



Product Service

SAO Y BẢN CHÍNH  
Ngày 20 tháng 09 năm 2017

# EC Certificate

## Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)  
(Devices in Class IIa, IIb or III)

No. G1 15 06 38393 015

**Manufacturer:** **MEDICOP d.o.o.**  
Obrtna ulica 43, p.p. 161  
9000 Murska Sobota  
SLOVENIA

**Facility(ies):** MEDICOP d.o.o.  
Obrtna ulica 43, p.p. 161, 9000 Murska Sobota, SLOVENIA

**Product Category(ies):** **Medical suction devices, terminal units for medical gases and vacuum, flowmeter and pressure regulators for medical gases, components for medical gas supply, medical supply units**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

**Report No.:** 713061011

**Valid from:** 2015-08-02

**Valid until:** 2020-08-01



*H.-H.*

Hans-Heiner Junker



GIÁM ĐỐC  
*Nguyễn Đức Sáu*

**Date,** 2015-07-01

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

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