

2-Day In-Person Seminar by Ex-FDA Official **FDA Recalls - Before You Start, and After You Finish**

By: **Casper (Cap) Uldriks**, Former Associate Center Director of FDA's CDRH

Location: Mebane, NC | May 16-17, 2019



SPEAKERS

Casper (Cap) Uldriks, Former Associate Center Director of FDA's CDRH

Casper (Cap) Uldriks, through his firm "Encore Insight LLC," brings over 32 years of experience from the FDA. He specialized in the FDA's medical device program as a field investigator, served as a senior manager in the Office of Compliance and an Associate Center Director for the Center for Devices and Radiological Health. He developed enforcement actions and participated in the implementation of new statutory requirements. His comments are candid, straightforward and of practical value. He understands how FDA thinks, how it operates and where it is headed. Based on his exceptionally broad experience and knowledge, he can synthesize FDA's domestic and international operational programs, institutional policy and thicket of legal variables into a coherent picture.

TOPIC BACKGROUND

The products regulated by the FDA can cause serious adverse health consequences or death. The FDA's recall program is designed to make sure firms' recalls can mitigate such problems, even when the adverse consequence seems remote. The FDA's recall program has remained relatively the same over several years. How the FDA and industry manage recalls and learn from their mistakes continues to evolve. In some ways recalls have become more efficient through the benefit of technology. In other ways the reason for recalls remains substantially the same. The FDA's recall procedures and regulatory management of any risk to health are still sufficient to determine whether a recall is effective and whether manufacturers learn from their mistakes.

LEARNING OBJECTIVES

- ✓ Understand FDA's recall authority and policy
- ✓ Learn how to manage recalls under FDA oversight
- ✓ Learn how to interact with FDA
- ✓ See how to develop health risk determinations
- ✓ Learn critical recall strategy components
- ✓ Manage possible FDA enforcement actions

COURSE DESCRIPTION

FDA's recall authority and program launches you into a project of crisis management. You will learn how to establish a roadmap for conducting recalls. The knowledge you gain will sharpen your recall management decisions and strategy. You will learn how to use the FDA's health risk criteria so you can develop effective recall procedures. One critical aspect of recalls involves the identification of the root cause of the recall and how you could or should prevent that problem from happening again. Your corrective and preventive action program (CAPA) and quality assurance functions require a rigorous approach to prevent a chronic history of recalls. Reiterative recalls lead the FDA to the conclusion that, "You don't get it."

Your compliance competency becomes a regulatory issue for the FDA if your recall is deemed ineffective. The seminar will cover critical performance targets for conducting an effective recall. You will learn how missteps in the recall process become an expensive problem in terms of money and a sloppy corporate image.

You will take away practical knowledge on how to work with FDA staff during a recall, and how you can prepare for inspectional follow up or a regulatory action, and in some cases a legal action. You will learn that your approach to recalls plays a major role in mitigating direct and indirect damage to your firm's business. A firm with a history of chronic recalls needs to learn how to get out of that downward spiral. Likewise, for established and new firms you will learn how you can reduce the negative impact of a recall with the use of proper planning.

