-rich islands.
Scanning electron microscopy (SEM) images clearly evidence these entities by their after polishing
relief and light grey contrast (as well as the Ta/Ta-rich islands by their bright light contrast), as shown
in Figs. 1(a) and 1S.

78 Increasing the VED induced the Nb/Mo/Ta-rich islands solubilization, improving the homogeneity of 79 the final alloy (Fig. 1(b, c)). Nevertheless, the solubilization of Nb/Mo/Ta into the matrix triggers the 80 vaporization of Ti, since the melting point of Ta is close to the boiling point of Ti (3287°C). As a consequence, typical processing defects, such as keyhole porosities form. The higher the VED, the 81 82 higher the keyhole porosities fraction was in the structure of the alloy. Their appearance frequency, 83 means the coalescence of the cavities while porosities facilitate the degradation of the mechanical properties of the alloy.¹² Contrariwise, cracks cause less dramatic effects on the mechanical properties 84 85 of the material. Cracking occurs either as a result of the residual stress relaxations (induced by the high cooling rate of the SLM) or the complex solidification mechanisms (involving the solidification range 86 87 and liquid viscosity) (Fig. 1(e, f)). The second case usually forms short/thin intergranular cracks, primarily along the high-angle grain boundaries (HAGBs). 88

The VED affects also the grain morphology, size and size distribution. The inverse pole figure (IPF) electron backscatter diffraction (EBSD) images recorded on the specimens, elaborated at the VED = 150, 450, and 550 J.mm⁻³, clearly evidence it (Fig. 2(a-c)). These images show heterogeneous grain shapes. The degree of heterogeneity decreases with the applied VED. For the VED = 550 J.mm⁻³, a relatively bimodal character was observed, while for the VED = 450 J.mm⁻³, a slight <100> texture was noted.

95 On a finer scale, the grains appeared as made up of dendritic cells whose sizes increased with 96 increasing the VED, from 429 up to 516, and 740 nm, for the VED = 150, 450 and 550 J.mm⁻³, 97 respectively (Fig. 2(d-f)). The statistical details of the microstructure of the alloys can be seen in Fig. 98 2(g, h). These observations are in agreement with the existing literature data,^{10,11} where it has been 99 reported that the walls between the cells are made of accumulated geometrically necessary 100 dislocations, inducing the disorientations in the order of 2 degrees. Furthermore, the dendritic cell 101 walls were rich in Zr, while Mo was accumulated mostly inside the cells (Fig. 2S).

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VED 150

VED 450

VED 550



FIG. 1. Nature of the processing defects observed as a function of the applied VED. (a, d) 150 J.mm⁻³, (b, e) 450 J.mm⁻³ and (c, f) 550 J.mm⁻³.

108	The number of the HAGBs increased with increasing the VED. The average degrees of
109	misorientations were 24, 28, and 32° for the VED = 150, 450, and 550 J.mm ⁻³ , respectively (Fig. 2(g)).
110	The high fraction of low angle grain boundary (LAGB) misorientation at low VED is in line with the
111	cells size given in Fig. 2(h). Indeed, smaller cell sizes (longer cell walls) increased the degree of the
112	low angle misorientation. The increase in the HAGBs' percentage coincided with the solidification
113	cracking probably due to the increase in the disorientation between the contiguous grains, in
114	agreement with the literature data. ¹²
115	Apart from the evolution of the size of the dendritic cells which increased with the VED (Fig. 2(d-f)),
116	the average grain size of the alloys (measured as area fraction) decreased as the VED increased. The
117	latter is attributed to the bimodality of the alloy structure (Fig. 2(a-c)).



FIG. 2. Microstructure quantification details. (a-c) IP EBSD images superimposed to the grain
boundaries map of the VED = 150, 450 and 550 J.mm⁻³ specimens, respectively, (d-f) SEM images of
the cellular substructure of each specimen and (g, h) distribution of the misorientation across the grain
boundaries, frequency of grain, and cellular size as a function of the applied VED (Z axis in Fig (g)
indicates the laser beam direction).

