Two Days workshop on “Medical Devices Frameworks” organized by Danish Medical Agency (DKMA)- CDSCO on 8-9 April’ 2026 at CDSCO (HQ), New Delhi

A two-day training programme was conducted under the Strategic Sector Cooperation initiative between Denmark and India to strengthen collaboration in the medical devices and In-vitro diagnostics sector and enhance regulatory capacity building. The welcome address was delivered by Dr. Rajeev Singh Raghuvanshi, DCGI (CDSCO) who highlighted the importance of international collaboration and regulatory harmonization in advancing medical device safety and innovation. The workshop brought together experts from the Danish Medicines Agency (DKMA) and the Central Drugs Standard Control Organization (CDSCO) to exchange knowledge on EU legislation and the Indian legislation regarding Medical Devices, In-vitro diagnostics, medical device regulatory framework, best practices, and future cooperation opportunities. The training programme strengthened mutual understanding of EU and Indian regulatory systems, encouraged harmonization efforts, and laid the foundation for continued collaboration and capacity building in medical device and In-vitro Diagnostics regulation between Denmark and India.