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SUBCHAPTER 7.1 - RECALLS

7.1.1 - DEFINITIONS

7.1.1.1 - Recall

A Recall is a firm's removal or correction of a marketed product that FDA considers to be in violation of the laws it administers, and against which the Agency would initiate legal action (e.g., seizure). Recall does not include a market withdrawal or a stock recovery. See the Agency recall policy outlined in [21 CFR 7.1/7.59- Enforcement Policy - General Provisions, Recalls \(Including Product Corrections\) - Guidance on Policy, Procedures and Industry Responsibilities](#).

7.1.1.2 - Recall Classification

Recall Classification is the numerical designation, i.e., I, II, or III, assigned by the FDA to a particular product recall to indicate the relative degree of health hazard presented by the product being recalled.

7.1.1.2.1 - CLASS I RECALL

Class I Recall is a situation in which there is a reasonable probability that the use of, or exposure to, a violative product will cause serious adverse health consequences or death.

7.1.1.2.2 - CLASS II RECALL

Class II Recall is a situation in which use of, or exposure to, a violative product may cause temporary or medically reversible adverse health consequences or where the probability of serious adverse health consequences is remote.

7.1.1.2.3 - CLASS III RECALL

Class III Recall is a situation in which use of, or exposure to, a violative product is not likely to cause adverse health consequences.

7.1.1.3 - Recall Type

Recall type is a designation based on whether the recall is Voluntary, FDA Requested (at the request of the Commissioner or his/her designee), or ordered under [section 518\(e\) of the FD & C Act](#) [21 U.S.C 360h (e)].

7.1.1.4 - Recall Strategy

Recall strategy is a planned specific course of action to be taken in conducting a specific recall, which addresses the depth of recall, need for public warnings, and extent of effectiveness checks for the recall.

7.1.1.5 - Depth of Recall

Depending on the product's degree of hazard and extent of distribution, the recall strategy will specify the level in the distribution chain to which the recall is to extend, i.e., wholesaler, retailer, user/consumer, which is known as the

depth of recall.

7.1.1.6 - Recall Number

The recall number is the number assigned, by a responsible Center, for each recalled product they initiate. This number consists first of a letter designating the responsible Center (see letter Codes below), a 3 or 4 digit sequential number indicating the number of recalls initiated by that Center during the fiscal year, and a 4-digit number (the Center for Devices and Radiological Health (CDRH) uses 2-digit numbers) indicating the fiscal year the recall was initiated. For example: F-100-2011 identifies the 100th recall initiated by the Center for Food Safety and Applied Nutrition (CFSAN) in FY-2011. The following letters are used to identify the Centers.

Letter Center/Office

F	Foods - CFSAN
D	Drugs - Center for Drug Evaluation and Research (CDER)
Z	Medical Devices & Radiological Health - CDRH
V	Veterinary Medicine - Center for Veterinary Medicine (CVM)
B	Biologics - Center for Biologics Evaluation and Research (CBER)
N	Medical Devices (Voluntary Safety Alerts and Notifications)

T Tobacco Products – Center for Tobacco Products

7.1.1.7 - Medical Device Notification Order

A medical device notification order is an order issued by FDA requiring notification under [section 518\(a\) of the FD & C Act](#) [21 U.S.C. 360h (a)]. The directive issues when FDA determines a device in commercial distribution, and intended for human use, presents an unreasonable risk of substantial harm to the public health. The notification is necessary to eliminate the unreasonable risk of such harm, and no more practicable means is available under the provisions of the Act to eliminate such risk.

7.1.1.8 - Medical Device Notification

A medical device notification is a communication issued by the manufacturer, distributor, or other responsible person in compliance with a Notification Order. It notifies health professionals and other appropriate persons of an unreasonable risk of substantial harm to the public health presented by a device in commercial distribution.

7.1.1.9 - Medical Device Safety Alert

A medical device safety alert is a communication voluntarily issued by a manufacturer, distributor, or other responsible person (including FDA). It informs health professionals and other

appropriate persons of a situation which may present an unreasonable risk to the public health by a device in commercial distribution.

NOTE: Medical Device Notifications and Safety Alerts as described in IOM 7.1.1.7, 7.1.1.8, and 7.1.1.9 are to be handled by the Districts as recalls. They will go through the stages of alert, recommendation, classification, field notification, firm notification letter, firm effectiveness checks and status reports, FDA audit checks and termination recommendations.

7.1.1.10 Sub-Recall

A sub-recall is an action taken by a recalling firm's account to notify their own accounts/consignees of the recall, as long as no changes were made to the recalled product.

If the recalling firm's account changed the recalled product (for example, if they used the product as a component of a new product, re-labeled the product to obscure the original product name and/or lot code, repackaged the product, etc.) the account has created a new product which could warrant a new recall, instead of a sub-recall.

7.1.1.11 Consignee

Anyone who received, purchased, or used the product being recalled.

7.1.1.12 Account

Location at which you are conducting the audit check.

SUBCHAPTER 7.2 - RECALL NOTIFICATION / INSPECTION

If FDA learns of a potentially violative product which may lead/has led to a class I or significant class II recall, an inspection should be made to determine the root cause(s) of the problem(s). If the firm has failed to take appropriate corrective and preventive action, violations should be documented for possible regulatory action.

NOTE: In all discussions of violative or potentially violative products with the responsible firm, make it clear FDA is not requesting recall action. FDA requested recalls are authorized only by ORA, or by delegation of authority such as Drug Efficacy Study Implementation (DESI) recall requests.

When an investigation determines there is no evidence of manufacturing or distribution problems, but a firm has removed products from the market as a result of actual or alleged tampering with individual units, the action will be considered a Market Withdrawal. In addition, product that has been reported by a firm to be stolen and is being removed by the market by the firm will also be considered a market withdrawal. A market withdrawal means a firm's removal or correction of a distributed product which involves a minor violation that would not be subject to legal action by the FDA or which involves no violation, e.g., normal stock rotation practices, routine equipment adjustments, repairs, theft, etc.

