

**1. RECALL INFORMATION**

a. RES NUMBER	b. RECALLING FIRM	c. RECALLED CODE(S)	d. PRODUCT(S)
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<b>2. PROGRAM DATA (FDA Users Only)</b>		<b>3. AUDIT ACCOUNTS</b>	
		a. DIRECT	b. SUB-ACCOUNT (SECONDARY)
a. MONITORING DIVISION	b. FEI NUMBER OF RECALLING FIRM	PHONE NO.:	PHONE NO.:
c. PAC CODE		c. SUB-ACCOUNT (TERTIARY)	PHONE NO.

<b>4. CONSIGNEE DATA</b>		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
Contacted by: <input type="checkbox"/> Phone <input type="checkbox"/> Visit <input type="checkbox"/> Other a. NAME OF PERSON CONTACTED & TITLE			

<b>5. NOTIFICATION DATA</b>		b. RECALL NOTIFICATION RECEIVED FROM	c. DATE NOTIFICATION RECEIVED (mm/dd/yyyy)
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 "Yes", explain in remarks.)		<input type="checkbox"/> Recalling Firm <input type="checkbox"/> Other (Specify below) <input type="checkbox"/> Direct Account <input type="text"/> <input type="checkbox"/> Sub-Account	d. TYPE OF NOTICE RECEIVED (e.g., letter, phone)

<b>6. ACTION AND STATUS DATA</b>		c. CURRENT STATUS OF RECALLED ITEMS	<b>7. SUB-RECALL NEEDED?</b> <i>Did consignee distribute to any other accounts? (If "Yes", collect information and/or provide details in "Remarks" or Memo.)</i>
a. DID CONSIGNEE FOLLOW THE RECALL INSTRUCTIONS? (If "No", discuss in "Remarks" action taken as a result of audit check.) <input type="checkbox"/> Yes <input type="checkbox"/> No		<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	<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	<b>8. AMOUNT OF RECALLED PRODUCT NOW ON HAND</b>

<b>9. INJURIES/COMPLAINTS</b>	<b>10. REMARKS</b> (Include action taken if product was still available for sale or use.)
a. IS CONSIGNEE AWARE OF ANY INJURIES, ILLNESS, OR COMPLAINTS? <input type="checkbox"/> Injury <input type="checkbox"/> Complaint <input type="checkbox"/> Illness <input type="checkbox"/> None If answer is other than "None", collect relevant information, document findings, and route per division procedures.	

<b>CHECK</b>		<b>FDA ENDORSEMENT</b>	
Signature		Signature	
Printed Name and Title		Printed Name and Title	
Date of Audit Check (mm/dd/yyyy)	FDA Division	Date of Endorsement (mm/dd/yyyy)	<input type="checkbox"/> <b>Effective</b> <input type="checkbox"/> <b>Out of Business</b>  <input type="checkbox"/> <b>Ineffective</b> (Indicate level) <input type="checkbox"/> Notifying Firm <input type="checkbox"/> Consignee  <input type="checkbox"/> <b>Other</b> (Specify): _____