



CERTIFICATE



This is to certify that the company

Black & Black Surgical, Inc.

5175 S. Royal Atlanta Drive
Tucker, GA, 30084
United States of America

has implemented and maintains a **Quality Management System**.

Scope of certification:

Design and development, manufacturing, service and distribution of liposuction aspirators, infiltration pump, reusable suction and irrigation cannulas, sterile suction and irrigation tubing and reusable surgical instruments, Monopolar/ Bipolar electrosurgical devices.

-CND, USA (a,b,c,d)

Through an audit, documented in a report, performed by DQS Medizinprodukte GmbH, it was verified that the management system fulfills the requirements of the following standard:

ISO 13485 : 2016

including applicable country-specific regulatory requirements, as indicated below the scope (full references of abbreviations are listed in the annex)

Certificate registration no.	515494 MDSAP16
Certificate unique ID	1000169029
Effective date	2024-05-19
Expiry date	2027-05-18
Frankfurt am Main	2024-05-15



DQS Medizinprodukte GmbH

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DQS Medizinprodukte GmbH is recognized under the Medical Devices Single Audit Program.

Visit <https://www.dqs.de/en/customer-database/> to validate this certificate.

The validity of this certificate can only be verified by the QR-code.



Annex to certificate
Certificate registration No.: 515494 MDSAP16
Certificate unique ID: 1000169029
Effective date: 2024-05-19

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Audited site

515494
Black & Black Surgical, Inc.
5175 S. Royal Atlanta Drive
Tucker, GA, 30084
United States of America

REPs FEI No.: site scope and country-specific requirements

Design and development, manufacturing,
service and distribution of liposuction
aspirators, infiltration pump, reusable suction
and irrigation cannulas, sterile suction and
irrigation tubing and reusable surgical
instruments, Monopolar/ Bipolar
electrosurgical devices.
-CND, USA (a,b,c,d)
REPs FEI No.: F000657



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Full references of country-specific requirements of MDSAP participating Regulatory Authorities

Abbreviation	Jurisdiction	Reference
AUS	Australia	(a) Therapeutic Goods (Medical Devices) Regulations 2002, Schedule 3, Part 1 – Full Quality Assurance Procedure (b) Therapeutic Goods (Medical Devices) Regulations 2002, Schedule 3, Part 4 – Production Quality Assurance Procedure
BRA	Brazil	RDC ANVISA n. 665/2022 RDC ANVISA n. 551/2021 RDC ANVISA n. 67/2009
CND	Canada	Medical Device Regulations SOR/98-282, Part 1
JPN	Japan	MHLW Ministerial Ordinance No. 169 (2004) as amended by MHLW Ordinance No. 128 (2014), Articles 4 to 68 Japan PMD Act (as applicable)
USA	United States	(a) 21 CFR Part 803 (b) 21 CFR Part 806 (c) 21 CFR Part 807 (d) 21 CFR Part 820 (e) 21 CFR Part 821