Medical Device Product Regulations

**CDRH Learn** is the FDA’s Center for Devices and Radiological Health (CDRH) web page for multimedia industry education. It consists of more than 80 learning modules describing many aspects of medical device and radiation emitting product regulations, covering both premarket and post-market topics. The presenters are FDA /CDRH staff.

This tool is intended to provide industry with information that is comprehensive, interactive, and easily accessible. Modules are provided in various formats, including videos, audio recordings, and slide presentations.

Categories of topics include:
- The Basics: Overview of Regulatory Requirements for Medical Devices
- How to Study and Market A Device
- Postmarket Activities
- Unique Device Identification (UDI) System
- Specialty Technical Topics
- Radiation-Emitting Products
- In Vitro Diagnostics (ICD)
- Industry Basics Workshop Presentations

**References:**
CDRH Learn website:  [http://www.fda.gov/Training/CDRHLearn/](http://www.fda.gov/Training/CDRHLearn/)