

**KOPIA**

# CERTIFICATE

Management system as per  
**PN-EN ISO 13485:2012**  
Medical devices - Quality management systems - Requirements for regulatory purposes

In accordance with TÜV NORD Polska Sp. z o.o. procedures, it is hereby certified that

**Aflofarm Farmacja Polska Sp. z o.o.**  
**ul. Partyzancka 133/151, PL / 95-200 Pabianice**

applies a management system in line with the above standard for the following scope

**Design, manufacturing, storage and sales  
of osmotic hydrogels, emulsions, syrups, pessaries and tablets.**

Regardless of the fact that TÜV NORD Polska Sp. z o.o. is a notified body No. 2274 in the area of medical devices, this Certificate is not a Certificate of Conformity within the meaning of Directive 93/42/EEC and is not a basis for CE marking.

Certificate Registration No. **AC090 MD/1181/2720/2016**  
Audit Report No. PL2720/2017

Valid from **14-03-2017**  
Valid until **13-03-2020**  
Initial certification: **14-03-2014**

Manager of Certification Body  
TÜV NORD Polska Sp. z o.o.

Katowice, 03-03-2017

This certification was conducted in accordance with the TÜV NORD Polska Sp. z o.o. auditing and certification procedures and is subject to regular surveillance audits.

TÜV NORD Polska Sp. z o.o.

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www.tuv-nord.pl



AC 090  
QMS