

SOUTH AFRICAN HEALTH PRODUCTS REGULATORY AUTHORITY



Licence number: 00004412MD

LICENCE TO DISTRIBUTE MEDICAL DEVICES

**In terms of section 22C(1)(b) of the Medicines and Related Substances Act, 1965
To act as a Distributor and Importer**

This licence is granted to:

Licence Holder

Mello Medical Supplies (Pty) Ltd

8 Century Road

Riversands Outlet Park, Knopjeslaagte

Midrand

1684

On the following terms and conditions:

The licence holder and the persons described and named in Annexure 1 shall at all times ensure that all medical devices distributed, irrespective of its registration status, comply with all the provisions of the Medicines and Related Substances Act, 1965, as amended and in particular with sections 14, 18, 18A, 18B, 18C, 19, 20, 22A, 22C, 22H, 23, 26, 28, 33 and the Regulations relating to Medical Devices 2, 3, 4, 5, 6, 13, 14, 17, 18, 19, 20, 21, 22, 23, 24, 25, 27, 28 and all relevant South African Health Products Regulatory Authority Guidelines.

This licence consists of 3 pages.

This facility is authorised to perform the activities listed in Annexure 1 to this licence.

Bathumelo Semele-Makokotela

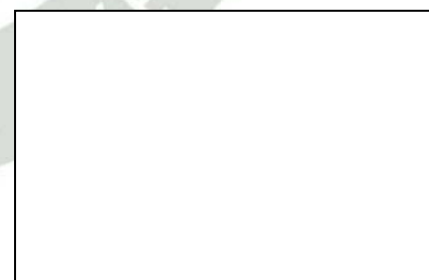
SAHPRA

CHIEF EXECUTIVE OFFICER

ORIGINAL DATE OF ISSUE: 25 February 2025

EXPIRY DATE: 25 February 2030

AMENDMENT DATE: N/A



This licence remains the property of the South African Health Products Regulatory Authority. Upon amendment, voluntary withdrawal, recall, suspension or revocation of the licence, the original licence must be returned to the Office of the Chief Executive Officer.

ANNEXURE 1

00004412MD

AUTHORISED DISTRIBUTION AND MATERIAL HANDLING ACTIVITIES

1. DISTRIBUTION ACTIVITIES	YES	NO
Distribution to hospitals and retail pharmacies and other clients: Class A	Yes	
Distribution to hospitals and retail pharmacies and other clients: Class B	Yes	
Distribution to hospitals and retail pharmacies and other clients: Class C		No
Distribution to hospitals and retail pharmacies and other clients: Class D		No
2. MATERIALS HANDLED OR STORED AT THIS SITE		
Combination medical devices with Penicillins	Yes	
Combination medical devices with Cephalosporins	Yes	
Combination medical devices with (other) Antibiotics (as specified):	Yes	
Combination medical devices with Hormones	Yes	
Combination medical devices with Cytostatics/Cytotoxics		No
Bulk Pesticides, Herbicides or Rodenticides		No
Radioactive material or Radioactive medical devices	Yes	
Other potent, toxic, sensitising or hazardous materials (as specified):		No
3. IMPORT		
Import Class A medical device	Yes	
Import Class B medical device	Yes	
Import Class C medical device		No
Import Class D medical device		No
Import Class A IVD	Yes	
Import Class B IVD	Yes	
Import Class C IVD		No
Import Class D IVD		No
Import RUO IVDs	Yes	
4. EXPORT		
Export Class A medical device		No
Export Class B medical device		No
Export Class C medical device		No
Export Class D medical device		No
Export Class A IVD		No
Export Class B IVD		No
Export Class C IVD		No
Export Class D IVD		No
Export RUO IVDs		No

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5. PARTICULARS OF THE PERSONNEL RESPONSIBLE FOR OPERATION ON THE PREMISES ON BEHALF OF THE LICENCE HOLDER

Authorised Representative	Manufacture / Import / Distribution / Export Control Person	Quality Control Person
Beverley Kedibone Zulu	Malesela Alphius Mello	Malesela Alphius Mello
Diploma: Business Administration	Diploma: Commercial Law	Diploma: Commercial Law

6. PARTICULARS OF THE LICENCE HOLDER CONTACT AND AUTHORISED REPRESENTATIVE (if not the same person)

Name	Contact Details	Address
Mr M.A. Mello (LH)	Tel: N/A Cell: N/A Fax: N/A Email: alfie.mello@etceteratec.co.za	8 Century Road Riversands Outlet Park Knopjeslaagte Midrand 1684
Miss B.K. Zulu (AR)	Tel: 120043268 Cell: 0727184248 Fax: N/A Email: zulubeverley025@gmail.com	8 Century Road Riversands Outlet Park Knopjeslaagte Midrand 1684

7. LICENCE SPECIFIC CONDITIONS

1. The holder of the licence shall conduct all manufacturing, distribution or wholesaling operations in respect of those medical devices for which a registration certificate has been obtained, so as to ensure that the medical devices shall conform to the standards of quality, safety and performance applicable to them in accordance with the specifications made by the person to whose order they are manufactured, distributed or wholesaled or the specifications under which the medical devices are sold or supplied.

8. ADDITIONAL LICENCE SPECIFIC CONDITIONS (IF REQUIRED)