

# CERTIFICATE

**The Certification Body TÜV Rheinland Italia S.r.l.**

certifies, in accordance with the TÜV Rheinland Group procedures, that the Company

**Kabomed Company for Medical Industries L.L.Co.**

**10<sup>th</sup> of Ramadan City, B4 Industrial Zone, Plot No. 174**

**Egypt**

has established and applies a quality management system  
for the following scope:

**Manufacturing, placing on the market of sterile parenteral bag and enteral bag.  
Production for third parties of sterile and non-sterile drainage bag, bloodline for  
haemodialysis, fistula needle for haemodialysis, blood transfusion set, solution infusion  
set, extension line, parenteral bag and enteral bag and non-sterile urine bag.**

Through an Audit proof has been furnished that the  
quality management system fulfils the requirements of the standard

**ISO 13485:2016**

Please refer to the Quality Manual for the details about  
the exclusions with respect to the requirements of the standard.

Certificate Registration No. **39 05 0832506**

This Certificate is valid from 2025/10/20 to 2027/12/22

The reference date for all the next audits is (day-month): 22/12

**Milan, 2025/10/20.** First Certification: 2009/12/22



The certification responsible: Cesare Gentile  
TÜV Rheinland Italia S.r.l., Via E. Mattei, 3 - I - 20005 Pogliano Milanese (MI)

This certificate does not represent proof that the statutory requirements of  
the Directives 93/42/EEC, 90/385/EEC, 98/79/EC or  
Regulations (UE) 2017/745, (UE) 2017/746 have been fulfilled



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