CERTIFICATE
FOR CE MARK CLASS I MD

™Acting as the Regulatory Authorized Representative in the EU for:

Truphatek International Ltd,
9 Haomanot Street
P.O. Box 8051
Netanya 42504 Israel

We confirm that the CE mark registration of the devices listed below have been completed, as required for Class I medical Devices under the MDD 93/42/EEC. The registration was made with the Medicinal and Healthcare Products Regulatory Agency in the UK (MHRA)

Class I devices registered:

- Truview™ , EVO2, PCD™ and PCD-R™ Range of blades, handles, camera, monitor and accessories, including
  - Truflex™ articulated stylet and Optishape™ stylet
  - PCD Optical View Tube
  - FibreClip™ for Truview PCD™
  - Power Supply for PCD screen
  - Truview Stand Organizer
  - Infant Intubation Manikin
- Shucman™ 2 range of Laryngoscopes, including fibre clip bundles
- Equip™ Range of Laryngoscopes
- EquipLED™ Range of Laryngoscopes
- Green Spec™ and Green Spec 2™ range of Laryngoscopes, including fibre clip bundles
- Tru-MR™ MR Conditional Range of Laryngoscopes with handle and battery packs
- Ultra Safe™, Standard Paediatric, Green SpeX™, GreenLed™ (including battery cartridge), Dolphin™ ranges of Handles
- Truled™ and Truled PCD ™ rechargeable Handle range including battery cartridges and power supply adaptors
- Equip Lite ™ Range of Laryngoscope blades
- Trulite™ Range of one -piece Disposable Laryngoscope blade/ handle combination
- Lite Blade™ and Lite Blade Slim™ Plastic Range of Laryngoscopes
- GreenLite™ Metal Disposable Range of Laryngoscopes
- DISPOLED™ and DISPOGRIP ™ disposable range of Laryngoscope handles
- Krypton and Xenon Bulbs

Registration number with the MHRA: **CA7730.**

Date: 23/08/15
Benny Arazy
General Manager
Medes Limited

MEDES LTD.

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Regulatory Authorized Representative

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