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TÜV SÜD

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
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ZLG-BS-244.10.08

TÜV SÜD Product Service

# EC Certificate

Product Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex VI
(Devices in class IIa or IIb)

**No. G3 034454 0013 Rev. 00**

**Manufacturer:**

**Biover AG**
Müliweg 2
6052 Hergiswil
SWITZERLAND

**Facility(ies):**

Biover AG
Müliweg 2, 6052 Hergiswil, SWITZERLAND

**Product Category(ies):**

**Microvascular clamps**

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

**Report no.:** 713142957

**Valid from:** 2019-03-16
**Valid until:** 2024-03-15

**Date,** 2019-02-14

Stefan Preiß


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

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