

DECLARATION OF CONFORMITY

according to annex II without 4, 93/42/EEC

- **Manufacturer:**

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We herewith declare under sole responsibility that the product

- **Name of product family:** **SLM Powders**
- **Name of products:** **WIRONIUM® RP**
- **REF number:** **50530**
- **Product class:** **Class IIa**

meets the relevant requirements of the EC Directive **93/42/EEC** concerning medical devices and is in conformity with the list of applied standards in the technical documentation.

This declaration of conformity is valid until expiration of the EC-Certificate 93/42/EEC (Number HD 60142369 0001) on 26 May 2024 and only together with the related batch release document.

- **Notified body:**

TÜV Rheinland LGA Products GmbH
 Tillystraße 2
 90431 Nürnberg, Germany

Notified body: **CE 0197**

Bremen, 28.05.2021

Place, Date


 Signature
 Chief Developer and Innovation
 Officer


 Signature
 Director of Quality Management
 and Regulatory Affairs


 Signature
 CEO