

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

No. 5-630-200-0911

Institute for Healthcare Quality Improvement and Hospital Engineering (EMKI)
certifies that the manufacturer

MEDIMA Sp. z o. o.
Al. Jerozolimskie 200
02-486 Warsaw
Poland

medima

for the products/product categories

Syringe infusion pumps
Volumetric infusion pumps
Docking stations for infusion pumps
Accessories for infusion pumps
Medical software

applies a quality system which ensures that the products conform to the provisions of the Directive which apply to them at every stage from design to final inspection.

Registry number of the related audit report: **42-131-2007**

This Certificate is valid until **2014-11-19** supposed that the results of the regular yearly surveillance audits are satisfactory.

Issued by EMKI as a Notified Body for the Council Directive 93/42/EEC with identification number

1011.

This certificate is valid only with the attachment.

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Budapest, 2011-03-03


General Director




Certification Office



EMKI 0372

The validity of the certificate is verifiable at the Certification Office of EMKI.

Egészségügyi Minőségfejlesztési és Kórháztechnikai Intézet
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