

# **SOUTH AFRICAN HEALTH PRODUCTS REGULATORY AUTHORITY**



DEPARTMENT OF HEALTH  
Republic of South Africa

**Licence number: 00000641MD**

## **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

**1 Stop Medical Supplies CC**

Unit 14, 997 Richards Drive

Halfway House, Midrand

Johannesburg

1685

### **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.**

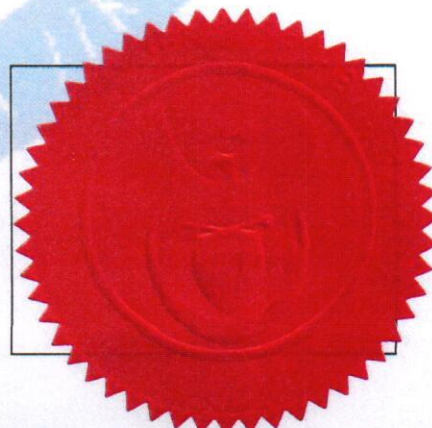
This facility is authorised to perform the manufacturing activities listed in Annexure 1 to this licence.

  
**ACTING CHIEF EXECUTIVE OFFICER**

**ORIGINAL DATE OF ISSUE: 11 May2018**

**EXPIRY DATE: 11 May 2023**

**AMENDMENT DATE: N/A**





## ANNEXURE 1

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<b>AUTHORISED MANUFACTURING AND MATERIAL HANDLING ACTIVITIES</b>
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1. MANUFACTURING ACTIVITIES	YES	NO
<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		No
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>		No
<b>Manufacture of Radioactive Medical Devices</b>		No
<b>Servicing and Refurbishment of Medical Devices</b>		No
<b>2. PACKAGING ACTIVITIES</b>		
Packaging of bulk product and labelling		No
Re-labelling or redressing	Yes	
Cartoning or secondary packaging	Yes	
Assembly or "kits" / procedure packs		No
<b>3. TESTING ACTIVITIES</b>		
Analytical		No
Microbiological		No
Sterility		No
Stability		No
Animal		No
Other Testing Activities (as specified):		No
<b>4. DISTRIBUTION ACTIVITIES</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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	YES	NO
<b>5. MATERIALS HANDLED OR STORED AT THIS SITE</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):		No
<b>6. IMPORT</b>		
Import Class A medical device	Yes	
Import Class B medical device	Yes	
Import Class C medical device	Yes	
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>		
Export Class A medical device	Yes	
Export Class B medical device	Yes	
Export Class C medical device	Yes	
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**

<b>Authorised Representative</b>	<b>Manufacture / Import / Distribution / Export Control Person</b>	<b>Quality Control Person</b>
Rajen Chetty	Rajen Chetty	Rajen Chetty
Diploma In Business Management	Diploma In Business Management	Diploma In Business Management

**9. PARTICULARS OF THE LICENCE HOLDER CONTACT (Other than the Authorised Representative)**

<b>Name</b>	<b>Contact Details</b>	<b>Address</b>
Mrs. F. Chetty	Tel: 011 312 9998 Cell: 076 053 8647 Fax: 086 224 9552 Email: fiona@1stopmedical.co.za	P O Box 50382 Randjesfontein Midrand 1683

**10. 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.

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

# **SOUTH AFRICAN HEALTH PRODUCTS REGULATORY AUTHORITY**

Republic of South Africa  
Private Bag X828  
PRETORIA  
0001



Republiek van Suid-Afrika  
Privaatsak X828  
PRETORIA  
0001

1 Stop Medical Supplies CC  
Unit 14, 997 Richards Drive  
Halfway House  
Midrand  
1685

Dear Sir/Madam,

## **LICENCE TO MANUFACTURE IN TERMS OF SECTION 22C(1)(b) OF THE MEDICINES AND RELATED SUBSTANCES ACT, 1965**

Licence Number      00000641MD

Your licence to manufacture in terms of section 22C(1)(b) of the Medicines and Related Substances Act has been approved and is attached herewith. This document replaces any licence document, for a medical device establishment, previously issued to you.

This licence authorises manufacture by the licence holder named; if the business should change hands, the company or person taking over the business will have to obtain a new licence before commencing the manufacture of medical devices.

This licence is subject to the limitations specified in the licence and to the statutory provisions contained in the Regulations to the Act.

Activities may only be carried out in accordance with the terms of the relevant product licence, unless a specified exemption applies, which allows it to take place other than in accordance with the licence.

This licence relates to the manufacture of medical devices on the premises and under the supervision of the persons specified. If any change of premises or of those persons takes place, prior approval must be sought from the South African Health Products Regulatory Authority. Any proposal to make structural alterations to the premises must also be notified to the South African Health Products Regulatory Authority.

The South African Health Products Regulatory Authority has power to revoke, suspend or amend licences in terms of section 22E.

Yours faithfully,



**ACTING CHIEF EXECUTIVE OFFICER**

Date: 11 May 2018

Faks/Fax: (012) 395 9201

Telefoon/Telephone: (012) 395 8126

Navrae/Enquiries: Ms. P. Nkambule