Final Rules of Unique Device Identification

Date : Feb 14, 2018 - 10:00 AM

Event URL : http://www.BostonEventsList.com/events/final-rules-of-unique-device-identification-feb-2018

Organizer : Netzealous LLC DBA - Compliance4All

Venue : Online Event

Location : 161 Mission Falls Lane, Suite 216,, Mission Falls Ln
Fremont, California, USA, ZIP: 94539
Phone: 8004479407

Ticket Price: One Dial-in One Attendee Price: $150.00
Overview:
This webinar will address the four key steps to compliance by device manufacturers. Also covered will be the Final Rule's provisions to address existing FG inventory, not properly labeled.

Why should you Attend:
It will also review the implementation schedule which is required of medical device companies selling products in the U.S. This is a major change for medical device manufacturers, with far-reaching effects in regulatory compliance, as envisioned by the FDA.

Areas Covered in the Session:
Learn the basic requirements of UDI Labeling and its Database
UDI / GUDID Implementation Schedules
Required steps for UDI / GUDID compliance by the Medical Device Company
Future Requirements

Who Will Benefit:
Process
Validations
CGMP Responsibilities
CROs and Clinicals Personnel

Medical Personnel
Other Healthcare Professionals
Staff and Office Personnel

Speaker Profile:
John E. Lincoln is a medical device and regulatory affairs consultant. He has helped companies to implement or modify their GMP systems and procedures, product risk management, U.S. FDA responses. In addition, he has successfully designed, written and run all types of process, equipment and software qualifications/validations, which have passed FDA audit or submission scrutiny, and described in peer-reviewed technical articles, and workshops, world wide.

Event Fee: One Dial-in One Attendee Price: US$150.00

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Event Categories :