



MINISTRY OF HEALTH

PHARMACY AND POISONS BOARD

(Section 3B(2)(e) of the Pharmacy and Poisons Act, Cap 244 Laws of Kenya)

MEDICAL DEVICE REGISTRATION CERTIFICATE

This Registration Certificate is issued to

ALPHA MEDICAL MANUFACTURERS LTD

for distribution and sale of

HYPODERMIC SYRINGES AND HYPODERMIC NEEDLES.

Registration Number:	MD/2023/635
Certificate Valid Until:	13th April 2028
Registration Date:	14th April 2023
Device Category:	Class B
GMDN:	
GMDN Term:	
Intended Purpose: INJECTING MEDICINE INTO THE HUMAN BODY AND WITHDRAWING FLUIDS FROM THE BODY.	
 	

MAH Details:

WUXI YUSHOU MEDICAL APPLIANCES CO., LTD.

Manufacturing Sites :

WUXI YUSHOU MEDICAL APPLIANCES CO. LTD.,

Device Accessories:

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Device Group:

Device Sub-group/Sub-sets:

AUTO-DISABLE SYRINGE- ALL SIZES. HYPODERMIC SYRINGE WITH NEEDLE AND WITHOUT NEEDLE- ALL SIZES. HYPODERMIC DISPOSABLE NEEDLES-ALL SIZES. INSULIN SYRINGES- ALL SIZES. STERILE DISPOSABLE SYRINGES 2 PART LUER SLIPS- ALL SIZES

Conditions

The above Medical Device has been entered on the Record subject to the following conditions:

- The granted approvals herein are in accordance to the Laws of Kenya, Health Amendment Law 2019 and the applicant is required to adhere to the stipulated conditions.
- Each applicant/MAH/Product Owner shall retain records of the distribution of all of the applicant's medical devices included in the records for Medical Devices. In the case of records relating to a Class Active Implantable Medical Device (AIMD) medical device, Class C medical device, or Class B medical device that is an implantable medical device, the distribution records shall be retained for a minimum period of 10 years. In the case of records relating to any other device, the distribution records shall be retained for a minimum period of 5 years.
- The applicant of a medical device shall keep an up to date log of information of all the medical devices registered.
- It is a condition of inclusion in the PPB that the applicant of a medical device that is an AIMD, Class for implantable Class B provides three consecutive annual reports to the Medical Devices Department , Directorate of Product Evaluation following registration of the Medical Device. Reports should be for the period 1 July to 30 June. The first report following the date of inclusion in the PPB records must be for a period of at least six months but no longer than 18 months. The annual report must include all complaints and adverse events received by the manufacturer relating to problems with the use of the device that have been received by them over the year.
- A applicant shall ensure that a medical device within their control is stored and transported in accordance with the instructions and information provided by the manufacturer.

Products Covered by This Entry

Product Specific Conditions

Pharmacy and Poisons Board
Head Office, Lenana Road
Po Box 27663-00506
Nairobi, Kenya



Serial NO: **aba02c9fca11135caddb62ea32aa6adb**
Registration Date: **14-04-2023**