Requirements for the registration of pharmaceutical companies:

1) Company registration form (**Form A1**) to be filled in one copy signed and stamped by the person responsible on the establishment.

2) GMP certificate Legalized from one of the following authorities:

- FDA (USA), Food and Drug Administration.
- HPFB (Canada), Health products and Food Branch, therapeutic products directorate.
- EMEA (EU or any of its countries), European Union, European Agency for the Evaluation of Medical products.
- MHLW (Health authority of Japan), Ministry of Health, Labor, and Welfare, pharmaceutical and Medical Safety Bureau.
- TGA (Therapeutic Goods Administration) Australia.
- MHRA (Medicines and Healthcare Products Regulatory Agency) United kingdom
- Swiss Medic.
- GCC (Gulf Cooperation Council).

3) Copy of the CPP for at least two of its products released from one of the following authorities:

- FDA (USA), Food and Drug Administration.
- HPFB (Canada), Health products and Food Branch, therapeutic products directorate.
- EMEA (EU or any of its countries), European Union, European Agency for the Evaluation of Medical products.
- MHLW (Health authority of Japan), Ministry of Health, Labor, and Welfare, pharmaceutical and Medical Safety Bureau.
- TGA (Therapeutic Goods Administration) Australia.
- MHRA (Medicines and Healthcare Products Regulatory Agency) United kingdom
- Swiss Medic.
- GCC (Gulf Cooperation Council).

4) Manufacturing license certificate legalized from responsible authority and that mention all the pharmaceutical production lines.