

## No.F.10-2/2017-DD (H&OTC) (20<sup>th</sup> Meeting) DIVISION OF HEALTH & OTC PRODUCTS DRUG REGULATORY AUTHORITY OF PAKISTAN

## FORM-6

[(See rule 7(16)]

## PROVISIONAL CERTIFICATE FOR ENLISTMENT AS MANUFACTURER

## E. No. 00466

M/s Bio Serve Pharma, Plot No. 165, 1<sup>st</sup> Floor, S.I.T.E. Super Highway, Phase-II, Karachi (Nutraceuticals) is hereby enlisted in the enlistment register as manufacturer [Tablet, Capsule, Sachet, Semi-Solid & Liquid] by the Authority [approved in 20<sup>th</sup> Meet ing of EEC and vide F.No.1-939/2016-DD (H&OTC)] subject to the following conditions namely:-

- a. The manufacturer shall be responsible for the quality, efficacy and safety of all the therapeutic goods manufactured and sold by him.
- b. He shall abide by all the provisions of the Drug Regulatory Authority of Pakistan Act, 2012 (XXI of 2012) except those exempted under the said Act.
- c. He shall immediately recall the defected therapeutic goods within 15 days after intimation to him and report the compliance to the Authority.
- d. He shall be responsible to withdraw the unsafe therapeutic goods from the market if so declared by the Authority. He shall also report adverse affect reports, (if any), to the Authority within 15 days period and notify focal person for such reporting.
- e. Any other relevant condition imposed by the Authority in future.
- This certificate shall not be valid after the date notified by the Authority through official gazette.
- g. Certificate of enlistment shall be surrendered to the Authority within 7 days if it is suspended, revoked, becomes invalid or its holder winds up his business.

Abdul Sattar Sohrani

Deputy Director

Division of Health and OTC Products

Drug Regulatory Authority of Pakistan

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Dated: 26-05-2017

ADIL SAEED

Assistant Director (H & OTC)
Drug Regulatory Authority of Pakistan
Ministry of National Health Services
Regulations and Coordination
Islamabad