Medical Device Manufacturer Registration in KMCA Requirements

1- Request Letter from Agent.

2- Registration Fee payment receipt.

3- Letter of Authorization.

4- Filling of KMCA Form “211-REGD-MD.docx” with attachments. 👉 Click this link to download the form.

5- Quality Certificate “GMP from reference country or ISO 13485:2016 from country of origin”.

6- Manufacturing License.

7- Certificate of Establishment.

8- FSC for each line from country of origin.

9- Letter of Invitation.

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).