

Downstream Effects

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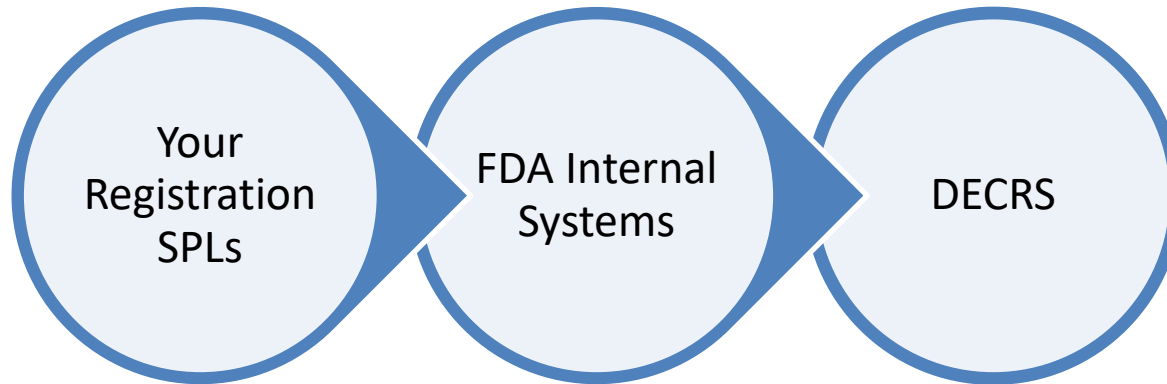
CDER Direct Workshop – September 28, 2023

Learning Objectives

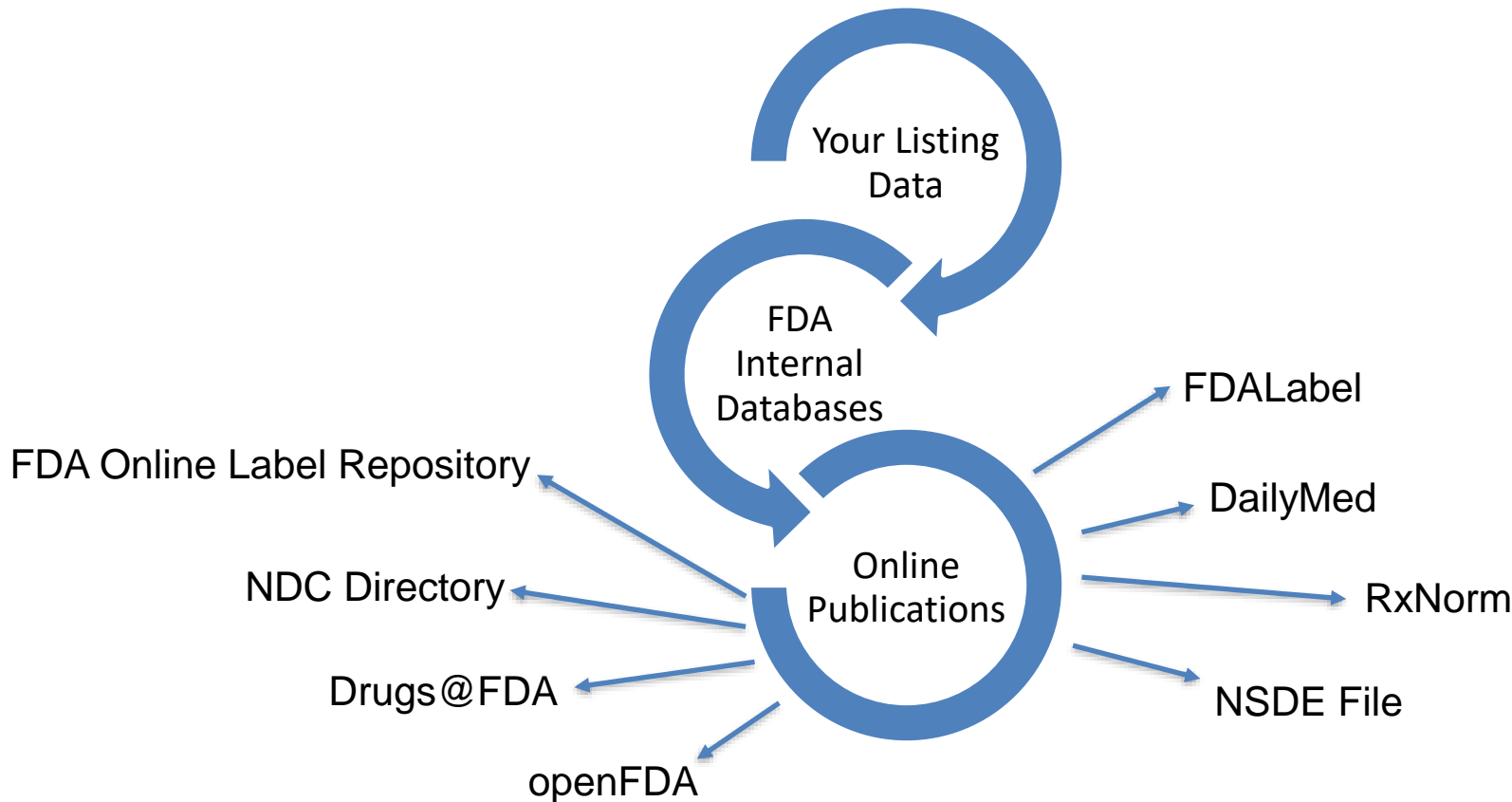
- Identify online publications
- Describe ways to report data discrepancies to FDA
- List potential real-life examples caused by data deficiency