The results of the compression and tension tests can be compared in Fig. 3(a, b). See also experimental details in the supplementary data. Fig. 3(a) depicts that all the as-processed alloys possess a high ability to get plastically deformed. Indeed, all the specimens exhibited nominal strains well above 70%. The yield stresses were found to be in the range from 900 to 1350 MPa, increasing with the VED increase. As expected, the above-mentioned processing defects did not compromise the use of these alloys in the loading conditions. Contrariwise, the tensile tests showed a high sensitivity to the microstructure. The higher the VED, the higher were the yield stress and yield strength values. For

instance, the VED = 550 J.mm^{-3} led to a relatively poor ductility. The related specimen is in the regime 133 134 with the highest keyhole porosities frequency, along with the solidification cracking. The coalescence 135 of these cavities during the straining and bridging of the microcracks into a three-dimensional 136 network, contributed to the observation of the lack of ductility. Fig. 3(c, d) illustrates the necking regions of the 550 and 450 J.mm⁻³ specimens, respectively. The elongation of the Ta-rich entities into 137 a lenticular shape in the latter is evident; since opposite to the 550 J.mm⁻³ specimen, here they 138 contributed to the overall ductility of the material. Images of the generated microcracks in the 450 139 J.mm⁻³ specimen are also given in Fig. 3(d, e). 140

Here the increase in the yield strength by increasing the VED can be explained by a Hall-Petch-like effect combined with a solid solution strengthening. The Hall-Petch-like effect comes from the reduction of the grain size while the origin of the solid solution strengthening comes from the solubilization of Nb/Mo/Ta in the matrix.

145 Table 1 summarizes the major compression and tensile loading parameters of the alloys. Here the Young's modulus E values (60 to 80 GPa) are much lower than those of the materials that are 146 147 currently used for the orthopedic applications (CP-Ti: 105, Ti-6Al-4V: 115 and Co-Cr: 240 GPa). The low E values are expected to eliminate the stress shielding effect of the material.^{13,14} Another 148 biomechanical indicator for the metallic implant materials, is the elastic strain deformation ε .¹⁵ It was 149 150 computed, for this study, from the tensile test data as $\varepsilon = \sigma_{YS}/E$. The obtained values were found to be 151 varying in the range from 0.87 to 1.22%, larger than those of the CP-Ti and Ti6Al4V (0.47 and 0.78%, respectively), but closer to that of the as-cast TiZrNbTa (1.26%).¹⁶ 152



FIG. 3. (a) Typical stress-strain compression test plots of the synthesized alloys along the Z axis/OXY
plane (building direction), (b) tensile test plots along the OYZ plane, (c, d) failure profiles of the
tensile tested 550 and 450 J.mm⁻³ specimens, respectively and (e) crack propagation along the HAGBs
in the tensile tested 450 J.mm⁻³ specimen.

160	Table 1. Some mechanical	parameters of the 150), $450 \text{ and } 550 \text{ J.mm}^{-3}$	specimens.
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specimen	Young modulus [GPa]	ultimate tensile stress [MPa]	yield strength [MPa]	strain to failure [%]	compression yield strength (MPa)
$\mathbf{VED} = 150 \ \mathbf{J.mm^{-3}}$	60	674	527	13.38	940
$\mathbf{VED} = 450 \ \mathbf{J.mm^{-3}}$	78	1016	950	17.99	1215
$\mathbf{VED} = 550 \ \mathbf{J.mm^{-3}}$	80	1061	964	3.23	1335

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Altogether, these results suggest that the 150 and 550 J.mm⁻³ specimens are the best according to the *E* and ε criterions, respectively; for the desired application, while the 450 J.mm⁻³ one offering finally the best compromise. In the same time, the 550 J.mm⁻³ specimen exhibited the best chemical homogeneity and then very probably a better corrosion performance. The persistence of a larger number of the Nb/Mo/Ta-rich islands in the 450 J.mm⁻³ specimen compared to the 550 J.mm⁻³ one, might favor pitting corrosion, as a consequence of the (electro)chemical reactivity differences between these islands and the matrix. Also differences between the biological behaviors might be expected between the islands and matrix, when contacted to bone cells. Consequently, to pursue our investigations, a
 focus was made on the 550 J.mm⁻³ specimen.

