



REPUBLIC OF LIBERIA

LIBERIA MEDICINES & HEALTH PRODUCTS REGULATORY AUTHORITY (LMHRA)

2nd & 3rd Floors, Clay Building, Sekou Toure Avenue, Mamba Point
1000 Monrovia, 10 Liberia – West Africa



Certificate No: 20-00142

**CERTIFICATE OF REGISTRATION OF
MEDICINES AND HEALTH PRODUCTS**

An Act to Establish the Liberia Medicines and Health Products Regulatory Authority
(LMHRA) of 2010 (Part V, Section 1 and 2)

This is to certify that

**FUNGUS GO TRIPPLE ACTION
(CLOTRIMAZOLE, USP 1%, BETAMETHASONE DIPROPIONATE 0.05% w/w
NEOMYCINULFATE USP 0.5%, CHLOROCRESOL USP1% 20gm CREAM)**

is registered with the LMHRA for use in Liberia and is hereby subject to the provisions of
the LMHRA Act of 2010.

Standard: USP Class: A

Applicant SUPER FORMULATION PVT. LTD.

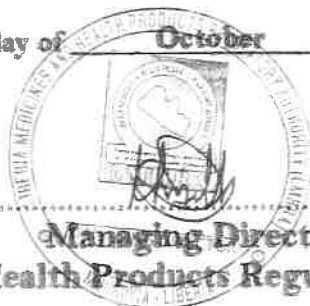
Manufacturer SUPER FORMULATION PVT. LTD.

Local Agent NAND PHARMACEUTICALS

Registration No: LMHRA/SM/20-00142 Date of Registration: 30 October 2020

The validity of this certificate shall continue until 29 October 2023 unless
otherwise suspended, revoked or varied as to the period of its validity.

Dated this 30th day of October, 2020



Managing Director

Liberia Medicines & Health Products Regulatory Authority (LMHRA)