7.2.1 - INSPECTION PROCEDURES

An important part of your job is to identify the root cause for the recall and assure the firm has implemented procedures to prevent it from recurring. In some cases, management will have conducted its own analysis and reached conclusions about the problem and its cause. The initial judgments about the problem are not always correct nor discriminating enough to identify the underlying causes. You need to verify the steps taken were sufficient in depth and scope and reflect the correct conclusions about both the problem and correction.

Determine if the firm conducted a failure analysis using techniques such as fault tree analysis or failure mode analyses. Did it consider things such as the length of time the product has been manufactured and sold, complaints or returns for the same or similar problems, any reworking of product prior to release or distribution which may have been due to the same or similar problems and, process or personnel changes which occurred about the time the problem appeared?

For all recall inspections, in addition to verifying the identification of the root cause:

1. Issue a Notice of Inspection (FDA 482);
2. Discuss the suspected problem with management and review the firm's complaint file;
3. Investigate all areas, control points and/or circumstances which may have a bearing on the product's deficiency;
4. Fully develop individual responsibility for the problem;
5. Review batch records, processing logs and/or other types of records for violative lots and associated lots;
6. Review and obtain copies of the firm's quality control/analytical data;
7. Determine any actions the firm has taken, is taking, or has planned to take to prevent similar occurrences. If corrective action is not underway, determine the firm's timetable for achieving correction; and
8. Determine what action the firm has taken or plans to take, and the time frames involved, regarding questionable product(s) remaining in commerce.

7.2.1.1 - Recall Decision Follow-up

If the firm has decided to recall, do the following:

1. Request that management obtain their FDA District's review of recall correspondence and any press releases before they are issued to prevent misunderstandings between the firm, its customers, and the FDA. This suggestion is voluntary on the part of the firm and is not required;
2. If the firm requests guidance in preparing recall communications, provide it in accordance with your District policy. See [Chapter 7 of the RPM](#) and IOM Exhibit 7-1 for an example of recall communications;
3. See [RPM Chapter 7-10, Attachment B "Recommendation for Recall Classification"](#) and [21 CFR 7.46a \(1\)-\(9\)](#) for information to be obtained;
4. Obtain an Official Sample of the recalled product. (See IOM 7.2.6 for the collection of samples for electronic products or medical devices.);
5. Obtain a complete distribution list of all shipments of the suspect lot(s), including foreign distribution;
6. Obtain specimens or copies of all labels and labeling associated with the recalled product;
7. Obtain complete copies of all recall communications issued or planned including the text of phone conversations, and submit them to your District's recall coordinator. Look in the Blue Pages for a list of District Recall Coordinators;
8. Advise the firm on how the returned products should be handled. FDA must witness or otherwise verify the reconditioning or destruction of the products returned under the recall; and
9. Take any other steps necessary in your judgment, or that your District requires.

NOTE: At this early stage there usually has not been a recall evaluation by the appropriate Center. In the absence of such an evaluation, avoid suggesting the firm extend its recall efforts.

7.2.2 - FOOD RECALLS

Experience with food recalls dictates specific information be obtained from firms which have used recalled material in the production of another product. This is necessary to decide if the recall must be extended to a new product(s). In those instances, the following are some areas to be covered:

1. Incoming ingredient quality control procedures;
2. Quality control over ingredients at the time of use, and the products in which the ingredients are used;
3. A detailed description of the methods used in preparation and packaging of the processed product;
4. How the finished product is stored and shipped;
5. Labeling of product, and any cooking instructions for consumer or purchaser;
6. Quality control testing of the finished product. Detail any test(s) performed by firm; and
7. For products produced in USDA plants, determine if the USDA was notified of the suspect incoming ingredient? Did USDA determine what testing was done by the firm?

This information must be evaluated by CFSAN (HFS-607) prior to the initiation of any sub-recall.

7.2.2.1 - Interstate Milk Shippers

The FDA will not ordinarily be involved in the classification and auditing of Interstate Milk Shippers (IMS) product recalls where such actions have been, or are being, handled expeditiously and appropriately by the State(s). However, the FDA District office in which the recalling firm is located must be assured that all States involved in an IMS plant's recall are participating in ensuring removal of the product from commerce and that, when appropriate, States issue warnings to protect the public health.

In the event that FDA determines that the States are unable to effect the recall actions necessary, the Agency will classify, publish, and audit the recall, including issuance of a public warning when indicated.

7.2.3 - MEDICAL DEVICE RECALLS

Medical device recalls may result from manufacturing defects, labeling deficiencies, failure to meet premarketing requirements [PMA, 510(k)], packaging defects or other nonconformance problems. How firms identify the causes of medical device recalls and corrective action activities is essential to the analysis of medical device failures and the determination of the effectiveness of the medical device GMP program. It is also useful in evaluating the medical device program, and for directing attention to problem areas during inspections. [21 CFR Part 806.1](#) requires device manufacturers and importers to report certain actions concerning device corrections and removals. They must also maintain records of all corrections and removals regardless of whether such corrections and removals are required to be reported to FDA. (See [21 CFR Part 806.20](#)). Failure to report as required by [21 CFR 806.10](#) is a violation and should be listed on the FDA-483, "Inspectional Observations." "Center concurrence is required prior to issuing warning letters for suspected violations of the user reporting regulations; to include Corrections or Removals regulations"

Each device manufacturer or importer must submit a written report to FDA of any correction or removal of a device initiated by such manufacturer or importer, if one was initiated:

1. To reduce a risk to health posed by the device; or
2. To remedy a violation of the Act caused by the device which may present a risk to health, unless the information has been provided according to [21 CFR 806.10](#) (f), or the correction or removal action is exempt from the reporting requirements under [21 CFR 806.1](#) (b).

