SAHPRA
SOUTH AFRICAN HEALTH PRODUCTS REGULATORY AUTHORITY

26/7/3/3/1/G0096/2020

CERTIFICATE OF GOOD MANUFACTURING PRACTICE (GMP)

I, the undersigned Medicines Control Officer of Medicines, South African Health Products Regulatory Authority of the Republic of South Africa, hereby certifies that the manufacturer of pharmaceutical products:

WRAPSA (PTY) LTD
C/O EDISON AND BELL CRESCENTS
HENNOPSPARK
CENTURION
PRETORIA
0046
REPUBLIC OF SOUTH AFRICA

has been authorised in accordance with section 22C (1)(b) of the Medicines and Related Substances Act, 1965 (Act 101 of 1965) under licence number 0000000180, to manufacture, test, pack, label, store and distribute tablets (Metronidazole, Ibuprofen, Ferrous Sulphate, Trimethoprim, Sulfamethoxazole, Paracetamol, Prednisone, Quinapril, Citalopram, Carvedilol, Cyproheptadine, Aspirin, Naproxen, Caffeine, Diazepam, Haloperidol, Metoclopramide, Magnesium Hydrochloride, Propranolol, Hydrochlorothiazide, TLD, TEE) capsules (Indomethacin, Oxytetracycline, Proxicam, Omeprazole), liquids, semi solids (Cream and Ointments), Aerosol, Colo Prep powders and complementary medicines.

From the knowledge gained during inspections of this manufacturer, it is considered that the company complies with the Good Manufacturing Practice requirements prescribed by South African Health Products Regulatory Authority. (S.A. Guide to GMP:www.sahpra.org.za)

ADDRESS OF CERTIFYING AUTHORITY
CHIEF EXECUTIVE OFFICER
SOUTH AFRICAN HEALTH PRODUCTS
REGULATORY AUTHORITY
CSIR CAMPUS
SAHPRA RECEPTION
BUILDING 38
MEIRING NAUDE DRIVE
BRUMMERIA
PRETORIA
0001

NAME OF AUTHORISED PERSON: Mr Bafana Malaza

TELEPHONE NUMBER: (012) 842-7544

SIGNATURE:

DEPT. OF HEALTH
REPUBLIC OF S.A.

ISSUE DATE: 29TH /07/2020

EXPIRY DATE: 20TH /07/2021