SAHPRA
SOUTH AFRICAN HEALTH PRODUCTS REGULATORY AUTHORITY

26/7/3/3/1/G0023/2022

CERTIFICATE OF GOOD MANUFACTURING PRACTICE (GMP)

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

PHARMACEUTICAL ENTERPRISES (PTY) LTD
49 MORNINGSIDE ROAD
NDABENI
CAPETOWN, 7405

Has been authorised in accordance to section 22C (1) (b) of the Medicines and Related Substances Act, 1965 (Act 101 of 1965) under licence number 0000000046MD to import, manufacture, test, pack, label and distribute Non-invasive Medical Device.

From the knowledge gained during inspection 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
LOFTUS PARK
BULDING A
2ND FLOOR
KIRKNESS STREET
ARCADIA
PRETORIA
0001

NAME OF AUTHORISED PERSON: Ms Naomi Pule TELEPHONE NO. 012 015 5442

SIGNATURE:

SAHPRA
SAHPRA South African Health Products Regulatory Authority
REPUBLIC OF SA

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