

# SOUTH AFRICAN HEALTH PRODUCTS REGULATORY AUTHORITY



**Licence number: 00004537MD**

## LICENCE TO MANUFACTURE MEDICAL DEVICES

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

This licence is granted to:

Licence Holder

**Cryosave SA (Pty) Ltd**

Acacia House, Green Hill Village Office Park  
c/o Botterklapper and Nentabos Street, The Willows  
Gauteng  
0041

### 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 4 pages.**

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

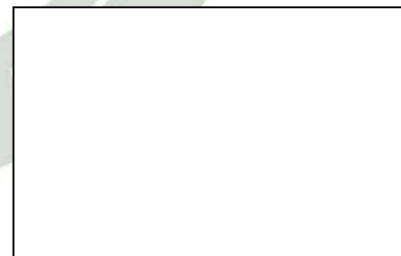
  
Boitumelo Senete-Makokotela

**CHIEF EXECUTIVE OFFICER**

**ORIGINAL DATE OF ISSUE: 11 April 2025**

**EXPIRY DATE: 11 April 2030**

**AMENDMENT DATE: N/A**



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**ANNEXURE 1**

**00004537MD**

**AUTHORISED MANUFACTURING AND MATERIAL HANDLING ACTIVITIES**

<b>1. MANUFACTURING ACTIVITIES</b>	<b>YES</b>	<b>NO</b>
<b>Sterile Medical Device Manufacture (includes primary packing, but not secondary packing such as cartoning or labelling)</b>		
Single use		No
Measuring medical devices		No
Non-invasive medical device		No
Invasive medical devices		No
Active medical devices	Yes	
Inactive medical devices		No
Contraceptive medical devices		No
Combination medical devices		No
Other sterile medical devices (as specified):		No
<b>Non-sterile Manufacture</b>		
Measuring medical devices		No
Non-invasive medical devices		No
Invasive medical devices		No
Active medical devices		No
Inactive medical devices		No
Contraceptive medical devices		No
Combination medical devices		No
Other non-sterile medical devices (as specified):		No
<b>Manufacture of In Vitro Devices (IVDs)</b>		
Class A IVD		No
Class B IVD		No
Class C IVD		No
Class D IVD		No
<b>End point Sterilisation of Medical Devices</b>	Yes	
<b>Manufacture of Radioactive Medical Devices</b>		No
<b>Servicing and Refurbishment of Medical Devices</b>		No
<b>2. PACKAGING ACTIVITIES</b>	<b>YES</b>	<b>NO</b>
Packaging of bulk product and labelling	Yes	
Re-labelling or redressing		No
Cartoning or secondary packaging	Yes	
Assembly or "kits" / procedure packs	Yes	
<b>3. TESTING ACTIVITIES</b>	<b>YES</b>	<b>NO</b>
Analytical	Yes	
Microbiological	Yes	
Sterility	Yes	
Stability	Yes	
Animal		No
Other Testing Activities (as specified):	Yes	
<b>4. DISTRIBUTION ACTIVITIES</b>	<b>YES</b>	<b>NO</b>
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	Yes	
Distribution to hospitals and retail pharmacies and other clients: Class D		No

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<b>5. MATERIALS HANDLED OR STORED AT THIS SITE</b>	<b>YES</b>	<b>NO</b>
Medical devices stored at licence holder site	Yes	
Combination medical devices with Penicillins		No
Combination medical devices with Cephalosporins		No
Combination medical devices with (other) Antibiotics (as specified):		No
Combination medical devices with Hormones		No
Combination medical devices with Cytostatics/Cytotoxics		No
Bulk Pesticides, Herbicides or Rodenticides		No
Radioactive material or Radioactive medical devices		No
Other potent, toxic, sensitising or hazardous materials (as specified):	Yes	
<b>6. IMPORT</b>		
	<b>YES</b>	<b>NO</b>
Import Class A medical device		No
Import Class B medical device		No
Import Class C medical device		No
Import Class D medical device	Yes	
Import Class A IVD		No
Import Class B IVD		No
Import Class C IVD		No
Import Class D IVD		No
Import RUO IVDs		No
<b>7. EXPORT</b>		
	<b>YES</b>	<b>NO</b>
Export Class A medical device		No
Export Class B medical device		No
Export Class C medical device		No
Export Class D medical device	Yes	
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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**8. 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
Lana du Plessis	Lana du Plessis	Petro Willemse
PhD: Human Genetics	PhD: Human Genetics	Certificate in Quality Management SABS

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

Name	Contact Details	Address
Mrs L Channon (LH)	Tel: N/A Cell: 082 656 6962 Fax: N/A Email: lizel@cryosave.co.za	Acacia House, Green Hill Village Office Park c/o Botterklapper and Nentabos Street, The Willows Gauteng 0041
Dr L du Plessis (AR)	Tel: N/A Cell: 0878080170 Fax: N/A Email: lana@cryosave.co.za	Acacia House, Green Hill Village Office Park c/o Botterklapper and Nentabos Street, The Willows Gauteng 0041

**10. LICENCE SPECIFIC CONDITIONS**

- 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.

**11. ADDITIONAL LICENCE SPECIFIC CONDITIONS (IF REQUIRED)**

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