Collection of complaint, PMA and 510(k) related information is necessary to determine compliance with the GMP requirements. During recall follow-up inspections, answers should be obtained to the questions below, in addition to routine recall information. For firms where it has been established a manufacturing defect led to the recall, conduct a complete GMP evaluation of the

manufacturing operations. Report such inspections into FACTS as "qualifying" GMP inspections.

7.2.3.1 - Problem Identification

1. How did the firm identify the nonconformance which led to the recall (e.g., complaint, in-house data, etc.)?
2. If the recall was due to a device defect, did the firm conduct a documented failure analysis of the device, using such techniques as fault tree or failure mode analyses? If so, report whether these results were provided for review.
 - a. Did the firm determine the failure mechanism (e.g., shorted component, incomplete weld, etc.)?
 - b. If not, how did firm determine the cause of the nonconformance?
 - c. If not, what rationale does the firm have for not conducting a failure analysis?
3. Did the firm determine at what phase of the device life cycle the nonconformance occurred (i.e., design, manufacturing, storage, use, etc.) and the actual cause of the nonconformance (i.e., software design error, process out of specifications, employee error, user misuse, etc.)? What evidence does the firm have to support the determination?
4. Did the firm determine if the nonconformance resulted in an injury or death?
5. If a component was responsible for the defect, determine if the same component was used in other devices manufactured by the firm. If so, has the firm conducted an analysis to assure the defect in the component will not have a deleterious effect on the operation of the other device(s)?
6. If a component was responsible for the device defect, what other device manufacturers use the same component (and especially the same lot number of the component)? Has the manufacturer of the recalled device notified the component manufacturer? Has the component manufacturer contacted its other customers about the problem?
7. Why was the component defective? Did the manufacturer of the component change the specifications without notifying the finished device manufacturer? Did the component fail to meet its release specifications?

NOTE: A visit to the component manufacturer may be needed to adequately answer questions 5, 6 and 7. Before doing so, confirm with CDRH and your supervisor that the matter is egregious enough to warrant this "next step."
8. Did the finished device manufacturer have an incoming component/raw material sampling and testing procedure? If not, why not?
9. If the manufacturer recalled the device because the labeling was inaccurate, or the wrong labeling was applied to the device (label mix-up) determine the following:
 - a. What quality system procedures should have been established to prevent the problem?
 - b. If the label or instructions for use were inaccurate, was the inaccuracy introduced in the design stage, or was it due to a printing problem?

10. If the device has been on the market for a year or more, and the manufacturer claims the problem is the result of design:
- Why is the problem just now showing up? How many reports concerning the problem did the firm receive before deciding a recall was necessary? Does the firm have a procedure established for determining if a recall is necessary, and if so, did it follow the procedure? Obtain a copy of the procedure.
 - If the firm doesn't provide rational answers to the above questions, determine if they explored other possible causes for the problem.
 - Was the design feature which caused the problem included in the design of the device that was the subject of a premarket submission?
 - If the design feature which caused the problem is part of the original design, did the manufacturer recall all products manufactured since the device was introduced to the market? If not, why not?
 - If the problem was introduced via a design change, did the manufacturer follow established design change or change control procedures? If yes, are the procedures adequate? Was the nature of the problem such that it should have been anticipated, and the design verification/ validation study fashioned to detect the problem?
 - Has the manufacturer recalled all products distributed since the design change was introduced? If not, why not?

7.2.3.2 - Corrective Action

- Describe the corrective action taken to correct the immediate problem, e.g., redesign, modify SOP, process validation, etc.
- Did the firm qualify/validate the corrective action?
- Did the firm establish responsibility to assure that the corrective action would be implemented and satisfactorily completed?
- What action did the firm take to prevent recurrence of the nonconformance, e.g., training, increased process monitoring, etc.?
- Was the nonconformance information provided to those responsible for the areas in which the nonconformance occurred?
- Did the firm determine if the nonconformance extended to other devices?
- Did the firm determine if changes were needed in procedures and, if so, did it validate and implement the changes?
- Has the manufacturer taken appropriate corrective action?

7.2.3.3 - Complaint and Medical Device Reporting (MDR) Reporting

Determine if adequate complaint investigations were performed as required by [21 CFR 820.198](#) (b). Also, determine if the investigation verified the complaint was a failure of the device to meet any or all of its specifications.

For complaints related to the recall, the firm should have made a determination whether the events are MDR reportable. Any event associated with a death or serious injury must be reported under MDR. Malfunctions likely to cause or contribute to a death or a serious injury are also reportable under MDR. Document the firm's explanations for the events they believe are nonreportable. Failure to submit required MDR reports are violations, and should be listed on the FDA-483 at the completion of the inspection.

Provide adequate documentation with the EIR to cross-reference complaints with associated MDRs.

Device Information - Obtain the 510(k) or PMA number for each device under recall. If there is no 510(k) or PMA, determine if the device is a pre-enactment device (i.e., in commercial distribution prior to May 26, 1976). If multiple devices are being recalled, obtain this information for each device model or catalog number under recall.

7.2.4 - DRUG RECALLS

7.2.4.1 - Recalls of Human Drug Products

If the recalled product is covered by a New Drug Application (NDA) or Abbreviated New Drug Application (ANDA), determine if the defective product involves the type of problems shown under [CFR 314.81](#) (b)(1)(i) and (ii). Also note whether or not the firm reported the problem to the FDA district office that is responsible for the firm within 3 working days of its receipt of the information, as required by that section.

7.2.4.2 - Recalls of Veterinary Drug Products

Veterinary Drug Products Recalls are classified by and health hazard evaluations are obtained through CVM's Division of Compliance (HFV-230), Eric Nelson, Director. To inquire about specific veterinary drug product recalls or to obtain information on how to proceed, contact the Division at 240-276-9200.

