Baxter 0.9% sodium chloride 500 mL Viaflex containers

On Mar. 25, 2015, Baxter recalled three lots (C926873, C928630, C929844) of **0.9% sodium chloride in 500mL Viaflex containers** (NDC # 00338-0049-03) after reports that some containers leaked, some had missing closures or both. The U.S. Food and Drug Administration (FDA) has not yet classed the recall, but Baxter set it at the consumer level, so it was treated as Class I.

The detailed information of the recalled product is listed below:

<table>
<thead>
<tr>
<th>Recalled Drug:</th>
<th>0.9% sodium chloride injection in 500mL Viaflex plastic containers</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>NDC Number:</strong></td>
<td>00338-0049-03</td>
</tr>
<tr>
<td><strong>Lot Numbers:</strong></td>
<td>C926873, C928630, C929844</td>
</tr>
<tr>
<td><strong>Manufacturer:</strong></td>
<td>Baxter</td>
</tr>
</tbody>
</table>

Baxter recommends that:

- Return and replacement for products from the recalled lot be arranged with the Baxter Healthcare Center for Service at 1.888.229.0001, Monday through Friday, between 7:00 a.m. and 6:00 p.m., CT.
- Patients with products from affected lots notify their physicians and return recalled sodium chloride to the pharmacies that filled it.
- Questions about the recall may be directed to onebaxter@baxter.com or to 1.800.422.9837, Monday through Friday, between 8:00 a.m. and 5:00 p.m. CT.
- Adverse events from the use of prescription drugs should be reported to FDA by visiting www.fda.gov/medwatch/report.htm or calling 1-800-FDA-0178.

Aurobindo 300mg gabapentin capsules

On Nov. 21, 2014, Aurobindo, recalled one lot (GESB14011-A) of **gabapentin capsules, 300mg**, which were distributed in 100-unit bottles under the Northstar label (NDC 16714-0662-01). Empty capsules had been reported in some bottles of the affected lot.

The recalled drugs and lot numbers are below.

<table>
<thead>
<tr>
<th>Recalled Drug:</th>
<th>gabapentin capsules, 300mg (Aurobindo, labeled as Northstar)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>NDC Number:</strong></td>
<td>16714-0662-01</td>
</tr>
<tr>
<td><strong>Recalled Lot Number:</strong></td>
<td>GESB14011-A</td>
</tr>
<tr>
<td><strong>Expiration Date:</strong></td>
<td>December 2015</td>
</tr>
</tbody>
</table>
Aurobindo recommends that:

- Product from the recalled lot should be quarantined.
- Aurobindo Pharma USA’s Pharmacovigilance group at PVG@aurobindousa.com or 732-839-9400, Option 2, should be contacted for more information about the recall.
- Adverse events from the use of prescription drugs should be reported to FDA by visiting www.fda.gov/medwatch or calling 1-800-332-1088.


**Cubicin® (daptomycin) for Injection**

Cubist Pharmaceuticals has announced the recall of multiple lots of Cubicin® (daptomycin) for injection because vials may contain glass particles. The affected vials are 500mg in 10 mL single-dose (NDC 67919-0011-01).


Cubist Pharmaceuticals recommends that:

- Recalled product should be quarantined.
- Customers who have recalled product should contact Cubist at 1-855-534-8309 Monday through Friday from 9 a.m. to 7 p.m. ET to arrange for instructions on how to return recalled product.
- For medical questions, contact Cubist Medical Information should be contacted at: 1-877-282-4786 between 8 a.m. and 5:30 p.m. ET, Monday through Friday.
- Adverse events from the use of prescription drugs should be reported to FDA by calling 1-800-332-0178 or visiting www.fda.gov/medwatch/report.htm.

**Advocate Redi-Code Plus Blood Glucose Test Strips**

On June 6, 2014, Diabetic Supply of Suncoast, Inc. recalled all lots of Advocate Redi-Code Plus Blood Glucose Test Strips (NDC – 94046-0001-57). Package labels for the recalled strips do not include the model number (BMB-BA006A). The strips were manufactured by BroadMaster Bio-Tech Corp. for specific blood glucose meters also made by BroadMaster. If the strips are used with similar meters from a different
company, blood glucose readings may be inaccurate, potentially leading to serious adverse effects.

**Specifics of the recall:**

<table>
<thead>
<tr>
<th>Product Description</th>
<th>NDC Number</th>
<th>Lot Number</th>
<th>Expiration Date</th>
<th>Manufacturer</th>
</tr>
</thead>
<tbody>
<tr>
<td>Advocate Redi-Code Plus Blood Glucose Test Strips</td>
<td>94046-0001-57</td>
<td>All</td>
<td>All</td>
<td>Diabetic Supply of Suncoast, Inc.</td>
</tr>
</tbody>
</table>

Diabetic Supply of Suncoast, Inc. recommends that:

- Recalled strips should not be used.
- Customers who have recalled strips should contact the company at 561-296-0488 between the hours of 9:00am and 5:00pm ET, Monday through Friday for instructions on how to have recalled product replaced.
- Patients or caregivers of patients who received recalled strips should talk with their prescribers.
- Adverse events from the use of prescription drugs should be reported to FDA by visiting [www.fda.gov/medwatch/report.htm](http://www.fda.gov/medwatch/report.htm) or calling 1-800-FDA-0178.