Registration Data

- Published on [Drug Establishments Current Registration Site | FDA](#)



Listing Data





Who Uses the Data

Inside FDA

- Facility inspection planning
- Statutory facility fee assessments
- Drug shortage initiatives
- Drug amount reporting
- Safety assessments

Outside FDA

- Other government agencies
- Pharmaceutical industry
- Drug databanks
- Academia and research
- Healthcare providers and patients

Why Data Integrity is Important

- Reliability of CDER's site selection model and product catalog

| Location | CY 2017 | CY 2018 | CY 2019 | CY 2020 | CY 2021 | CY 2022 |
|----------|---------|---------|---------|---------|---------|---------|
| Domestic | 3,479 | 3,297 | 3,139 | 879 | 1,311 | 2,061 |
| Foreign | 1,457 | 1,321 | 1,200 | 204 | 57 | 381 |
| Total | 4,936 | 4,618 | 4,339 | 1,083 | 1,368 | 2,442 |

CGMP Inspections of Registered Domestic and Foreign Drug and Device Establishments

[Source: Annual Report on Inspections of Establishments CY 2022](#)

Why Data Integrity is Important

- Inaccurate listing or NDC assignment can lead to medication error
 - A drug is co-packaged with two injectable vials
 - Drug A and Drug B have the same NDC
 - Patient was injected with the same drug twice



Why Data Integrity is Important

- Healthcare providers might miss key labeling information
 - A repackager drug listing was not updated with a boxed warning after the source drug's labeling update



Why Data Integrity is Important



- Incorrect packaging information can lead to reimbursement or reporting mistakes

Comprehensive NDC SPL Data Elements File (NSDE)

[NSDE | FDA](#)



CDER NextGen

Important Safety Initiatives



- We play a key role in preventing incidents related to substandard, falsified, and harmful drugs:
 - Diethylene glycol/ethylene glycol contamination [WHO urges action to protect children from contaminated medicines](#)
 - FDA ophthalmic drugs initiative [FDA Issues Warning Letters to Firms Marketing Unapproved Eye Products | FDA](#)



Poll Question

It is in the best interest of the pharmaceutical company, as well as public health, to submit correct registration and listing data to FDA.

- A. True
- B. False



Find Data Discrepancy in FDA Online Publications?

- Contact FDA:
 - Drug Registration and Listing Branch: eDRLS@fda.hhs.gov
 - Division of Drug Information: DrugInfo@fda.hhs.gov

Challenge Question #1



Which one of these publications is managed by FDA?

- A. Comprehensive NDC SPL Data Elements File
- B. Drug Establishment Current Registration Site
- C. Neither
- D. Both



Challenge Question #2

Which of the following statements is NOT true?

- A. Anyone can contact FDA about data discrepancy found in its published registration and listing data.
- B. FDA can make corrections to its online published registration and listing data.
- C. FDA's listing data is used by other government agencies for coding, reimbursement, prescribing, and dispensing of drugs.
- D. NDC assignment is not part of drug approval but is an important safety aspect of a drug.



U.S. FOOD & DRUG
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Questions?

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