IMPORTANT INFORMATION REGARDING

Recall of Linezolid Injection by AuroMedics Pharma

Effective Date: December 22, 2017

AuroMedics Pharma

Recall of Linezolid Injection

On December 22, 2017, AuroMedics Pharma announced a voluntary recall of one lot of Linezolid Injection because of a product complaint in which one bag was found to contain white particulate matter identified as mold.

Per AuroMedics Pharma, use of a non-sterile injectable product could result in fatal infections.

OptumRx notified members who may be affected by this recall. These members were provided information to help them identify if their product is being recalled and advised not to use the recalled medicine. In the future, these members may inquire about the Linezolid Injection prescription(s) you dispensed.

If you have any questions, please call Aurobindo (affiliate of AuroMedics Pharma) at (866) 850-2876 (9:00 AM – 5:00 PM EST, Monday through Friday) for more information about this recall.

Linezolid Injection Recalled by AuroMedics Pharma

<table>
<thead>
<tr>
<th>Product Description</th>
<th>NDC #</th>
<th>Lot # (expiration date)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Linezolid Injection 600 mg/300 mL flexible bags</td>
<td>55150-242-51</td>
<td>CLZ160007 (August 2018)</td>
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TO REDUCE PROCESSING ERRORS, PLEASE CONFIRM THE INFORMATION ON MEMBER’S ID CARD PRIOR TO SUBMITTING PRESCRIPTION CLAIMS.

Should you have any questions or require assistance, please contact AuroMedics Pharma:
(866) 850-2876 (9:00 AM – 5:00 PM EST, Monday through Friday)

For questions regarding this communication or other pharmacy related claims processing issues call: Provider Relations (877) 633-4701 or e-mail provider.relations@optum.com

Please distribute immediately.