



Pharmacy Department





#### **Drug Recall**

are actions taken by the firm to remove a product from the market. (U.S. FDA) Recalls can be conducted by firm's own initiative ,or by FDA request or by FDA order under statutory authority. (U.S. FDA)

## Products for Drug Recall

### **1. Near expiry drugs**

2. Suspected adulterated drugs

3. Drugs and medical supplies not properly sealed

# 4. Drugs with adverse reaction

5. Drugs declared by companies for drug recall



1.Distributors/ Companies should be informed first regarding the status of their products.

2. For any adverse drug event, an investigation will be conducted through the Pharmacy and **Therapeutics Committee.** 

2.1 After the decision has been made, the company concerned will be called for.



1. Products for drug recall should be placed in a separate shelf properly labeled to avoid dispensing. 2.Distributors/ Companies should be called for to check the products and retrieve particular invoice. 3. Investigation and confirmation will be done by the company.

**4.** After the confirmation, a report will be furnished in two copies one for the company and one for the Chief Pharmacist before the product is recalled.

5. Following the return and refund policy, the company has to refund the drugs/products recalled.

#### COMPLIANCE:

Quantity	Name of Drug	Expiry Date	Date Recalled
4 bots	Salbutamol/Guiafenesin 1mg/50mg /5ml expectorant 60 ml	Oct.2016	Sept.2016
4 bots.	Co-Amoxiclav 156.25mg/ 5ml 60 ml	Oct.2016	Sept. 2016
4 bots	Clarithromycin 125mg/5ml 25ml	Oct.2016	Sept.2016

