

CERTIFICATE

**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:**

Aflofarm Farmacja Polska Sp. z o.o.
ul. Partyzancka 133/151, 95-200 Pabianice
Single Registration Number (SRN): PL-MF-000001809

Report no.: **S/703/BM/2022**

The date of issue of the Certificate: 12.12.2024

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POLISH CENTRE FOR
TESTING AND CERTIFICATION

No. 1434-MDR-025/2024

**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: Calominal

Risk Classification: III

Basic Code UDI-DI: 590280270026AT

Intended use: treatment of overweight, body weight control, reduction of cholesterol absorption

Certificate validity conditions and/or limitations: **not applicable**

Revision History including Changes: 1 / 12.12.2024 / S/703/BM/2022

Certificate bears the qualified signature PCBC S.A.
Warsaw, 12/12/2024

FBM-R-24-MDR-E/7 of 27.09.2024

Polish Centre for Testing and Certification

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