The electrochemical measurements were thus performed on both 150 and 550 J.mm⁻³ specimens to 171 investigate their corrosion performance. The specimens were exposed to a 0.1 M NaCl (25°C) solution 172 and to a 37°C deoxygenated Dulbecco's Modified Eagle mammal cell culture medium (DMEM) or 173 DMEM supplemented with 20% of fetal bovine serum (FBS) (experimental details in the 174 175 supplementary data). After ~ 24 h of exposure, their potentiodynamic polarization curves and impedance plots were recorded (Figs. 4(a, b) and 3S(a-d)). Each open circuit potential (OCP) value, 176 177 given in the inset of the figures is the average potential of at least three electrodes, before being polarized. The corrosion potential (E_{corr}) , as well as the corrosion current density (I_{corr}) values for each 178 179 coupon in each electrolyte are given in these figures. Along with the potentiodynamic polarization 180 curves, some selected SEM images of the pits nucleated during the polarization of the coupons in the DMEM and NaCl electrolytes can be seen in Figs. 4(a) and 3S(b). (more images can be found in Fig. 181 4S). 182

183 Interestingly, the anodic branches of all the curves (Figs. 4(a) and 3S(b)) exhibited the generation of a large oxidation current, especially in $\sim +0.7$ V/SHE. This indicates that by polarizing the coupons 184 185 towards more positive potentials, a considerable degree of dissolution occurs on their surfaces. The OCP of the 550 J.mm⁻³ alloy in the NaCl ($pH_{(25^{\circ}C)} = 6.55$) and DMEM + FBS ($pH_{(25^{\circ})} = 7.70$) solutions 186 187 is the most positive (+0.150 V/SHE). This potential in the DMEM ($pH_{(25^{\circ}C)} = 7.70$) solution changes to -0.035 V/SHE. The curves in Fig. 4(a) also depict that although the I_{corr} recorded in the NaCl solution 188 189 is slightly larger than those of the DMEM and DMEM + FBS solutions, the alloy generated a smaller 190 oxidation current in this solution, compared to the two others.

191 The recorded Nyquist plots of the 550 J.mm⁻³ coupons, show that the resistivity of the alloy in the 192 three solutions is in the order DMEM + FBS > DMEM > NaCl. This can be seen in the polarization 193 resistance and imaginary impedance values in Fig. 4(b). The plot recorded in the NaCl solution has an 194 incomplete/deformed semi-circle which can be fitted with an electrical circuit model based on a 195 constant phase element (*CPE*), parallel to a charge transfer resistance (R_{ct}) element; both in series with 196 a solution resistivity (R_s) element. In DMEM, the recorded plot consists of only a deformed semi-

- 197 infinite Warburg linear diffusion impedance, that is represented with a Warburg element (*W*) in series 198 with a R_s (Fig. 4(b)). Here it should be noted that the slope of a classic Warburg plot is 45° and plots 199 with degrees > 45° indicate the existence of a greater mass transport limitation. In DMEM + FBS, the 200 recorded plot has an incomplete/deformed semi-circle in the high frequency region, followed with a 201 linear diffusion Warburg impedance, which can be modelled with an electrical circuit based on a *W* 202 element in series with a R_{ct} and parallel to a *CPE*.
- Comparing the Bode total impedance (|Z|) versus frequency plots of the two alloy coupons in the three electrolytes (Fig. 3S(a, d)) allows evidencing that the |Z| of the 550 J.mm⁻³ coupon at the minimum applied frequency is the maximum and minimum for the DMEM and NaCl solutions, respectively. While, in the case of the 150 J.mm⁻³ coupon the largest and smallest minimum frequency |Z| belongs to the plots recorded in the DMEM and DMEM + FBS.
- To resume the overall electrochemical results, it is clear that the 550 J.mm⁻³ alloy, has a better resistivity to corrosion compared to the 150 J.mm⁻³ alloy, especially in the DMEM and DMEM + FBS media, at least for the 24 h of exposure, which is useful for the desired application.





