TÜVRheinland®

EC Certificate

Directive 93/42/EEC Annex II, excluding Section 4
Full Quality Assurance System
Medical Devices

Registration No.: HD 60135612 0001

Report No.: 26300444 002

Manufacturer:
Oxipit, UAB
Sauletekio al. 15
10224 Vilnius
Lithuania

Products:
Class IIa:
Standalone computer assisted diagnosis medical device software for chest X-ray images analysis and reporting

Expiry Date: 2023-11-21

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II, section 4 is required.

Effective Date: 2019-01-28

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TÜV Rheinland LGA Products GmbH Zertifizierungsstelle

Notified Body

Sebastian Mniszek

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg
TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.