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Food and Drug ...

**FDA Regulation of
Medical Devices
(Part 1 of 3)** (Part 1 of
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3) General overview of **medical device regulation.**

FDA 101 for Medical Devices Registrar Corp's webinar provides industry with important information regarding U.S. **FDA regulation of medical devices, ...**

How to work in Regulatory Affairs (Drug and Medical Devices) ► Webpage:

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For those that want to work on a Regulatory Affairs department, the path ...

**4.2 Regulatory
Affairs Basics -
Medical Devices**

***The 5 most relevant
changes the Medical
Device Regulation
MDR introduces,
that you must know***

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The Medical Device Regulation MDR replaces both, the Medical Device Directive (MDD, 93/42/EEC) and the Directive for Active ...

***Classification
Medical Device in EU
(Medical Device
Regulation MDR
2017/745)*** Medical
Device Regulation
Training - Within less
than 10 minutes you'll
learn how to classify a

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**Medical Device News
- March 2020**

**Regulatory Review
(EU MDR) ►** Webpage:
[https://podcast.easyme
dicaldevice.com/70/](https://podcast.easymedicaldevice.com/70/)

March 2020 is really a
period with a lot of
information to provide
for ...

**What Is a Medical
Device? (New**

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(2017/745)** Medical
Device Regulation
Training - What is a
Medical Device in
Europe?

The new Medical
Device Regulation EU
MDR 2017/745 ...

***Transitioning from
the Medical Device
Directives (MDD) to
the Medical Device
Regulation (MDR)***

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The move from the
Medical Device
Directive (MDD) to the
Medical Device
Regulation (MDR) is
one of the most
daunting ...

***What is ISO 13485
for medical devices?***

A brief introduction to
this ISO Standard for
medical devices. ISO
13485:2016.

***Regulatory
Standards & Risk***

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**Management in
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Standards
and Risk Management
in Medical Devices**

The webinar highlights
the speaker's unique
career paths to ...

**Best ISO 13485:2016
Starter Video [For
Medical Devices]**

Easy Medical Device - <https://easymedicaldevice.com> is a blog to learn
about the Medical
Device Regulations and

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Standards ...
Regulatory Affairs

***What is Post
Marketing***

***Surveillance for
Medical Devices?
(MDR 2017/745) ►***

Webpage: <https://podcast.easymedicaldevice.com/41>

Her nickname is the
PMS Queen as she can
tell you all you need to
know ...

Surviving a
Page 14/28

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Regulatory Interview

This program is about interviewing techniques.

Specifically, the program will show the incorrect and correct ways to handle ...

**Introduction to
Medical Device
Labeling Symbols** To

thrive in a global market place, it is crucial to communicate important product information in an

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understandable format.
It's also...

The 5 most important steps to CE certification - The EU medical device approval process

This video introduces
the European medical
device regulations, in
particular the Medical
Device Regulation
MDR, the conformity ...

Overview of the USA FDA Classification

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Process Classification is arguably one of the most important steps in the US **FDA medical device** approval process. Understanding how the ...

What is a European Authorized Representative (EU MDR 2017/745) What is a European Authorized Representative or EC Rep or CE Representative? (EU

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This ...

***FDA Regulations and
Medical Device
Pathways to Market***

Russ King, President of
Method Sense,
provides a high level
overview of **FDA
regulations** as part of
the
commercialization ...

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**Medical Device
Regulatory Update -
February 2020**

On this Regulatory Update, we will talk about many diverse topics but here is an outline.

First, we will review the ...

**Medical Device
Regulation
Transition Timeline
(MDR 2017/745)**

Medical Device
Regulation Training:

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Learn about the hard
and the soft transition
for your Medical
Devices within the MDR
2017 ...

***Medical Device News
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shaking the Medical
Device World*** In this
episode of the Medical
Device made Easy
Podcast, we will
review;
- How Coronavirus is
shaking the Medical
Device World ...

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***[Webinar] Preparing
for the New EU
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Regulation***

The
upcoming European
Union (EU) **Medical
Device Regulation**
(MDR) will replace the
Medical Devices
Directive and the
Active ...

***#Cybersecurity for
Medical Devices with
Erik Vollebregt (MDR
& IVDR)*** Let's see if
what is shown in

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movies is correct. Can
some hackers take
control of a Hospital or
kill you by hacking an
insulin ...

***How to prepare your
MDSAP certification?
(Medical Devices) ►***

Podcast Webpage: <http://podcast.easymedicaldevice.com/10/>

On today's show, I
invited Angelina Hakim
the CEO of Qunique ...

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**EU Medical Device
Regulation Webinar**

Claigan Environmental
Inc. will present a
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restricted materials
requirements in the EU
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