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POLISH CENTRE FOR
TESTING AND CERTIFICATION

CERTIFICATE

No. 1434-MDR-002/2025

of the EU technical documentation assessment with the requirements of Chapter II of Annex IX to the Regulation (EU) 2017/745 issued for medical devices manufactured by:

VUP Medical, a.s.
Šujanovo náměstí 302/3, 602 00 Brno, Czech Republic
Single Registration Number (SRN): CZ-MF-000042015

Polish Centre for Testing and Certification S.A. (Notified Body no. 1434) certifies that the technical documentation of the medical devices covered by this certificate complies with the requirements specified in Annexes II and III to the Regulation (EU) 2017/745. Details of the devices covered by this certificate are described on the following pages. The technical documentation report referenced below summarises the result of the assessment and includes reference to the relevant common specifications, harmonized standards and test reports.

Report no.: **S/711/BM/2022**

Validity of the Certificate: from 04.03.2025 to 03.03.2030
The date of issue of the Certificate: 04.03.2025

[redacted]

Certificate bears the qualified signature PCBC S.A.
Warsaw, 04.03.2025

Polish Centre for Testing and Certification 469, Puławska Street, PL-02-844 Warsaw tel.: +48 22 45 46 200, pcbc@pcbc.gov.pl

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No. 1434-MDR-002/2025

of the Assessment of the technical documentation with the requirements of Chapter II of Annex IX to the Regulation (EU) 2017/745

Devices covered by the certificate: Knitted vascular prosthesis with collagen Ra K

Types:

- **Ra lv K**
- **Ra bv K**

Risk Classification: III

Basic Code UDI-DI: 8591954RAKMV

Intended use: Knitted vascular prostheses impregnated with bovine collagen are suitable for vascular reconstruction wherever an application of an artificial vascular graft of the corresponding lumen is indicated.

Certificate validity conditions and/or limitations: N/A

Revision History including Changes: 1/04.03.2025 /S/711/BM/2022

Certificate bears the qualified signature PCBC S.A.
Warsaw, 04.03.2025

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