Life-Assist
Helping Heroes Save Lives

Baxter Recall EXPANDED- Presence of Particulate Matter

Thursday, January 18, 2018

Dear Valued Life-Assist Customer,

According to our records you may have purchased an item that has been recalled by the manufacturer. Please examine your stock to determine if you have the following product, with an affected lot number in your possession.

NEXTERONE (amiodarone HCl) Premixed Injection, 150 mg/100 mL

Life-Assist, Inc. Product Code	Baxter Product Code	NDC	Lot #
DR0150-10	2G3451	43066-150-10	NC109123
DR0150-10	2G3451	43066-150-10	NC109925

Reason for Recall:

Baxter Healthcare Corporation is expanding the voluntary drug recall of NEXTERONE Injection issued November 10, 2017 to include one additional lot of product (NC109123) – distributed in the United States to wholesalers/distributors and healthcare facilities – due to the potential presence of particulate matter.

Please see the enclosed recall letter from the manufacturer for more information.

Action Required:

1. Immediately forward this notification to any pertinent personnel, department, and/or location within your organization.
2. Check your inventory and quarantine any product with an affected lot number. If the affected stock has been depleted, you may disregard this notification.
3. Contact Life-Assist Customer Service at 800-824-6016 or salesservice@life-assist.com to obtain a Return Authorization.

Customers with questions regarding this recall can contact Baxter Corporate Product Surveillance at 800-437-5176, Monday through Friday, between 8 a.m. and 5 p.m. Central Time. We apologize for any inconvenience.