

EC Certificate



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

Registration No.: HD 1169866-1

Manufacturer: Ziehm Imaging GmbH
Lina-Ammon-Str. 10
90471 Nürnberg
Germany

Products: mobile Radiographic/Fluoroscopic Units (mobile C-arms)

Replaces Certificate, Registration No.: HD 60144926 0001

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.

Report No.: 1083421-20

Effective date: 2021-03-11

Expiry date: 2024-05-26

Issue date: 2021-03-11




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