SOUTH AFRICAN HEALTH PRODUCTS REGULATORY AUTHORITY

Licence number: 00005565MD

LICENCE TO WHOLESALE MEDICAL DEVICES

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

This licence is granted to:

Licence Holder
Ligwena Security
5 Fred Els Ave
Norkem Park
Kempton Park
Gauteng
1618

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 product 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.

Boitumelo Semete-Makokotlela

CHIEF EXECUTIVE OFFICER

ORIGINAL DATE OF ISSUE: 17 April 2026

EXPIRY DATE: 17 April 2031

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 **00005565MD**

AUTHORISED WHOLESALING AND MATERIAL HANDLING ACTIVITIES

1. SPECIFIC ACTIVITIES	**YES**	**NO**
Imported medical devices are stored and handled at this site.		No
Medical devices are stored on behalf of licenced local manufacturers or distributors of medical devices.		No
Medical devices are transmitted, consigned, conveyed or delivered from local manufacturers or distributors of medical devices.	Yes	
Custom made / "Special" manufactured medical devices are handled at this site.		No
Medical devices are exported from this site on behalf of manufacturers or distributors of medical devices.		No

2. METHOD OF ONWARD DISPATCH	**YES**	**NO**
Post	Yes	
Courier / Van Service	Yes	
Own Courier / Van Service	Yes	
Customer Collection	Yes	
Other (as specified):		No

3. CATEGORIES OF MEDICAL DEVICES HANDLED AT THIS SITE						
Class Medical Device / Type Medical Device		**A**	**B**	**C**	**D**	**RUO**
Sterile		X	X	X	X	-
Measuring Medical Device		X	X	X	X	-
Blood storage / transportation product		X	X	X	X	-
Non-Invasive		X	X	X	-	-
Invasive Medical Device	< 60 min	X	X	X	-	-
	>60 min & < 30 days	X	X	X	-	-
	>30 days	X	X	X	-	-
Inactive Medical Device		X	X	X	X	-
Active Medical Device		X	X	X	X	-
Contraceptive		-	X	X	X	-
Combination Medical Device		-	-	-	-	-
IVD		-	-	X	X	-

4. HOLDER OF AN ADDITIONAL SAHPRA LICENCE	**YES**	**NO**
Licence to manufacture / distribute medical devices (as specified):		No
Licence to manufacture / wholesale medicine (as specified):		No

5. SITE LISTING ON A SAHPRA LICENCE HELD BY ANOTHER LICENCE HOLDER	**YES**	**NO**
Medical Device Establishment Licence to Manufacture		No
Medical Device Establishment Licence to Distribute		No

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6. PARTICULARS OF THE PERSONNEL RESPONSIBLE FOR OPERATIONS ON THE PREMISES ON BEHALF OF THE LICENCE HOLDER

Authorised Representative
Pfariso Makhwathana
Diploma Financial Accounting

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

Name	Contact Details	Address
Mr P Makhwathana (LH/AR)	Tel: 015 962 2449 Cell: 081 818 6285 Fax: N/A Email: ligwenasec@gmail.com	5 Fred Els Ave Norkem Park Kempton Park Gauteng 1618

8. 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.

9. ADDITIONAL LICENCE SPECIFIC CONDITIONS (IF REQUIRED)

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