AGENDA

Day One (8:30 AM - 4:30 PM)	
<p>08.30 AM - 09.00 AM: Registration</p> <p>09.00 AM: Session Start</p> <p>Day 1 - Morning</p> <p>FDA's Regulatory Authority</p> <ul style="list-style-type: none"> ✓ Recall Regulations <ul style="list-style-type: none"> ▶ Voluntary recall: 21 Code of Federal Regulations (C.F.R.) Part 7 ▶ Mandatory recall actions <ul style="list-style-type: none"> ▶ 21 C.F.R. Part 810 ▶ 21 C.F.R. Part 806 ✓ Recall Classification <ul style="list-style-type: none"> ▶ Violation of the law <p>Break (10:30 AM – 10:45 AM)</p> <ul style="list-style-type: none"> ✓ Risk to Health ✓ Precedents ✓ Exemptions <ul style="list-style-type: none"> ▶ Stock Recovery ▶ Product Withdrawal ▶ Product Improvement <p>Lunch (12:00 PM to 1:00 PM)</p> <p>Day 1 / Afternoon</p> <p>Recalls and risk to health</p> <ul style="list-style-type: none"> ✓ Risk to health categories <ul style="list-style-type: none"> ▶ Death ▶ Serious injury / serious illness ▶ Non-reversible / reversible ▶ May cause, if it were to recur ▶ Remote possibility 	<p>Break (2:30 PM – 2:45 PM)</p> <ul style="list-style-type: none"> ✓ Health Hazard Evaluation for Recall Classification <ul style="list-style-type: none"> ▶ FDA's internal evaluation ▶ Vulnerable subpopulations ▶ Scoring ▶ Participants ▶ Industry HHE equivalent ▶ FDA's recall database
Day Two (8:30 AM - 4:30 PM)	
<p>Day 2 - Morning</p> <p>FDA's Recall Procedures</p> <ul style="list-style-type: none"> ✓ Understanding FDA's program and implementation ✓ FDA's agency-wide recall procedures ✓ The FDA's investigator's job ✓ Preparing a recall strategy <p>Break (10:30 AM – 10:45 AM)</p> <ul style="list-style-type: none"> ✓ Preparing for FDA oversight ✓ Recall notification to FDA's District Office ✓ Recall notification to the public <p>Lunch (12:00 PM to 1:00 PM)</p> <p>Day 2 / Afternoon</p> <ul style="list-style-type: none"> ✓ Root cause identification ✓ Correction and Prevent Action (CAPA) <p>Break (2:30 PM – 2:45 PM)</p> <ul style="list-style-type: none"> ✓ FDA inspectional follow up ✓ Enforcement: FDA administrative and legal remedies ✓ End 	

WHO WILL BENEFIT

- ✓ Recall managers
- ✓ Quality assurance managers
- ✓ Regulatory affairs directors
- ✓ Risk and product liability managers
- ✓ Manufacturers' sales and marketing managers
- ✓ Own label distributors

Companies and departments:

- ✓ Manufacturers
- ✓ Own Label Distributors
- ✓ Importers
- ✓ Healthcare institutions
- ✓ Nursing homes
- ✓ Medical practice groups

..... **Registration Form**

Registration Information:

- ✓ **Register Online.** Use your American Express, Visa or MasterCard.
- ✓ Get your group to attend the seminar at a discounted price call +1-646-416-9016.
- ✓ Your Registration for the seminar is subject to following terms and conditions. If you need any clarification before registering for this seminar please call us +1-646-416-9016 or email us at info@salesmist.com

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Written cancellations through fax or email (from the person who has registered for this conference) received at least 10 calendar days prior to the start date of the event will receive a refund – less a \$300 administration fee. No cancellations will be accepted – nor refunds issued – within 10 calendar days from the start date of the event. On request by email (before the seminar) a credit for the amount paid minus administration fees (\$300) will be transferred to any future SalesMist event and a credit note will be issued. Substitutions may be made at any time. No-shows will be charged the full amount. We discourage onsite registrations, however if you wish to register onsite payment to happen through credit card immediately or check to be submitted onsite. Conference material will be given on the spot if it is available after distributing to other attendees. In case it is not available we will send the material after the conference is over. In the event SalesMist cancels the seminar, SalesMist is not responsible for any airfare, hotel, other costs or losses incurred by registrants. Some topics and speakers may be subject to change without notice.

Seminar Topic: FDA's Medical Device Software Regulation Strategy
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Date & Location:

Attendee Details:

	Name	Title	Email
Attendee 1			
Attendee 2			
Attendee 3			
Attendee 4			

Email address (so you can receive order acknowledgements, updated news, product information and special offers)

Company Information

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Address

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City

State Zip.....

Country

Phone Fax

Payment Options

Charge to: Visa MasterCard American Express

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