№ 000079

Ministry of Health
of the Republic of Uzbekistan

PHARM AGENCY
Agency on Development of
Pharmaceutical Industry

Center of good practices

GMP – GOOD MANUFACTURING PRACTICE

CERTIFICATE

No. GMP_UZ – 04:2025

Is issued on the basis of a completed pharmaceutical inspection conducted in accordance with the regulation on the procedure for conducting inspections for compliance with the requirements of good manufacturing practice (GMP).

STATE ENTITY
"CENTER OF GOOD PRACTICES" APPROVES

located at

str. Cetatea Alba 176, mun.Chisinau MD 2002, Republica Moldova

COMMERCIAL COMPANY
"FLUMED-FARM" LLC

Compliance with the requirements of
O'zDSt 2766:2018 –"Good Manufacturing Practice - GMP"

The basis for pharmaceutical inspection was application of **Commercial Company "FLUMED-FARM" LLC** dated **13th February, 2024** for pharmaceutical inspection in accordance with the requirements of O'zDSt 2766:2018 - "Good Manufacturing Practice-GMP".

GOOD MANUFACTURING PRACTICE — GMP CERTIFICATE APPENDIX

I. Sterile Products

1. Aseptically prepared (list of dosage forms):

☐ large volume liquids
☐ small volume liquids
☐ dispersions
☐ lyophilisates
☐ solids
☐ semi-solids
☐ other aseptically prepared products:

_____.

(the type of medicine or the type of activity is shown).

2. Medicines subject to sterilization at the end of production:

☐ large volume liquids
☐ small volume liquids
☐ solids and implants
☐ semi-solids
☐ lyophilisates
☐ other terminally sterilised prepared products:

_____.

(the type of medicine or the type of activity is shown).

II. Non-sterile products

☐ capsules, hard shell
☐ capsules, soft shell
☐ chewing gums
☐ impregnated matrices
√ **liquids for external use (nasal/oropharyngeal spray, drops, solution, emulsion)**
☐ liquids for internal use
☐ medicinal gases
☐ other solid dosage forms
☐ pressurised preparations
☐ radionuclide generators
☐ semi-solids
☐ suppositories
☐ tablets
☐ transdermal patches
☐ intraruminal devices
☐ other non-sterile medicinal product:

_____.

(the type of medicine or the type of activity is shown).

III. Biological medicinal products

☐ blood products
☐ immunobiological products
☐ cell therapy products
☐ gene therapy products
☐ tissue engineered products
☐ biotechnology products
☐ animal extracted products
☐ other biological medicinal products:

_____.

(the type of medicine or the type of activity is shown).

IV.Other products or manufacturing activity

☐ herbal products

☐ homoeopathic products

☐ other product

_____.

(the type of medicine or the type of activity is shown).

*Based on the information obtained during the pharmaceutical inspection conducted on **06-08.08.2024** and **17.01.2025 (online)** the applicant complies with the requirements of the Good Manufacturing Practice - GMP. The certificate is valid if all its pages (both main pages and additional pages) are presented. The validity of this certificate can be checked from the database of the State entity "Center of Good Practices". If the certificate is not provided in the indicated database, it is necessary to contact the working body that issued it.*

The GMP_UZ – 04:2025 Good Manufacturing Practice - GMP certificate
validity period from ***22.01.2025*** to ***21.01.2028***

Director
of the SE "Center of Good Practices" _____ (signature) **Dusmatov A.F.** (full name)

S.P.