

QUALITY MANAGEMENT SYSTEM CERTIFICATE

No. 4-510-135-1911

The **Directorate of Device Testing and Clinical Engineering (EMKI)**
as a Certification Body with ID No. NAH-4-0096/2016
accredited by the National Accreditation Authority for management system certification
certifies that the quality management system applied by

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

meets the requirements of standard

EN ISO 13485:2016

in the field:

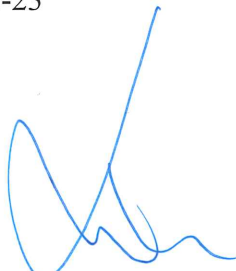
**Developing, manufacturing, marketing and service
of syringe infusion pumps, volumetric infusion pumps, docking stations,
medical software and infusion sets**

Registry number of the related audit report: **43-131-2007**

This certificate is valid until **2022-11-22** supposed that the results of the regular yearly surveillance audits are satisfactory.

The Company has been certified by EMKI since 2004-11-22.

Budapest, 2019-11-23



Head of EMKI



EMKI 2266

The authenticity and validity of the certificate are verifiable at EMKI.

Eszközminősítő és Kórháztechnikai Igazgatóság
Directorate of Device Testing and Clinical Engineering

H-1097 Budapest, Albert Flórián út 3/A, Telefon: +36 20 268 75 95, Fax: +36 1 886 93 33

E-mail: cert@emki.hu, Web: www.emki.hu

H-1051 Budapest, Zrínyi u. 3. (1372 P.O. Box 450.)

EMKI