7.2.5 - HUMAN CELLS, TISSUES, AND CELLULAR AND TISSUE BASED PRODUCTS (HCT/PS) FOR IMPLANTATION, TRANSPLANTATION, INFUSION, OR TRANSFER

The agency may consider an order of retention, recall, destruction, or cessation of manufacturing when any of the conditions specified in [21 CFR 1271.440](#) (a)(1) to (3) exist. The conditions include an agency finding that:

- The HCT/P is infected or contaminated so as to be a source of dangerous infection to humans; or
- An establishment is in violation of the regulations in this part and, therefore does not provide adequate protections against the risks of communicable disease transmission.

In addition to the conditions noted above, the agency may issue an order of cessation of manufacturing until compliance with the regulations has been achieved, as stated in [21 CFR 1271.440](#) (a)(3), when the FDA determines there are reasonable grounds to believe there is a danger to health. An order to cease manufacturing would be issued where violations create an urgent situation involving a communicable disease, because an establishment is in violation of the regulations in Part 1271 and, therefore, does not provide adequate protections against the risks of communicable disease transmission. An order to cease manufacturing is a remedial action taken to put important protections in place to prevent communicable disease transmission.

NOTE: FDA will not issue an order for the destruction of reproductive HCT/Ps, nor will FDA carry out such destruction itself ([21 CFR 1271.440](#) (f)).

7.2.6 – TOBACCO PRODUCT RECALLS

When you become aware of a possible tobacco product recalls or obtain information, contact the Center for Tobacco, Office of Compliance and Enforcement, Division of Enforcement and Manufacturing. See CTP's intranet site for contact information." <http://inside.fda.gov:9003/CTP/ucm249908.htm>

7.2.7 - SAMPLE COLLECTION

Collection of samples for regulatory consideration is at the discretion of District management. Consult your supervisor and/or compliance branch for guidance. If a sample is indicated, only collect documentary samples for electronic products or medical devices, unless otherwise instructed.

If, after consulting with the Centers, it is determined that a product must be examined physically for health hazard evaluation, ship an appropriate sample to the designated Center office by the most expeditious and practical means available. Notify the Center of the time and method you sent the product and its estimated time of arrival.

7.2.8 - RECALL ALERT

When a District learns of or confirms a recall situation exists or is planned, the district R&E Coordinator will send the appropriate Center Recall Office and OEIO/DE a twenty-four hour alert through the Recall Enterprise System (RES) with basic information regarding the recall. See [RPM Chapter 7-10, Attachment A](#), "Recall Alert Information."

7.2.9 - RECOMMENDATION FOR RECALL NUMBER

The district should prepare a memorandum, containing the recall recommendation [RPM Chapter 7, Attachment B](#) as soon as possible. From the time the recall alert is sent to

the appropriate Center, the district has five days to submit the Recall Recommendation (ten days if the recall is completed). Do not wait for writing, typing and submission of the EIR or investigative memorandum. If the memorandum is prepared by the Investigator, a copy of the memo should be attached to the EIR or investigative memo so the information need not be repeated in the body of the report.

SUBCHAPTER 7.3 - MONITORING RECALLS

7.3.1 - INSPECTIONS TO MONITOR RECALL PROGRESS

It may be necessary to re-inspect the firm between the initiation and closeout of a recall to monitor its progress and verify the recalled product's disposition. These visits are limited inspections; issue an FDA-482, Notice of Inspection, at each one. Request recalling firms submit periodic status reports to FDA. See [21 CFR 7.53](#).

7.3.2 - FDA RECALL AUDIT CHECKS

7.3.2.1 - Definition

A recall audit check is a personal visit, telephone call, letter, or a combination thereof, to a consignee of a recalling firm, or a user or consumer in the chain of distribution. It is made to verify all consignees at the recall depth specified by the strategy have received notification about the recall and have taken appropriate action.

7.3.2.2 - Level of Audit Checks

Level A - 100% of the total number of consignees to be contacted.

Level B - Greater than 10% but less than 100% of the total number of consignees to be contacted.

Level C - 10% of the total number of consignees to be contacted.

Level D - 2% of the total number of consignees to be contacted.

Level E - No audit checks. (0% of the total number)

NOTE: The audit check levels listed in an assignment may vary depending on the type of consignee being audited.

Follow the levels indicated in your assignment. If you cannot meet the levels indicated in your assignment, contact your district's Recall Coordinator for further instruction.

7.3.2.3 - Conducting a Recall Audit Check

The purpose of a recall audit check is to confirm the account received the recall notification from the notifying firm and followed all instructions included in the notification. The notifying firm may be the recalling firm, or a downstream account that received the recalled product and is conducting a sub-recall (such as a distributor).

Prior to conducting a recall audit check, review the recall audit check assignment given to you. Your assignment will contain the necessary details of the recall, recall strategy, and a list of accounts to be audited (Please Note: The assignment may list specific accounts to be audited or may provide a list of accounts to choose from). Pay particular attention to the type of product recalled, the labeling of the product, and the recall notification attached to the assignment which the recalling firm sent to their accounts. Take note of the depth of the recall listed in the assignment (i.e. wholesale, retail, consumer level). Your responsibility is to verify the account received the same recall information, they followed the instructions in the recall notification, and that the recall has been carried out to the appropriate depth listed in the assignment. The assignment will include how checks will be conducted, i.e., visit, phone calls, etc. as well as detailed instructions specific to the recall. During your review of the assignment, try to gain an understanding of the list of accounts, and whether those listed actually received or may have received the recalled lot. This information affects the endorsement for the audit check. If the list is specific to the recalled product lot, the account should have received it. If the list is not so specific, or the account you are auditing does not know or remember if they received the recalled lot, the account should still follow the instructions in the recall notification and initiate a sub-recall of the product, if needed. This information affects the endorsement of the recall (see section 7.3.2.6 Endorsing the Recall Audit Check).

