

CDER SMALL BUSINESS AND INDUSTRY ASSISTANCE (SBIA)

# ELECTRONIC DRUG REGISTRATION AND LISTING (eDRLS)

Using CDER Direct 2023



SEPT 28

VIA WEBCAST | [www.fda.gov/CDERSBIA](http://www.fda.gov/CDERSBIA)

Version 6, August 28, 2023  
(use link below to check for updates)

For files and resources, please visit

[The Event Page on SBIAevents.com](https://www.fda.gov/sbiaevents)

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## AGENDA

All times are Eastern (UTC-5)

[View Start Time on World Clock](#)

Thursday, September 28, 2023

8:45 - 9:00

### Welcome and Overview

**Brenda Stodart, PharmD, BCGP, RAC-US**

*Captain, United States Public Health Service  
Director, Small Business, and Industry Assistance (SBIA)  
Division of Drug Information (DDI) | Office of Communications (OCOMM)  
Center for Drug Evaluation and Research (CDER) | FDA*

9:00 – 9:15

### Keynote

**Jill Furman, JD**

*Director  
Office of Compliance (OC)  
CDER | FDA*

### Your SBIA Host

**Forest "Ray" Ford, Jr., PharmD, BCPS**

*CAPT, USPHS, Pharmacist  
SBIA | DDI | OCOMM | CDER | FDA*

Thursday, September 28, 2023

**9:15 – 10:00**

**Registering Your Drug Manufacturing Establishment Using CDER Direct**

Topics include:

- CDER Direct Establishment Registration Demo
- Using the Appropriate Business Operation(s) and Business Qualifier(s)
- Registration Renewals, Updates, and Deregistration
- US Agents and Official Contacts

**Regie Samuel**

*Technical Information Specialist*  
 Drug Registration and Listing Branch (DRLB)  
 Division of Labeling, Registration and Unapproved Drugs (DLRUD)  
 Office of Unapproved Drugs and Labeling Compliance (OUDLC)  
 Office of Compliance (OC)  
 CDER | FDA

**Jose Cabrera**

*Information Technology Specialist*  
 DRLB | DLRUD | OUDLC | OC | CDER | FDA

**10:00 – 10:45**

**Requesting a Labeler Code from FDA**

Topics include:

- CDER Direct Labeler Code Request Demo
- Who Will be Assigned a Labeler Code and Who Won't
- Labeler Code Inactivation and Reactivation
- Updates, Mergers and Acquisitions

**Soo Jin Park**

*LCDR, USPHS*  
*Regulatory Officer*  
 DRLB | DLRUD | OUDLC | OC | CDER | FDA

**Laurie Simonds, GWCPM**

*Technical Information Specialist*  
 DRLB | DLRUD | OUDLC | OC | CDER | FDA

**Lalnunpuii Huber**

*Technical Information Specialist*  
 DRLB | DLRUD | OUDLC | OC | CDER | FDA

**10:45 – 11:00: BREAK**

**11:00 – 12:00**

**Listing Your Drug Using CDER Direct**

Topics include demonstrations of:

- CDER Direct Drug Listing Demo
- Listing a Combination Product
- Strength Conversion in Drug Listing
- Listing Updates and Delisting
- Blanket No Change Certification

**Troy Cu**

*Technical Information Specialist*  
 DRLB | DLRUD | OUDLC | OC | CDER | FDA

**Leyla Rahjou-Esfandiary**

*Lead Consumer Safety Officer*  
 DRLB | DLRUD | OUDLC | OC | CDER | FDA

**Yogesh Paruthi**

*Consumer Safety Officer*  
 DRLB | DLRUD | OUDLC | OC | CDER | FDA

**Vikas Arora**

*Pharmacist*  
 DRLB | DLRUD | OUDLC | OC | CDER | FDA

Thursday, September 28, 2023

12:00 – 12:15

**503B Registration and Product Reporting Using CDER Direct**

Topics include:

- Demo
- Updates

**Huascar Batista**

*Senior Advisor*

Office of Compounding Quality and Compliance (OCQO)  
Office of Compliance (OC) | CDER | FDA

12:15 – 12:45

**Q&A Panel**

**All Speakers**

12:45 – 1:40: LUNCH BREAK

1:40 – 1:45

**SBIA Welcome Back & CE Reminders**

**Forest "Ray" Ford, Jr., PharmD, BCPS**

*CAPT, USPHS, Pharmacist*  
SBIA | DDI | OCOMM | CDER

1:45 – 2:00

**OMUFA Updates**

**Yajun (Jason) Tu, PharmD, PhD, BCSCP**

*LCDR, USPHS*

Program Management Officer  
Policy and Operations Branch (POB)  
Division of User Fee Management (DUFM)  
Office of Management (OM) | CDER | US FDA

2:00 – 2:30

**National Drug Code**

Topics include:

- NDC Reservation
- Future Format of the National Drug Code
- NDC Assignment to Drugs

**David Mazyck**

*Consumer Safety Officer*

DRLB | DLRUD | OUDLC | OC | CDER | FDA

**Julian Chun**

*Pharmacist*

DRLB | DLRUD | OUDLC | OC | CDER | FDA

**Soo Jin Park**

*LCDR, USPHS*

*Regulatory Officer*

DRLB | DLRUD | OUDLC | OC | CDER | FDA

Thursday, September 28, 2023

2:30 – 3:15

**Registration and Listing Compliance Program**

Topics include:

- Untitled Letters and Warning Letters
- Data Inactivation
- Data Removals and Flags
- Downstream Effects

**Tasneem Hussain**

*Pharmacist*

DRLB | DLRUD | OUDLC | OC | CDER | FDA

**Vikas Arora**

*Pharmacist*

DRLB | DLRUD | OUDLC | OC | CDER | FDA

**Leyla Rahjou-Esfandiary**

*Lead Consumer Safety Officer*

DRLB | DLRUD | OUDLC | OC | CDER | FDA

3:15 – 3:30: BREAK

3:30 – 3:45

**Recent Automated Validation Rules**

**Lalnunpuii Huber**

*Technical Information Specialist*

DRLB | DLRUD | OUDLC | OC | CDER | FDA

3:45 – 4:15

**Case Studies**

**Julian Chun**

*Pharmacist*

DRLB | DLRUD | OUDLC | OC | CDER | FDA

4:15 – 4:40

**Q&A Panel**

**All Speakers**

4:40 – 4:45

**SBIA Closing**

**Forest "Ray" Ford, Jr., PharmD, BCPS**

*CAPT, USPHS, Pharmacist*

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4:45 - ADJOURN