

# EOS Titanium Ti64 ELI

EOS Titanium Ti64ELI is a titanium alloy powder intended for processing on EOS DMLS<sup>™</sup> machines. This document provides information and data for parts built using EOS Titanium Ti64ELI powder (EOS art.-no. 9011-0040) on the following system setup:

- EOS DMLS<sup>™</sup> system: EOS M400 SF
  - HSS recoating blade
  - Argon atmosphere
  - IPCM M extra sieving module with  $63 \mu m$  mesh recommended
- EOSPRINT v.1.5/HCS v.2.4 or newer
- EOS Parameter set Ti64ELI\_030\_FlexM400\_100

## Description

Parts built in EOS Titanium Ti64ELI have a chemical composition corresponding to ASTM F136 and ASTM F3001.

Ti64ELI is well-known light alloy, characterized by having excellent mechanical properties and corrosion resistance combined with low specific weight and biocompatibility. This material is ideal for many high-performance applications.

Parts built with EOS Titanium Ti64ELI powder can be machined, shot-peened and polished in asbuilt and heat treated states. Due to the layerwise building method, the parts have a certain anisotropy.

## **Quality Assurance**

The quality of the EOS Titanium Ti64 powder lots is ensured by the Quality Assurance procedures. The procedures include sampling (ASTM B215), PSD analysis (ISO 13320), chemical analyses (ASTM E2371, ASTM E1409, ASTM E1941, ASTM E1447), and mechanical testing (ISO 6892-1).

The results of the quality assurance tests are given in the lot specific Mill Test Certificates (MTC) according to EN 10204 type 3.1.

Electro Optical Systems Finland Oy

Robert-Stirling-Ring 1 D-82152 Krailling / München

FIN-20520 Turku Telephone: +358 23358119 Telefax: +358 20 7659141

Lemminkäisenkatu 36



## **Technical Data**

### **Powder properties**

laterial composition [wt.%]	Element	Min	Max
	AI	5.50	6.50
	V	3.50	4.50
	0	-	0.13
	Ν	-	0.05
	С	-	0.08
	Н	-	0.012
	Fe	_	0.25
	Y	-	0.005
	Other element each	5, _	0.10
	Other element total	5, _	0.40
	Ti		bal.

Particle size	
d50 [1]	39 ± 3 μm

[1] Particle size distribution analysis according to ISO 13320

### General process data

Layer thickness	30 μm
Volume rate [2]	5 mm³/s (18 cm³/h)

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Telefax: +358 20 7659141

EOS GmbH Electro Optical Systems

Robert-Stirling-Ring 1 D-82152 Krailling / München

Telephone: +49 (0)89 / 893 36-0

Telefax: +49 (0)89 / 893 36-285

www.eos.info

Internet:



[2] The volume rate is a measure of build speed during laser exposure of the skin area. The total build speed depends on this volume rate and many other factors such as exposure parameters of contours, supports, up and downskin, recoating time, Home-In or LPM settings.

### Physical properties of parts\*

Part density [3]	4.4 g/cm3
Surface roughness after shot peening [4]	Approx. R₂ 5-10 μm; R₂ 15-30 μm
Hardness as built [5]	typ. 340 HV5
[3] Weighing in air and water according to ISO 3369.	

[4] The numbers were measured at the horizontal (up-facing) and all vertical surfaces of test cubes. Due to the layerwise building the roughness strongly depends on the orientation of the surface, for example sloping and curved surfaces exhibit a stair-step effect.

[5] Hardness measurement according to standard EN ISO 6507-1 with load 5kgf (HV5)

### Tensile data at room temeprature\* [6,7]

	As built	Heat treated [8]
Ultimate tensile strength	typ. 1270 MPa	typ. 1040 MPa
Yield strength, Rp0.2%	typ. 1100 MPa	typ. 930 MPa
Elongation at break A	typ. 8,7 %	typ. 14,0 %

[6] The numbers are average values and are determined from samples with horizontal and vertical orientation.

[7] Tensile testing according to ISO 6892-1 A14, proportional test pieces, diameter of the neck area 5 mm (0.2 inch), original gauge length 20 mm (0,79 inch).

[8] Heat treatment procedure: 2 hours at 800°C in Argon atmosphere.

#### Abbreviations

min. minimum

max. maximum

wt. weight

#### Electro Optical Systems Finland Oy

Lemminkäisenkatu 36 FIN-20520 Turku

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This powder has not been developed, tested or certified as a medical device according to Directive 93/42/EEC (MDD) or Regulation (EU) 2017/745 (MDR) and is not intended to be used as a medical device, in particular for the purposes specified in Art. 2 No. 1 MDR. Insofar as you intend to use the powder as raw material for the manufacture of pharmaceutical products or medical devices (e.g. as raw material which as a material must meet the requirements of Annex 1, Chapter II MDR), the responsibility and liability for all analyses, tests, evaluations, procedures, risk assessments, conformity assessments, approval and certification procedures as well as for all other official and regulatory measures required for this purpose shall lie solely with you both with regard to the pharmaceutical product and/or medical device manufactured by you and with regard to the properties, suitability, testing, evaluation, risk assessment, other requirements for use of the powder as raw material. This also applies to applications with food contact. In this respect, the limitations of liability pursuant to our General Terms and Conditions and the system sales or material contracts shall apply.

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Electro Optical Systems Finland Oy

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