When initiating a recall audit check, attempt to make contact with an individual at the account who has knowledge of the receipt of recall notifications and the disposition of recalled products. In hospitals, this responsibility may be held within the Risk Management or Safety departments. PLEASE NOTE: In the case of an audit check at the consumer level, attempt to verify you are speaking with the individual who was indicated as having received the product before disclosing the name of the recalled product and verifying they received notification of the recall.

If the account did not have any knowledge of the recall prior to your recall audit check, inform them of the recall, provide them with a copy of the press release (if available) and recall notification letter, encourage them to follow the recall instructions, and document that you did so. DO NOT give the account a copy of your recall assignment.

If your audit check discloses the account did not follow the recall instructions (for example, recalled product being held for sale, or a requested sub-recall has not been initiated), encourage the account to follow the recalling firm's instructions. If the account chooses not to follow the recall instructions, document the title/responsibility of the

individual at the account who chose not to follow the recall instructions and reason.

When you conduct an audit check by visit, it is important to examine the storage sites where the recalled product is stored and check the shelf stock to ensure all recalled product has been identified, removed from areas of use, and properly quarantined. This is especially important in Class I recalls.

For some recalls, the strategy may be a correction instead of a removal. Recall audit check assignments for field corrections may instruct you to verify that either the field correction has been completed, or to assess whether the recalling firm issued the initial instructions to discontinue and/or modify the use of the product, and the account followed those instructions. Detail the status of the correction in the remarks section of your FDA 3177.

If you encounter a refusal to permit entry or provide information during a recall audit check, document the name and title of the individual who refused, and the reason why they refused the audit check. Contact your supervisor for additional instruction.

Do not conduct recall audit checks at DOD and VA facilities, as the FDA has a Memorandum of Understanding with them, and they have their own procedures for recalls. If during your audit check you find that the consignee used the FDA regulated product to manufacture USDA-regulated product complete the recall audit check. Provide the information to your district Recall Coordinator, who will forward it to OEIO (Office of Enforcement and Import Operations), who will share it with the USDA.

During your audit check, verify that the consignee has conducted a sub-recall to the level specified in the assignment. If the consignee is unsure if they handled the recalled product, then collect the distribution list. Inform the consignee that a sub-recall may be necessary. If an account has not conducted a sub-recall, follow the procedures outlined in IOM 7.3.2.5 Ineffective Recalls.

If the account did not have any knowledge of the recall prior to your recall audit check, inform them of the recall, provide them with a copy of the press release (if available) and recall notification letter, encourage them to follow the recall instructions, and document that you did so. DO NOT give the account a copy of your recall assignment.

7.3.2.4 - Audit Check Reporting

The results of your audit check should be reported on an FDA 3177, "Recall Audit Check Report" form. See IOM Exhibit 7-3. Districts have the option of completing the FDA 3177 electronically or as a hard copy. Directions for completing the FDA 3177 can be found in section 7.3.2.3 Conducting the Recall Audit Check. The FDA 3177 will be routed to your district's Recall Coordinator following your district policy.

Identified exhibits should be submitted with your FDA 3177. Identify each page with the following information:

- RES Event number (as listed in your assignment)
- Direct account name or sub-account name, whichever is applicable
- Investigator's initials and date of the audit check
- Exhibit and page numbers

FACTS allows you to enter the amount of time spent conducting your audit check. When you complete a recall audit check, you should report your time using the "Miscellaneous Operations Accomplishment Hours" screen using the code OP 17.

7.3.2.5 - Ineffective Recalls

An audit check is considered ineffective if one of the following conditions were found:

- A. The account did not receive formal notification from the notifying firm. Note: in instances where the account was not formally notified but still took action based on information learned about the recall from a source other than the notifying firm, the audit check is still ineffective.
- B. The account did not follow the instructions provided by the notifying firm.
- C. The account distributed the recalled product, but did not conduct a sub-recall, if applicable.
- D. The account received the type of product under recall, but cannot determine whether they received the specific recalled lots.

7.3.2.6 Endorsing the Recall Audit Check

Recall audit checks should be endorsed by either the Supervisory Investigator or Recall Coordinator as per district policy based on the information collected during the audit check.

The audit check should be endorsed based on conditions found when the audit check was conducted and not based on the account's actions to correct ineffectiveness. Choose the endorsement that is best described by one of the scenarios below.

An audit check should be endorsed "Effective" if the account was notified of the recall by the appropriate notifying firm and followed, or is in the process of following, the instructions in the recall notification.

The following are examples of ineffective recall audit checks:

- A. "Ineffective – Recalling Firm"

- The account did not receive formal notification from the notifying firm. Note: in instances where the account was not formally notified but still took action based on information learned about the recall from a source other than the notifying firm, the audit check is still ineffective.

B. "Ineffective – Consignee"

- The account did not follow the instructions provided by the notifying firm.
- The account distributed the recalled product, but did not conduct a sub-recall, if applicable.
- The account received the type of product under recall, but cannot determine whether they received the specific recalled lots.

Your district's Recall Coordinator can assist you if you need help evaluating if an account must conduct a sub-recall. In some instances (e.g. field corrections) the effectiveness of the recall audit check may be determined by the assignment and discussion with the recall coordinator.

If the account assigned for a recall audit check is out of business, endorse the audit check as "Out of Business".

Endorse as "Other" if the account cannot remember whether or not they received the recall notification and does not carry the recalled product.

Do not endorse recall audit checks as "Does Not Carry Product" as this does not give sufficient information as to if notification of the recall was received.

7.3.3 - RECALL TERMINATED/RECALL COMPLETED

7.3.3.1 - Definitions

Recall Terminated - A recall will be terminated when the FDA determines that all reasonable efforts have been made to remove or correct the violative product in accordance with the recall strategy, and when it is reasonable to assume that the product subject to the recall has been removed and proper disposition or correction has been made commensurate with the degree of hazard of the recalled product. Written notification that a recall is terminated will be issued by the appropriate District office to the recalling firm.

