November 18, 2013

Dear Director of Pharmacy:

<table>
<thead>
<tr>
<th>Affected Product</th>
<th>Lot</th>
<th>Product Description</th>
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| 2B7729
NDC: 0338-1136-03 | P287045 | CLINIMIX 4.25/25 sulfite-free (4.25% Amino Acid in 25% Dextrose) Injection in CLARITY Dual Chamber Container |
| 2B7717
NDC: 0338-1115-04 | P275883 | CLINIMIX E 4.25/10 sulfite-free (4.25% Amino Acid with Electrolytes in 10% Dextrose with Calcium) Injection in CLARITY Dual Chamber Container |
| 2B7709
NDC: 0338-1099-04 | P285122 | CLINIMIX 5/15 sulfite-free (5% Amino Acid in 15% Dextrose) Injection in CLARITY Dual Chamber Container |

Problem Description
Baxter Healthcare Corporation has received complaints reporting the presence of various types of particulate matter associated with CLINIMIX and CLINIMIX E products. Baxter Healthcare Corporation is issuing a voluntary recall for the above-referenced lots. Baxter distributed this product to customers between May 2012 and October 2013.

Hazard Involved
The intravenous infusion of a solution with particulate matter could cause vein irritation or lead to blood vessel occlusion, including a pulmonary embolism. Baxter has not been made aware of any adverse events associated with these lots.

Action to be taken if product was purchased directly from Baxter
Baxter is requesting that you take the following actions:

1. Check all locations; remove all affected product from your facility and quarantine product subject to recall. The product code and lot number can be found on the individual product package or shipping carton.

2. Contact Baxter Healthcare Center for Service to arrange for return and credit/replacement product. The Center for Service can be reached at 1-888-229-0001 between the hours of 7:00 am and 6:00 pm Central Time. Please have your Baxter eight-digit ship-to account number ready when calling.

3. If your product was purchased from Baxter, complete the attached Reply Form (Attachment 1) and return it to Baxter either by faxing the Form to 1-224-270-5457 or scanning and e-mailing it to fca@baxter.com. Returning the Reply Form promptly will prevent you from receiving repeat notifications.

4. Please notify other facilities, departments, or services if they distribute, transfer or sell the affected product.

5. If you are a dealer, wholesaler, or distributor/reseller that distributed any product to other facilities, please conduct a recall to notify your end user customers in accordance with your customary procedures.
Action to be taken if product was purchased from a distributor or reseller

1. Locate and remove all affected product from your facility (the product code and lot number can be found on the individual product package or shipping carton).

2. Contact Baxter Healthcare Center for Service to arrange for return and credit. The Center for Service can be reached at 1-888-229-0001 between the hours of 7:00 am and 6:00 pm Central Time. Please have your Baxter eight digit ship-to account number ready when calling.

3. Follow your supplier’s reply and recall process. Please do not return your supplier’s reply form to Baxter.

Further Information and support

- If you have clinical questions regarding this communication, please call Medical Information Services at Baxter at 1-800-422-2751 during the hours of 8:00 am and 4:30 pm Central Time.

- For general questions regarding this communication, contact The Center for One Baxter at 1-800-422-9837, Monday through Friday, during the hours of 8:00 am and 5:00 pm Central Time.

The CLINIMIX and CLINIMIX E product information states that the products should be inspected visually for particulate matter and discoloration prior to administration, whenever solution and container permit. The labeling also recommends the use of a final filter during administration of the parenteral solutions, where possible.

The United States Food and Drug Administration has been notified of this action. Any side effects or quality problems experienced with the use of these products may be reported using one of the following options:

- Calling Baxter Corporate Product Surveillance at 1-800-437-5176
- Emailing Baxter at: corporate_product_complaints_round_lake@baxter.com
- Linking to the MedWatch website at www.fda.gov/medwatch/report.htm
- Calling the FDA at 1-800-FDA-1088
- Faxing to the FDA: 1-800-FDA-0178
- Mailing to: MedWatch, FDA, 5600 Fishers Lane, Rockville, MD 20852-9787

We apologize for any inconvenience this may cause you. We look forward to continuing to serve your nutrition needs and we thank you for your cooperation.

Sincerely,

Pedro Rivera
Vice President, Quality
Medical Products
Baxter Healthcare

cc: Director of Nursing
Attachment 1: Reply Form