The Board of Pharmacy has received notice of the following product recall. The Board strongly encourages pharmacies to immediately review their quality assurance and recall policies and procedures to determine if any corrective action is required.

<table>
<thead>
<tr>
<th>Description</th>
<th>Lot #</th>
<th>Exp Date</th>
<th>NDC</th>
<th>UPC</th>
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</thead>
<tbody>
<tr>
<td>ARTIFICIAL TEAR OINT AKOR 3.5GM</td>
<td>9K82B</td>
<td>09/30/2022</td>
<td>17478006235</td>
<td>31747806235</td>
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<td></td>
<td>9K82A</td>
<td>09/30/2022</td>
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<td></td>
<td>9H32A</td>
<td>07/31/2022</td>
<td></td>
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<tr>
<td></td>
<td>9G01A</td>
<td>06/30/2022</td>
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</tr>
</tbody>
</table>

Akorn is recalling the above item/lots due to a confirmed out of specification identified during sterility testing observed during 12-month controlled room temperature stability testing for the annual stability lot; Akorn has decided in an abundance of caution to recall all lots manufactured in 2019. This recall is to the retail level. Affected product started shipping October 22, 2019.