Recall Completed - For monitoring purposes, the FDA classifies a recall action "Completed" when all outstanding product, which could reasonably be expected is recovered, impounded, or corrected.

7.3.3.2 - Closeout Inspection

The final monitoring step is a limited inspection made to verify recall closeout by the recalling firm. A memorandum or limited EIR should be prepared. See [RPM Chapter 7, Attachments B1](#), "Recommendation for Recall Classification and Termination" and [Attachment C](#), "Recall Termination or Recommendation for Termination" for the format. Portions of this format (i.e., Section II and certain items in Section III) will be completed by your supervisor, R&E Coordinator, or compliance officer, depending upon your District's policy.

During the closeout inspection, you should witness destruction or reconditioning of the recalled product when possible. If you are unable to witness the destruction or reconditioning, obtain written documentation from the firm and/or any state or local government agencies which may have witnessed or otherwise verified product disposition. The disposal of large amounts of contaminated or hazardous items may require the firm to file an Environmental Impact Statement (EIS), or pre-disposal processing to render the goods harmless. Do not agree to witness destruction without resolution of these issues. Obtain a "Letter of Voluntary Destruction" from the firm whenever you witness this operation. See IOM 2.6.4.1.

SUBCHAPTER 7.4 - SPECIAL RECALL SITUATIONS

7.4.1 - General

There are several special recall situations which may require you to deviate from the normal recall procedures. Seek your supervisor's or R&E Coordinator's guidance on these. Examples include:

1. Products in the possession of U.S. Defense Installations;
2. NDA and ANDA withdrawals;
3. National Academy of Science (NAS)/Nuclear Regulatory Commission (NRC) (DESI) recalls of drugs judged ineffective; and
4. Recalls involving jurisdiction of more than one Federal Agency (e.g., FDA/EPA, FDA/Consumer Product Safety Commission (CPSC), etc.).

MODEL DRUG RECALL LETTER

John Doe Laboratories
Somewhere, U.S.A. 12345

Control Division
Date _____

(red print) --URGENT: DRUG RECALL -- Nonsterile injectable

Re: List 1234, Cyanocobalamin Injection Lot No. 4321

Recent tests showed that the above lot number of this product is not sterile and therefore, represents a potential public health hazard. Consequently, we are recalling this lot from the market. Other lot numbers are not involved.

Please examine your stocks immediately to determine if you have any of Lot 4321 on hand. If so, discontinue dispensing the lot and promptly return via parcel post, to our New York City Plant; ATENTION RETURNED GOODS.

(NOTE: If a sub-recall is indicated in a particular situation, the following paragraph should be added:)

“If you have distributed any of lot 4321, please immediately contact your accounts, advise them of the recall situation, and have them return their outstanding recalled stocks to you. Return these stocks as indicated above.”

You will be reimbursed by check or credit memo for the returned goods and postage.

Please return the enclosed card immediately providing the requested information.

This recall is being made with the knowledge of the Food and Drug Administration. The FDA has classified this recall as class _____ (if classified).

We appreciate your assistance.

John Doe
President

PLEASE FILL OUT AND RETURN

We do not have any stock of List 1234, Cyanocobalamin

Injection Lot No. 4321 on hand

We have requested our accounts to return their stocks of this merchandise to us.

We are returning _____ bottles of List 1234, Lot No. 4321

Name _____

Address _____

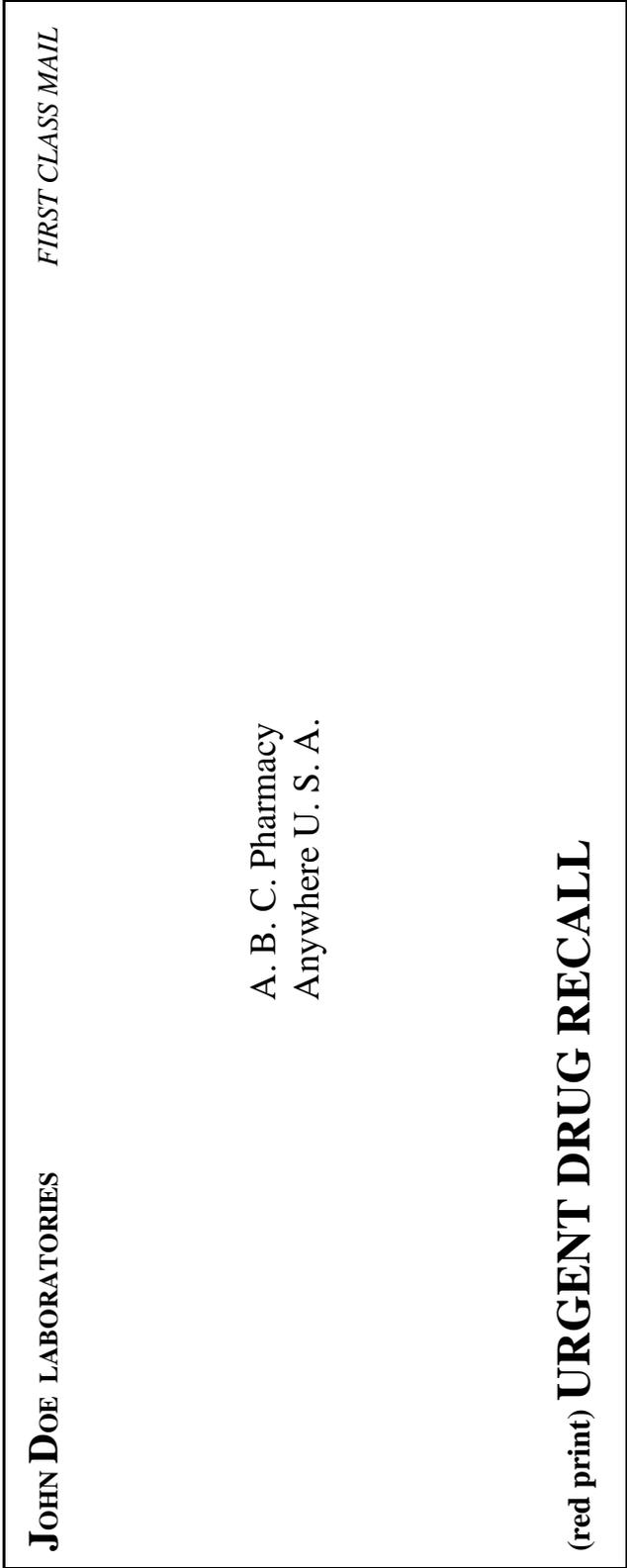
First Class
Permit No. 2

BUSINESS REPLY MAIL
No Postage Stamp Necessary if mailed in U.S.A.

