Government of Pakistan
Ministry of National Health Services
Regulations & Coordination
Drug Regulatory Authority of Pakistan

Sr. No. 388

FORM-4

[see rule 5(2)]

LICENCE TO IMPORT MEDICAL DEVICES

Licence No. ELI-00314
Date of issue: 25/02/2019
F.No: 12-71/2019-MD

M/s Bajwa Sons, is hereby licensed to import registered medical devices at the following premises: 13- Muslim Street, House No 3, 1st and 2nd Floor, Mayo Hospital Road, Lahore.

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

Name	Address	CNIC
Mr. Farhat Munawar Bajwa	House No. E-5/13-A, St. 6-A, Cavalry Ground Lahore Cantt	35201-2792838-5
Mr. Usman Ahmed Bajwa	House No. E-5/13-A, St. 6-A, Cavalry Ground Lahore Cantt	35201-2816947-1

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
Miss. Nimra Khalid	House No. 437-38, Nizam Block, Allama Iqbal Town, Lahore	35202-2273974-8

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

House No. 437-38, Nizam Block, Allama Iqbal Town, Lahore

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: 24/02/2024

(DR. GHAZANFAR ALI KHAN)
Additional Director (MD&MC)
Drug Regulatory Authority of Pakistan
M/o National Health Services,
Regulations and Coordination