

## **Practitioner Diversion Awareness Conference**

#### Inventories, Records and Reports Lynnette Wingert, Unit Chief Policy Unit







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## **Course Objectives**

**s** Discuss who is responsible for maintaining controlled substance records.

**§** General recordkeeping requirements.

- **§** Basic inventory requirements.
- **s** Required records of receipt and distribution.

**s** Determine when and what reports are required to be submitted.



## **Questions To Discuss**

At the completion of this block of instruction you will be able to answer the following questions:

- 1. Practitioners are required to keep records of all controlled substances on hand, to include samples such as Lyrica®?
- 2. Who is ultimately responsible for keeping records of controlled substances at the registered location?



## **Questions To Discuss**

- **3.** If a practitioner writes controlled substance prescriptions, using their special "x" identifier for maintenance and detoxification they must keep a record of the prescription information?
- 4. How many controlled substances can a practitioner distribute in a calendar year to other dispensers before the practitioner must register as a distributor?
- 5. If a practitioner discovers a theft or significant loss of controlled substances, when must the theft or loss be reported?







## Who Must Keep Records

- S A practitioner who handles controlled substances, other than prescriptions. 21 CFR § 1304.03(b)
  - Except EPCS prescriptions. 21 CFR § 1304.03(c)
- S A practitioner who prescribes an FDA approved CIII-V narcotic controlled substance for opioid maintenance and/or detoxification. 21 CFR § 1304.03(c) & (d)



#### Responsible Party 21 CFR § 1304.03(a)

The DEA registrant is the person who is responsible for keeping controlled substance records.

s Not your nurse
s Not your office manager
s Not your corporation
s Not your vendor
s Not your employer



## **EPCS PRESCRIBERS**

A practitioner who issues electronic prescriptions for controlled substances (EPCS) must use an electronic prescription application that retains the following information:



- The digitally signed record of the information specified in <u>21 C.F.R. Part 1306</u>.
   21 C.F.R. 1304.06(a)(1)
- The internal audit trail and any auditable event identified by the internal audit as required by 21 C.F.R. § 1311.150.
   21 C.F.R. 1304.06(a)(2)



- S An institutional practitioner must retain a record of identity proofing and issuance of the two-factor authentication credential, where applicable, as required by 21 C.F.R. § 1311.110. 21 C.F.R. 1304.06(b)
- Solution Security Security incident Security Filed with the Administration pursuant to 21 C.F.R § 1311.150 and § 1311.215. 21 C.F.R. § 1304.06(d)



- S An electronic prescription or pharmacy application provider must retain third party audit or certification reports as required by 21 C.F.R § 1311.300. 21 C.F.R. § 1304.06(e)
- An application provider must retain a copy of any notification to the Administration regarding an adverse audit or certification report filed with the Administration on problems identified by the third-party audit or certification as required by 21 C.F.R. § 1311.300. 21 C.F.R. 1304.06(f)



#### S Unless otherwise specified, records and reports must be retained for two years. 21 C.F.R. § 1304.06(g)



## Maintenance and Detox Prescribers

- § Prescription records are required for prescribers who issue controlled substance prescriptions for maintenance or detoxification.
   21 C.F.R. § 1304.03(c)
- s Records of prescription information must be maintained separate from all other required records and readily retrievable. 21 C.F.R. § 1304.04(g)



## Maintenance and Detox Prescribers

§ Practitioners who dispense controlled substances for maintenance and/or detoxification must maintain the same records as any other dispensing practitioner. <u>21 C.F.R. § 1304.22(c)</u>



# General Recordkeeping



## General Record Keeping Requirements

**Requirements that apply to all controlled substance records required to be kept:** 

- **s** Must be complete and accurate. <u>21 C.F.R. § 1304.21(a)</u>
- **§** Must be stored at the registered location. <u>21 C.F.R. § 1304.21(b)</u>
- **s** Must be kept for two years. <u>21 C.F.R. § 1304.04(a)</u>



## General Record Keeping Requirements

- S Must be readily retrievable. 21 C.F.R. § 1304.04(f)(2)
- s Records must be kept for each separate DEA registered activity. <u>21 C.F.R. § 1304.21(c)</u>
- **§** Must be kept for each DEA registered location. 21 C.F.R. § 1304.21(b)



## Inventories



## **Inventory Requirements**

#### s Is a "Physical Count"

§ Must include all controlled substances "On Hand" (In possession/under the control of). (21 CFR §1304.11(a)



### **Inventory Requirements**

## **s** Inventory date must reflect the date of the actual inventory.

§ Maintained in Written, Typewritten, or Printed Form at the Registered Location. 21 C.F.R. § 1304.11(a)



## **Separate Inventories**

#### s Separate inventories are required for each registered location. 21 C.F.R. § 1304.11(a)

§ Must be taken at the Beginning of Business (BOB) or Close of Business (COB). 21 C.F.R. § 1304.11(a)

Separate inventories for each independent activity. <u>21 C.F.R. § 1304.11(a)</u>



### **Initial Inventories**

§ Inventory of all stocks of controlled substances.

Son the date you first engage in the manufacture, distribution, or dispensing of controlled substances.

s Best if labeled "Initial Inventory."

s If nothing on hand record "0."



## **Biennial Inventories**

#### S The biennial inventory is required to be taken on any date within two years of a previous required inventory.

s Best if labeled "Biennial Inventory."



