On March 9, 2016, Teva Pharmaceuticals announced a recall of one lot of amikacin 1 gm/4 mL injection because of the potential presence of glass particulate matter.

Per Teva Pharmaceuticals, the intravenous administration of glass particulate may result in local irritation, swelling, or more serious outcomes such as blocked blood vessels and organ damage.

Catamaran, an OptumRx Company, notified members who may be affected by this recall. These members were provided information to help them identify if their medication is being recalled. In the future, these members may inquire about the amikacin injection prescription(s) you dispensed.

If you have any questions, please call Teva Pharmaceuticals at 888-838-2872, option 3, then, option 4, (8:00 am – 5:00 pm EST, Monday through Friday), for more information about this recall.

Recalled Lot of Amikacin Injection by Teva Pharmaceuticals

<table>
<thead>
<tr>
<th>Product Description</th>
<th>NDC</th>
<th>Lot #</th>
<th>Expiration Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>Amikacin sulfate injection, 1 gm/4 mL (250 mg/mL)</td>
<td>0703-9040-01 (individual pack); 0703-9040-03 (shelf pack)</td>
<td>4750915</td>
<td>9/2017</td>
</tr>
</tbody>
</table>

Help Desk phone number:

Teva Pharmaceuticals at (888)-838-2872, option 3, then option 4

Thank you for your continued support. Please distribute immediately.