



Product Service

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 14 10 63179 021

Manufacturer: **STORZ & BICKEL GMBH & CO. KG**
Rote Strasse 1
78532 Tuttlingen
GERMANY

Facility(ies): **STORZ & BICKEL GMBH & CO. KG**
Rote Strasse 1, 78532 Tuttlingen, GERMANY

Product Category(ies): **Cannabinoid Vaporizer
for medical purpose**

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.: 713045268

Valid from: 2014-10-09
Valid until: 2019-10-08

Date, 2014-08-19

Hans-Heiner Junker



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

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