



America

CERTIFICATE

No. QS6 063179 0068 Rev. 03

Certificate Holder: **STORZ & BICKEL GMBH**
In Grubenäcker 5-9
78532 Tuttlingen
GERMANY

Certification Mark:



Scope of Certificate: **Design and Development, Manufacturing and Distribution of Vaporizer Systems for Release of Cannabinoids out of Cannabis Blossoms (Cannabis Flos), Cannabis Extracts or Pure Cannabinoids for Medical Purpose such as Chronic Pain, Spasticity and Muscle Spasms, Anorexia and Chachexia, Nausea and Emesis**

Standard(s): **ISO 13485:2016**

Regulatory Authority(ies): **Australia TGA, Health Canada, USA FDA. See attached for listing of specific regulatory requirements.**

The Certification Body of TÜV SÜD America Inc. certifies that the quality management system of the manufacturer listed above has been audited against the stated criteria and found to conform to those criteria for the scope of certification listed. For details and certificate validity see:

[www.tuvsud.com/ps-cert?q=cert:QS6 063179 0068 Rev. 03](http://www.tuvsud.com/ps-cert?q=cert:QS6_063179_0068_Rev.03)

TÜV SÜD America Inc. is an MDSAP Recognized Auditing Organization.

REPs Facility ID: **F001736**
Report No.: **713336089**
Effective Date: **2024-10-22**
Expiry Date: **2027-10-21**

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Date of Issue: 2024-08-23

(Renee Walker)



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Regulatory Requirements: Audit/Certification Criteria

Australia

Therapeutic Goods (Medical Devices) Regulations 2002
 - Schedule 3, Part 1 (excluding Part 1.6) – Full Quality Assurance Procedure

Canada

- Medical Device Regulations – Part 1- SOR 98/282

United States

- 21 CFR Part 803
 - 21 CFR Part 806
 - 21 CFR Part 807 – Subparts A to D
 - 21 CFR Part 820

Facility(ies):

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