

in accordance with § 23 of Law No. 56/2018 on Technical Requirements for Products and on Conformity Assessment and on amendments to certain laws, as amended

and with Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices, amending Directive 2001/83/EC, Regulation (EC) No 178/2002 and Regulation (EC) No 1223/2009 and repealing Council Directives 90/385/EEC and 93/42/EEC as amended (MDR).

UNIMED PHARMA, spol. s r.o., Orišková 11, 821 05 Bratislava, Slovak republic
single registration number (SRN): SK-MF-000003111

Medical device type:	Eye drops, solution
Trade name of medical device:	OCUflash® eye drops, solution
Variant:	1 × 10 ml; 2 × 10 ml
Basic UDI-DI:	8588009045OCUflash7K
Risk class of the Medical Device:	IIB, according to the Rule 21 of the EU Medical Device Regulation (MDR) 2017/745, Annex VIII
Intended purpose:	OCUflash eye drops, solution is intended to flush irritating particles, chemicals and foreign bodies from the eye, to flush the eye in case ocular surface injuries, and to drip into the eye when exposed to environmental factors (e.g. pollution, computer use, air conditioning) as prevention and complementary symptomatic treatment of non-infectious inflammation of the eyelids, conjunctiva and chronic conjunctiva infections, as well for their prevention.
Used procedure of conformity assessment:	Procedure of conformity assessment according to Regulation (EU) 2017/745 on medical devices, Annex IX Conformity assessment based on a quality management system and on assessment of technical documentation
Applied CS:	N/A
Name and number of Notified body (NB):	3EC International a.s. NB: 2265
Certificates issued:	Certificate: 2024-MDR/QS-022/A, revision 02 Issued: 11.12.2025 Expires: 22.02.2029

We issue the EU declaration of conformity under the sole responsibility of the manufacturer UNIMED PHARMA, spol. s r.o. and declare that the above-mentioned medical device is in accordance with the conformity assessment of the quality management system pursuant to Article 52 (4), the technical documentation pursuant to Annex IX and meets the general safety and performance requirements pursuant to Annex I of Regulation (EU) 2017/745 the European Parliament and of the Council of 5 April 2017 on medical devices and the device is in conformity with this Regulation. This medical device is marked with CE 2265.

The EU declaration of conformity was signed by the person responsible for regulatory compliance for and on behalf of the manufacturer UNIMED PHARMA, spol. s r.o.

In Bratislava, date: 2026-01-07
(Date and place of issue of the declaration)

Name and function of the responsible person
representing the manufacturer:

PharmDr. Vladimír Jančuš



Signature of the responsible person
representing the manufacturer
Company stamp