Pharmaceutical Manufacturer & Healthcare Product Manufacturer Registration in KIMADIA Requirements

1- Letter of Authorization from Manufacturing Company to Scientific Drug Bureau in Iraq.

2- Filling of Registration Form "Appendix 3" with attachments. 👉 Click this link to download the form.

3- Israel Boycott Letter.

4- Manufacturing License.

5- Quality Certificate; **cGMP** from country of origin for Pharmaceuticals, **ISO 22000:2018** for Dietary Supplements, **ISO 13485:2016** for Medical Device and **ISO 22716:2007** for Cosmetic Products.

6- Product list & Catalogue of Products.

7- Site Master File & Site Master Plan.

8- Invitation Letter for Inspection Team visit.

9- Product Registration Certificate or CoPP or FSC for one product from each line of production mentioned in Quality Certificate issued from Country of Origin.

10- Incorporation Certificate (Establishment Certificate).

11- Registration Certificate for one product from each line of production from other countries.

Note; for further information / clarification / explanation on any point please feel free to contact us through our e-mail (info@daylapharm.com). We reply immediately through social messaging apps from our phone numbers (+964-771-854-6622) & (+964-751-541-8516), available on our website (www.daylapharm.com).