Postage will be paid by:

JOHN DOE LABORATORIES
Somewhere, U.S.A. 12345-0909

Henry Doe



1. RECALL INFORMATION

a. RES/RECALL NUMBER(S)	b. RECALLING FIRM	c. RECALLED CODE(S)	d. PRODUCT(S)
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2. PROGRAM DATA

a. HOME DISTRICT	b. FEI NUMBER OF RECALLING FIRM
c. PAC CODE	d. HOURS

3. AUDIT ACCOUNTS

a. DIRECT	b. SUB-ACCOUNT (SECONDARY)
PHONE NO.:	PHONE NO.:
c. SUB-ACCOUNT (TERTIARY)	
PHONE NO.	

4. CONSIGNEE DATA

Contacted by: <input type="checkbox"/> Phone <input type="checkbox"/> Visit <input type="checkbox"/> Other a. NAME OF PERSON CONTACTED & TITLE	b. TYPE CONSIGNEE <input type="checkbox"/> Distributor <input type="checkbox"/> Consumer <input type="checkbox"/> Pharmacy <input type="checkbox"/> Retailer <input type="checkbox"/> Physician <input type="checkbox"/> Restaurant <input type="checkbox"/> Processor <input type="checkbox"/> Hospital <input type="checkbox"/> School <input type="checkbox"/> Other: _____	c. DOES (DID) THE CONSIGNEE HANDLE RECALLED PRODUCT? <input type="checkbox"/> Yes <input type="checkbox"/> No
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5. NOTIFICATION DATA

a. FORMAL RECALL NOTICE RECEIVED? (If "No", skip to item 6c.) <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Cannot be determined (If answer is other than "No", explain in remarks.)	b. RECALL NOTIFICATION RECEIVED FROM <input type="checkbox"/> Recalling Firm <input type="checkbox"/> Other (Specify below) <input type="checkbox"/> Direct Account <input style="width:50px; height:20px;" type="text"/> <input type="checkbox"/> Sub-Account	c. DATE NOTIFIED (mm/dd/yyyy) d. TYPE OF NOTICE RECEIVED (e.g., letter, phone)
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6. ACTION AND STATUS DATA

a. DID CONSIGNEE FOLLOW THE RECALL INSTRUCTIONS? (If "No", discuss in "Remarks" action taken upon FDA contact.) <input type="checkbox"/> Yes <input type="checkbox"/> No	c. CURRENT STATUS OF RECALLED ITEMS <input type="checkbox"/> Returned <input type="checkbox"/> None on Hand <input type="checkbox"/> Corrected <input type="checkbox"/> Was Still Held for Sale/Use* <input type="checkbox"/> Destroyed <input type="checkbox"/> Held for Return/Correction* * = Ensure Proper Quarantine/Action	7. SUB-RECALL NEEDED? Did consignee distribute to any other accounts? (If "Yes", collect information and/or provide details in "Remarks" or Memo.) <input type="checkbox"/> Yes <input type="checkbox"/> No
b. AMOUNT OF RECALLED PRODUCT ON HAND AT TIME OF NOTIFICATION	d. DATE AND METHOD OF DISPOSITION	8. AMOUNT OF RECALLED PRODUCT NOW ON HAND

9. INJURIES/COMPLAINTS

a. IS CONSIGNEE AWARE OF ANY INJURIES, ILLNESS, OR COMPLAINTS?

Injury Complaint
 Illness None

If answer is other than "None", report details in a separate memo to monitoring district and copy to OEO (HFA-615).

10. REMARKS (Include action taken if product was still available for sale or use.)

CHECK		ENDORSEMENT	
INVESTIGATOR	SCSO OR R&E COORDINATOR	<input type="checkbox"/> Effective <input type="checkbox"/> Does Not Carry Product <input type="checkbox"/> Ineffective (Indicate level) <input type="checkbox"/> Out of Business <input type="checkbox"/> Recalling Firm <input type="checkbox"/> Consignee <input type="checkbox"/> Other (Specify): _____	
Signature	Signature		
Printed Name	Printed Name		
Date of Check (mm/dd/yyyy)	District	Date of Endorsement (mm/dd/yyyy)	

Completing the FDA 3177 Recall Audit Check Report Form

Note: Obtain as much information as possible in order to successfully complete the FDA 3177 Recall Audit Check Report Form as follows:

1. RECALL INFORMATION

- a. RES/RECALL NUMBER(S) – Enter the recall number and Recall Enterprise System (RES) number as listed in your assignment. If more than one recall number is involved, enter the lead number. In some cases, a recall number may not have been assigned yet. In these cases, 000 or 0000 will be entered as the recall number (for example, a medical device recall that does not yet have a recall number assigned, will be listed here as “Z-0000-20XX” where “XX” is the fiscal year).
- b. RECALLING FIRM – Provide the name and address of the firm listed in your assignment as the recalling firm.
- c. RECALLED CODE(S) – Provide the lot, batch, or serial number indicated as the recalled product in your assignment. If there are more numbers than can fit in the space, state that there are numerous lots under recall and refer to the assignment.
- d. PRODUCT(S) – Provide the name of the recalled product as indicated in your assignment. If numerous products are involved, use a generic term (such as ice cream, dried fruit, etc.).

