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)