IMPORTANT INFORMATION REGARDING

Recall of Pravastatin by International Laboratories

Effective Date: August 10, 2017

International Laboratories

Recall of Pravastatin

On August 10, 2017, International Laboratories announced a recall of one lot of Pravastatin 40 mg tablets because a bottle was mislabeled as Pravastatin 40 mg but contained Bupropion XL 300 mg tablets.

Per International Laboratories, if a patient mistakenly takes Bupropion, the side effects may include: nausea, vomiting, dry mouth, headache, constipation, sweating, sore throat, diarrhea, dizziness, restlessness, and blurry vision. Other more serious reactions include lowering of seizure threshold in patients with epilepsy, hypertensive crisis in patients on concomitant monoamine oxidase inhibitor therapy, and life threatening allergic reactions.

OptumRx notified members who may be affected by this recall. These members were provided information to help them identify if their medication is being recalled, advised not to use the recalled product, and to return any recalled product to their pharmacy for a replacement or refund. In the future, these members may inquire about the pravastatin prescription(s) you dispensed.

You may also call International Laboratories at 727-322-7146 (8:00 AM – 5:00 PM EST, Monday through Friday) for more information about this recall.

### Pravastatin Recalled by International Laboratories

<table>
<thead>
<tr>
<th>Product Description</th>
<th>NDC #</th>
<th>Lot #</th>
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<tbody>
<tr>
<td>Pravastatin sodium 40 mg in bottles of 30 tablets</td>
<td>54458-925-16</td>
<td>115698A</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 International Laboratories:
(727) 322-7146 (8:00 a.m. – 5:00 p.m. EST, Monday through Friday)

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

Please distribute immediately.