## Newly Scheduled Controlled Substances

Solution System 4 Sector 2 Sector 2

s Must be taken at the Beginning of Business or Close of Business.



## Records



#### Schedule II controlled substance records shall be maintained separately from all other records.



- s Records of schedules III-V controlled substances must be kept separate from all other records or readily retrievable.
- § Records that are readily retrievable can be separated out in a reasonable time. <u>21 C.F.R. § 1300.03</u>



#### § Some examples of ways to render your records readily retrievable include but not limited to: <u>21 C.F.R. § 1300.01</u>

- s Items asterisk
- s Redlined
- **s** Or in some manner which sets them visually apart.



## **DEA Form 222**

- S The DEA Form 222 is used for the acquisition and distribution of schedule II controlled substances.
- **s** The DEA Form 222 must be filled out completely and accurately.
- **s** Power of Attorney authorizing who may execute a DEA Form 222.



## **Power Of Attorney**

Solution of the DEA application or renewal is the individual authorized to execute DEA Form 222's.

§ All others authorized who wish to execute a DEA Form 222 must first obtain a Power of Attorney from the signer.



#### § Must immediately inventory all schedule III-V controlled substances when received.

s Annotate the date received on the record of receipt.

## **Dispensing Log/Patient File**

Dispensing of controlled substances can be entered in patient chart or can be maintain in a dispensing log with the below information:

- s Actual Name of Controlled Substance, Form, Quantity, Strength;
- s Number of Units or Volume of Finished Form Dispensed;
- s Name, Address of the Person to Whom It Was Dispensed;
- s Date of Dispensing.



## **Transferring Controlled Substances**

## What to do if you need to transfer controlled substances to another DEA Registrant.

**§** Must use a DEA Form 222 (CII). <u>21 CFR 1307.11(a)(1)(iii)</u>

§ Must use a sales invoice for (CIII-CV). 21 CFR 1307.11(a)(1)(ii)



## **Transferring Controlled Substances**

#### § 5% of your yearly total. <u>21 CFR 1307.11(a)(1)(iv)</u>

§ If more you must register as a distributor. 21 CFR 1307.11(b)



## Reports

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## **Theft and Loss**

- s Theft, or Significant Loss.
- s Not an Inventory Adjustment.
- s Loss (Unexplained Disappearance).

S Any discovered shortage which the practitioner cannot convincingly establish to have been diverted after reasonable review/investigation should generally be considered a loss.



Solution Solution States St

s Must complete a DEA form 106, online, once your investigation is complete.
<u>21 C.F.R. § 1301.76(b)</u>

s Registrants are encouraged to immediately report theft and losses to your local law enforcement and state regulatory agency.

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- **s** DEA Form 41 shall be used to record the destruction of all controlled substances using an on-site method that renders the controlled substances non-retrievable.
- § DEA Form 41 shall include the names and signatures of the two employees who witnessed the destruction. 21 CFR § 1317.95(d)



#### S The DEA Form 41 no longer needs to be sent to the Division Office, maintain with your other records.



**Exceptions for DEA Form 41:** 

§ Destruction of a controlled substance dispensed by a practitioner for immediate administration at the practitioner's registered location, when the substance is not fully exhausted (i.e. wastage) shall be properly recorded in accordance with § 1304.22(c), and such record need not be maintained on a Form 41. 21 C.F.R. § 1304.21(e)



#### **Exceptions for DEA Form 41:**

§ Transfers by registrant to a reverse distributor must be recorded in accordance with § <u>1304.22(c)</u>, and such record need not be maintained on a Form 41. <u>21 C.F.R. § 1304.22(e)</u>





- S Registrants are required to provide effective controls and procedures to guard against theft and diversion of controlled substances. <u>21 C.F.R. § 1301.71(a)</u>
- S Registrant cannot employ anyone who has a felony drug conviction who will have access to controlled substance, without a DEA approved employment waiver. 21 C.F.R. § 1301.76(a)





#### § Practitioners are required to store stocks of CII-CV controlled substances in a securely locked, substantially constructed cabinet. <u>21 C.F.R. 1301.75(b)</u>



## **State Regulations**

- s Also consult your state regulating agency for more strict recordkeeping requirements.
- s Example some state boards require records be kept for 7 years.
- **§** Stricter Law Provision.



1. Practitioners are required to keep records of all controlled substances on hand, to include samples such as Lyrica®?

A. TrueB. False

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- 2. Who is ultimately responsible for keeping records of controlled substances at the registered location?
  - A. Licensed Practical Nurse
    B. Office Manager
    C. DEA Registrant
    D. Corporation



**3.** If a practitioner writes controlled substance prescriptions, using their special "x" identifier for maintenance and detoxification they must keep a record of the prescription information?

A. TrueB. False



4. How many controlled substances can a practitioner distribute in a calendar year to other dispensers before the practitioner must register as a distributor?

A. 40%
B. 20%
C. 60%
D. 5%



- 5. If a practitioner discovers a theft or significant loss of controlled substances, when must the theft or loss be reported?
  - A. On a DEA Form 106 upon completion of the investigation of the theft or loss.
  - **B.** In writing to DEA within 1 business day of discovery of the theft and loss.
  - C. DEA must be notified upon completion of the local police departments investigations.
  - **D. DEA** is not required to be notified.



# Thank-you for your time and attention!



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