# ANNEX I TO EU QUALITY ASSURANCE CERTIFICATE
## No. 2024-MDR/QS-022

issued for the company

**UNIMED PHARMA spol. s r.o.**

Oriešková 11, 821 05 Bratislava, Slovak Republic

**List of medical devices covered by the EU Quality Assurance Certificate:**

| MD type | Trade names | Models / variants |
|---|---|---|
| OCUflash® eye drops, solution | **OCUflash®** | 1 x 10 ml |
| | **OCUflash®** | 2 x 10 ml |



Notified body
3EC international
NB 2265
NOTIFIKOVANÁ OSOBA

**Katarína Tomin Srdošová, PhD.**
*Director of NB2265*

In Bratislava, Slovakia, 22.02.2024
Valid until 22.02.2029

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## No. 2024-MDR/QS-022

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**Intended purpose of medical devices covered by the EU Quality Assurance Certificate:**

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 chronical conjunctiva infections.



Notified body 3EC international NB 2265 NOTIFIKOVANÁ OSOBA

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# ANNEX III TO EU QUALITY ASSURANCE CERTIFICATE
# No. 2024-MDR/QS-022

issued for the company

## UNIMED PHARMA spol. s r.o.

Oriešková 11, 821 05 Bratislava, Slovak Republic

**Certificate history:**

| Revision | EU QA Certificate reference | Date of issue | Application for Conformity Assessment of MD number | Description |
|---|---|---|---|---|
| 00 | 2024-MDR/QS-022 | 22.02.2024 | MDR287_2023 | Initially granted certification |



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