2. PROGRAM DATA – Complete as per District policy.

- a. HOME DISTRICT – Monitoring district for the recall, as listed in your assignment.
- b. FEI NUMBER OF RECALLING FIRM – FEI number of the recalling firm as listed in your assignment.
- c. PAC CODE – PAC code given in your assignment.
- d. HOURS – Hours spent preparing, conducting, and completing the FDA 3177 for the audit check.

3. AUDIT ACCOUNTS

- a. DIRECT – The name, address, and telephone number of the account that was listed in your assignment as receiving the product directly from the recalling establishment. This may or may not be the same account at which you are conducting your audit check.
- b. SUB-ACCOUNT (SECONDARY) – If the Direct account indicates the recalled product(s) were further distributed, complete this section for each sub-account audited as well as the DIRECT account section with the name, address, and telephone number of the applicable establishments.
- c. SUB-ACCOUNT (TERTIARY) – If the Secondary account indicates the recalled product(s) were further distributed, complete this section for each sub-account audited, the SUB-ACCOUNT (SECONDARY) section, and the DIRECT account section with the name, address, and telephone number of the applicable accounts.

4. CONSIGNEE DATA

Contacted by: The method used to conduct the audit check (check the appropriate box).

- a. NAME OF PERSON CONTACTED & TITLE – The name and title of the person at the account being audited who provided the most information during the audit check.
- b. TYPE CONSIGNEE – The type of establishment at which you are conducting your audit check (check the appropriate box – if none, check “Other” and describe the type of establishment).
- c. DOES (DID) THE CONSIGNEE HANDLE RECALLED PRODUCT? – If the account at which you are conducting the audit check never received the recalled product, indicate “No”. If the account received or may have received the recalled product, indicate “Yes”. This includes if the company is unsure they received the recalled lot.

5. NOTIFICATION DATA

- a. **FORMAL RECALL NOTICE RECEIVED?** – Indicate if the account received formal notification of the recall (check the appropriate box). Formal notification may be received from the recalling firm, or the secondary/tertiary firm. If notification is received informally e.g. press release, subscription service, or social media, indicate “No” and explain in Remarks how the account received notification. If there is some reason why you cannot determine if a notification was received (for example, it may have been discarded) indicate “Cannot be determined” and explain in Remarks
- b. **RECALL NOTIFICATION RECEIVED FROM** – The firm that formally notified the account at which you are conducting your audit check (check the appropriate box).
- c. **DATE NOTIFIED** – Date the account received the formal notification.
- d. **TYPE OF NOTICE RECEIVED** – How the formal notification was received (letter, phone, e-mail, automated messaging system, etc.).

6. ACTION AND STATUS DATA

- a. **DID CONSIGNEE FOLLOW THE RECALL INSTRUCTIONS?** – If the account followed or is following all of the recall instructions prior to your audit check, indicate “Yes”. If the account did not follow or has not begun to follow the recall instructions prior to your audit check, indicate “No”. Explain what was/was not done in Remarks, and if the account took action as a result of your audit check.
- b. **AMOUNT OF RECALLED PRODUCT ON HAND AT TIME OF NOTIFICATION** – The amount of recalled product the account had at the time they received formal notification from the notifying firm.
- c. **CURRENT STATUS OF RECALLED ITEMS** – Indicate the status of the recalled items at the account at the time of your audit check (check the appropriate box). If the recalled product is still being held for sale/use, or was being held for return/correction, ensure that the account properly quarantined the product (if applicable) and followed the recall instructions. Include details in the Remarks
- d. **DATE AND METHOD OF DISPOSITION** – Indicate the date and method the recalled product was destroyed/returned/corrected.

7. SUB-RECALL NEEDED? – If during the course of an audit check, you find the recalled product has been further distributed, and your audit check for the recall has not reached the depth indicated in your assignment, a sub-recall may be needed. For example, if your assignment indicates the recall depth is at the retail level, and you are auditing a wholesaler, the wholesaler should conduct a sub-recall to reach the retail level.

Determine if the consignee followed the instructions and conducted a sub-recall. If they did not, then inquire with the consignee about their willingness to continue the recall to the depth specified in the recall strategy and gather as much distribution information as possible. Indicate “Yes” in this section and add as much detail as possible in Remarks.

In some cases, if the consignee has re-labeled, repackaged, or remanufactured the recalled product, a new recall may be needed instead of a sub-recall. However, a new recall may not be needed, if the consignee has manipulated the recalled product in a way that corrects the initial reason for the recall (e.g. if the consignee re-labels the product so the labeling issue is no longer a concern, or if the consignee heat treats the product adequately to eliminate the hazard causing the original recall). If you determine a new recall is needed, or are unsure, collect all relevant information, including labeling to be evaluated with the assistance of your district’s Recall Coordinator (refer to section 7.3.2.4 for labeling instructions of attachments).

Indicate “No” in this section if the product has not been further distributed and your evaluation finds that a sub-recall is not necessary.

8. AMOUNT OF RECALLED PRODUCT NOW ON HAND – The amount of recalled product still at the

account during your audit check.

9. INJURIES/COMPLAINTS

a. **IS CONSIGNEE AWARE OF ANY INJURIES, ILLNESS, OR COMPLAINTS?** – Ask the consignee if they have firsthand knowledge of any injuries, illness, or complaints pertaining to the recalled product. Collect relevant information and report the findings to your supervisor.

10. REMARKS – Use this section to provide details that could not be addressed in the previous sections, or to give additional information.

CHECK – Place a handwritten or electronic signature, followed by your name printed or typed, the date your audit check completed, and your district.

ENDORSEMENT – Follow section 7.3.2.6 Endorsing the Audit Check.