

EC Certificate**Full Quality Assurance System according to
Medical Devices Directive 93/42/EEC Annex-II Section 3****Certificate Number: 1984-MDD-17-428**

We hereby declare that an examination of the under mentioned full quality assurance system has been carried out following the requirements of the national legislation to which the undersigned is subjected, transposing annex II (with the exemption of section 4) of the Directive 93/42/EEC on medical devices. We certify that the full quality assurance system conforms with the relevant provisions of the aforementioned directive.

Organization:**OKUMAN MEDİKAL SİSTEMLER ANONİM ŞİRKETİ**

HQ: Zübeyde Hanım Mahallesi Kazım Karabekir Caddesi No:95/95
06060 İskitler, Ankara, Turkey

Factory: İvedik Organize San. Bölgesi Arı Sanayi Sitesi 1.Etap 1417 Sok.
No:51 Yenimahalle, Ankara, Turkey

Products: Intensive Care Phototherapy, Radiant Warmer, Infant Incubator, Multi-Parameter Patient Monitors, Infant Ventilator, Transport Incubator, Defibrillator Monitor, Warming Bed

The products defined at the enclosure which is the part of this certificate and contains one (1) pages. The certificate is valid till expiration date, subject to successful completion of periodical surveillance audits. Please contact Kiwa for details.

Report Number: M.3222.10
Date of first issue: 07 March 2017
Date of last issue: 30 April 2021
Revision Number: 06
Expiry Date: 27 May 2024



Muhteşem Gökhan Yücel
Head of Notified Body

30 April 2021, Istanbul, Turkey

Enclosure of the EC Certificate:

Page 1/1

Full Quality Assurance System according to

Medical Devices Directive 93/42/EEC Annex-II Section 3

Certificate Number: 1984-MDD-17-428, Revision Number: 06

Concerned medical devices;

Product: Intensive Care Phototherapy

Model Number: BiliCare

Product: Radiant Warmer

Model Number: OKM 730

Product: Infant Incubator

Model Number: OKM 801

Product: Infant Ventilator

Model Number: OKM IBS

Product: Transport Incubator

Model Number: TR 203

Product: Multi-Parameter Patient Monitors

Model Number: OKM VS3, OKM 300, OKM 500, OKM 600, OKM 700, OKM 800, OKM 860, OKM 900, OKM 84, OKM 104, OKM 121, OKM 8800

Product: Defibrillator Monitor

Model Number: DFM600, DFM800

Product: Warming Bed

Model Number: OKM 740

Kiwa Belgelendirme Hizmetleri A.Ş. is Notified Body under Council Directive 93/42/EEC concerning medical devices with identification number: 1984

A handwritten signature in black ink, appearing to read "Muhteşem Gökhan Yücel".

30 April 2021, Istanbul, Turkey

Muhteşem Gökhan Yücel
Head of Notified Body