



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**

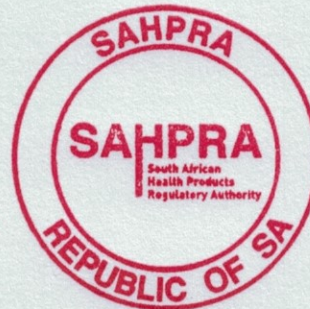
Has been authorised in accordance to section 22C (1) (b) of the Medicines and Related Substances Act, 1965 (Act 101 of 1965) under licence number 0000000046MD to import, manufacture, test, pack, label and distribute Non-invasive Medical Device.

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](http://www.sahpra.org.za) ).

**ADDRESS OF CERTIFYING AUTHORITY:  
CHIEF EXECUTIVE OFFICER  
SOUTH AFRICAN HEALTH PRODUCTS  
REGULATORY AUTHORITY  
LOFTUS PARK  
BUILDING A  
2<sup>ND</sup> FLOOR  
KIRKNESS STREET  
ARCADIA  
PRETORIA  
0001**

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

**SIGNATURE:**



**ISSUE DATE: 2022/02/11**

**EXPIRY DATE: 2023/02/11**