a deoxygenated DMEM + FBS (80:20 (v/v)) (37° C) media. Inset: SEM images of the pits, nucleated during the polarization of the alloy in DMEM.

Finally, the toxicity of the 550 J.mm⁻³ alloy was evaluated by studying the cell proliferation and morphology in the alloy conditioned and normal media. The conditioned media were prepared by immersing a freshly prepared 550 J.mm⁻³ coupon for 24 h in a 10 mL DMEM-F₁₂ medium.

Cells were thus stained (their nucleus and actin cytoskeleton in yellow and purple, respectively) (the experimental details in the supplementary data) and optical images were collected and compared (Fig. 5). The images in Fig. 5 revealed no morphological differences between the cells cultured in the control medium compared to those cultured in the medium conditioned by the alloy. This was consistent with the study of the cell proliferation over 48 h. The number of the viable cells is nearly identical between the cell populations, regardless of whether they were exposed to the conditioned medium or not.

228 These observations are a strong indication of no acute toxicity of the alloy, suggesting its229 biocompatibility.

Since it is commonly accepted that the cell proliferation performance gets enhanced by modification of the surface morphology,^{17,18} the effect of the surface microstructure, in term of roughness, will be further investigated. Application of the bioactive peptide coatings is also currently under investigation to improve the cell viability, adhesion and proliferation.¹⁹ Up to now, only nude coupons were tested. Here despite the lack of the surface optimization, the biological preliminary results supported the choice of the elaborated alloy as an implant material.

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FIG. 5. Optical microscopic images of the MC_3T_3 A A pre-osteoblasts cells cultured in the nonconditioned ((a) 24 h, (b) 48 h) and conditioned media ((c) 24 h, (d) 48 h) and (e) the proliferation graphic (the line within the box represents the median). ANOVA test results: n = 3, * p < 0.05, ** p < 0.01.

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In summary, the L-PBF processed $Ti_{35}Nb_{30}Zr_{29}Mo_3Ta_3$ alloys were investigated, optimizing the elaboration parameters, mainly VED, to tentatively reach a valuable microstructure, mechanical properties, electrochemical corrosion parameters and a biological compatibility toward mammal osteoblast cell proliferation, for a successful integration as bone bioimplants.

247 Increasing the VED induced the Nb/Mo/Ta solubilization, improving the chemical homogeneity of the 248 processed alloys. Besides, the alloys had a high ability to get plastically deformed. They also exhibited nominal strains of >> 70%, Young modulus values much lower than those of the conventional implant 249 250 alloys, elastic strains close to those of bones and yield stresses from 900 to 1350 MPa, that increased 251 with increasing the VED, proving the capability of the alloy to be used as a bioimplant material. 252 Focusing on the specimens processed at high VED, their OCPs in the NaCl and DMEM + FBS media 253 were found to be more positive, compared to DMEM, while the impedance parameters of the alloy were much more desirable in contact with the DMEM and DMEM + FBS. Finally, the same 254

specimen, did not induce acute toxicity and did not inhibit cell proliferation, proving itsbiocompatibility, supporting their valuability as an implant material.

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258 Supplementary Material

The supplementary data includes the detailed information about the elaboration process,
microstructure/mechanical characterization, electrochemical corrosion performance and cytotoxicity
behavior of the synthesized alloys.

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268 Author Contributions

269 Equal: conceptualization, Equal: data curation, formal analysis, writing original draft and project270 administration.

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272 Data Availability

The data that support the findings of this study are available within the letter and its supplementarymaterial.

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