Sr. No. 3383

Government of Pakistan
Ministry of National Health Services
Regulation & Coordination

Drug Regulatory Authority of Pakistan

FORM-4

[see rule 5(2)]

LICENCE TO IMPORT MEDICAL DEVICES

Licence No. ELI-00966
Date of Issue: 13/10/2022
F.No: 03-141/2022-MD

M/s. Lifetek Traders (Pvt) Ltd, is hereby licensed to import registered medical devices at the following premises: Office No. B-8, Second Floor, Siraj Plaza (Gulzari Optics) Opp. Rashid Nursing Home Saidpur Road, Rawalpindi

2. Name(s) of proprietor(s) along with the residential address and CNIC Number(s)

Name	Address	CNIC
Mr. Abdul Rehman Khalid	House No. 5/587, Mohallah Gorah, Mandi Bahauddin	34402-4739495-9
Mr. Abdul Shakoor	Model Town, Humak Kahuta Road, House No. 926-A, Islamabad	61101-3300258-9
Mr. Abdul Rahim	House No.122-D, Street No.55, Sector G-6/4, Islamabad	34402-6025484-9

3. Name(s) of the person(s) incharge who will personally supervise the import and sale of medical devices by way of wholesale along with registration No, residential address and CNIC No.

Name	Address	CNIC
Mr. Abdul Rehman Khalid	House No. 5/587, Mohallah Gorah, Mandi Bahauddin	34402-4739495-9

4. Addresses of godowns , if any, where medical devices shall be stored

Rawalpindi Office No. B-8, Second Floor, Siraj Plaza (Gulzari Optics) Opp. Rashid Nursing Home Saidpur Road,

5. The licence will be in force for a period of five years from the date of issue unless earlier suspended or cancelled.

6. This licence shall, in addition to the conditions specified in the Medical Devices Rules, 2017 made under the DRAP Act, 2012, be subject to the following conditions namely:-

(i) The persons mentioned above shall personally supervise the sale of medical devices.

(ii) The licence and registration certificate from the Pharmacy council of the person(s) incharge, personally supervising the sale of medical devices shall be displayed in a prominent place in the premises open to public.

(iii) No medical device requiring special storage conditions of temperature and humidity shall be stored or sold unless the precaution necessary for preventing the properties of the components have been observed throughout the period during which it remained in possession of the licensee.

(iv) Importer shall be responsible for labeling requirements as per Medical Devices Rules, 2017 including Importer Licence details, Products Registration Numbers and MRP.

Renewal Date: 12/10/2027

BABAR KHAN
Additional Director (MD&MC) Secretary MDB
Drug Regulatory Authority of Pakistan
Ministry of National Health Services
Regulations & Coordination
Islamabad

13/10/2022