Barr / Duramed Pharmaceuticals / (Teva Pharmaceuticals USA Inc.)
URGENT MEDICAL DEVICE RECALL – RETAIL LEVEL
Initiated 3/30/2012

ViaSpan Cold Storage Solution 1000mL Bag

MANUFACTURED BY:
Fresenius Kabi (Barr / Duramed Pharmaceuticals)
Graz, Austria

RECALLED BY:
Teva Pharmaceuticals USA
Sellersville, PA 18960

<table>
<thead>
<tr>
<th>Lot #</th>
<th>Exp. Date</th>
<th>Product Code</th>
<th>Size</th>
</tr>
</thead>
<tbody>
<tr>
<td>16EK0007</td>
<td>10/2012</td>
<td>1000-46-06</td>
<td>10 x 1000mL Bags</td>
</tr>
<tr>
<td>16EK0193</td>
<td>10/2012</td>
<td>1000-46-06</td>
<td>10 x 1000mL Bags</td>
</tr>
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Dear Customer:

Teva Pharmaceuticals USA Inc. is voluntarily recalling the above mentioned lots of ViaSpan Cold Storage Solution Bags distributed under the Barr/Duramed Pharmaceuticals label. This recall is being carried out to the RETAIL LEVEL as a precautionary measure due to a lack of assurance of sterility.

Wholesalers / Distributors - Please perform the following activities:

- Examine your inventory immediately for the above lots of ViaSpan Cold Storage Solution Bags.
- Our records indicate we shipped this product between December 1, 2011 and March 19, 2012.
- Immediately discontinue distribution of the specific lots being recalled.
- **Please perform a SUB-RECALL to your RETAIL accounts using this Recall Notification and Stock Response Form.**
- Promptly complete the attached recall stock response and reply via fax number 817-868-5362 or mail, even if you have no product to return.

Completed Recall Stock Response forms can be mailed, emailed, or sent via FAX to Inmar Attn: Recall Coordinator, 4332 Empire Road Suite 200, Fort Worth, TX 76155. Inmar Email address: recallnotice@inmar.com. Inmar FAX: 817-868-5362. Inmar will send a Return Goods Authorization label and shipping label. Appropriate credit for product returns, plus handling and shipping expenses, will be issued upon receipt of said product with the Return Goods Authorization form. All recalled product returned without a Return Goods Authorization label may delay the issuance of your credit.

This recall is being made with the knowledge of the Food & Drug Administration. Your cooperation and prompt response to this notice is appreciated. If you have Customer Service related questions, please contact Teva Customer Services at 800-545-8800. Medical related questions, please contact Teva Medical Affairs at 215-641-6974. If you need a Recall Stock Response form, contact Inmar at 800-967-5952 or acquire it from clsnetlink.com.

Sincerely,

Christopher A. Murdock, PhD
Sr. Director, Regulatory Compliance
Teva Pharmaceuticals USA, Inc.
**URGENT MEDICAL DEVICE RECALL – RETAIL LEVEL**

**Initiated 3/30/2012**

**ViaSpan Cold Storage Solution 1000mL Bag**

**Please fill out completely**

**Date:** ______________

**DIRECT CUSTOMERS ONLY:** Does this response include all DC locations?  YES □  NO □

Customer/Store Name: ____________________________  DEA #: ____________________________

*DEA # is required; if not provided the processing of your form will be delayed.

Address:________________________________________

City:_________  State:_________  Zip:_________

Contact Name (please print) ____________________________  Telephone #: ____________________________

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**I have checked my stock and:**

___ I do not have stock of the recalled item(s) OR ___ I do have stock of the recalled item(s) listed above.

Please send me___________ shipping box labels

**NON DIRECT CUSTOMERS ONLY:** Please complete the following:

Purchased From (Wholesaler name): ____________________________  DEA #: ____________________________

City:_________  State:_________

**Inquiries regarding this recall are to be directed to the following:**

Recall Stock Response forms - If your return kit is not received between 7-10 business days contact Inmar at 800-967-5952, Option 1 then Option 3. Please do not resubmit response form.

Customer service related questions - contact Teva Customer Services at 800-545-8800

Medical related questions - contact Teva Medical Affairs 215-641-6974

Please fax this form to: 817-868-5362 or E-mail at: recallnotice@inmar.com