Dear [Your Hospital Name],

We, B. Braun Melsungen AG, have decided to recall the following products in the context of a FIELD SAFETY CORRECTIVE ACTION from the market:

<table>
<thead>
<tr>
<th>Article Number</th>
<th>Article Name</th>
<th>Batch</th>
</tr>
</thead>
<tbody>
<tr>
<td>4417930</td>
<td>UREOFIX 500 CLASSIC, EB.2.0L, TUBE 120CM</td>
<td>16F01E8SUA</td>
</tr>
<tr>
<td>4417920</td>
<td>UREOFIX 500 CLASSIC, EB.1.5L, TUBE 170CM</td>
<td>16F02E8SUA</td>
</tr>
<tr>
<td>4417910</td>
<td>UREOFIX 500 CLASSIC, EB.1.5L, TUBE 120CM</td>
<td>16F06E8SUA</td>
</tr>
</tbody>
</table>

**Reason for the Recall**

In the course of complaints analyses we identified a defect in UREOFIX 500 Classic. The 1st compartment empties itself in the second one before being full which makes the reading of the diuresis more difficult. Three (3) batches show this defect.

**Actions to be taken by the USER**

Our records show that your hospital has received potentially affected UREOFIX 500 Classic products as specified in the table above.

We kindly ask you to initiate the following activities immediately and with priority:

- Identify, quarantine and return affected devices.
- Do not use affected devices anymore.
- Inform the responsible personnel in the affected facilities.
- Confirm the receipt of this information.

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TO WHOM IT MAY CONCERN
Page 2 to the letter of October 24, 2016 to

If more information is needed, please contact

Local contact 1
Name
Title
Email
telephone

Local contact 2

Kindly accept our apologies for any inconveniences.

Yours sincerely,