

## How to Comply with both in the Same Organization

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**Date :** Feb 16, 2018 - 10:00 AM

**Event URL :** <http://www.BostonEventsList.com/events/how-to-comply-with-both-in-the-same-organization-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

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**Ticket Price: One Dial-in One Attendee Price: \$150.00**

Overview:

This ISO 13485:2016 webinar is a standard for Quality Management Systems of medical device manufacturers and suppliers, and is used worldwide for developing and maintaining the system that caters to the needs of the market requirements for medical devices.

Why should you Attend:

For people in the medical devices industry which either develops or planning to market medical devices to USA and Europe and plan to implant a QMS that complies with both regulatory authorities and ask themselves how to do it correct and efficiently.

Areas Covered in the Session:

Introduction

Brief review of regulations and standard: status and history

ISO 13485:2016 short review

Principals

Quality management system

Management responsibility

Resource management

Product realization

Measurement, analysis and improvement

**Who Will Benefit:**

Medical Devices Quality Personal (QA and QC)

Laboratory Managers

R&D Researchers of Medical Devices and Combined Products

Quality Control Staff

Regulatory Affairs (RA) Staff

**Speaker Profile:**

Eyal Lerner is owner of ELC Consulting Services which offers the pharmaceutical and medical devices industries support in all quality related issues: Preparation for inspections, CAPA and RCA, quality manual QSM, self-inspection, compliance gap analysis, 3rd party audits, Management Review, GMP, GLP, MD: ISO 13485:2016 adjustments, DHF, DMR, V&V, 510k submission preparation and ISO 17025 and Validation.

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

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**Event Categories :**