Medical Devices Registration in Bangladesh

We help you to prepare Medical Device Technical File (MDR) as per the regulatory requirement of EU MDR 745/2017 and submit to notified body for review and approval of CE certification. In Bangladesh, the Directorate General of Drug Administration (DGDA) oversees **medical devices registration**. Operating under the Ministry of Health and Family Welfare, DGDA regulates the import, production, distribution, and sale of medical devices, emphasizing quality and safety standards.