



Phone: (614) 492-8177  
Toll Free Phone: (800) 707-4621

**URGENT PRODUCT RECALL**  
**UPDATED – RETAIL RECALL**  
Converted from market withdrawal to recall

February 4, 2026

Dear Valued Customer,

American Health Packaging is initiating a voluntary recall to the **RETAIL LEVEL** for **AHP Oxycodone Hydrochloride Tablets (CII), 5 mg, 100 UD; Carton NDC#: 68084-354-01, (Individual Dose NDC: 68084-354-11)**, for the lots listed below:

Product Description	AHP Lot No.	Expiration Date	Ship Dates of Product
<b>AHP Oxycodone Hydrochloride Tablets (CII) 5 mg, 100 UD</b>	1027932	06/30/2027	12/15/2025 to 12/18/2025
<b>Carton NDC#: 68084-354-01 (Individual Dose NDC: 68084-354-11)</b>	1028360	08/31/2027	12/18/2025 to 01/05/2026
<b>REASON</b>	American Health Packaging has received multiple customer complaints for card seal defects observed on the subject lots. Customers reported card seal defects (weak/non-existent seals), leading to the tablets falling out of their cavities.		
<b>PRODUCT INDICATION</b>	Oxycodone hydrochloride is indicated for the management of pain severe enough to require an opioid analgesic and for which alternative treatments are inadequate.		
<b>ACTIONS REQUIRED</b>			
<ol style="list-style-type: none"> <li><b>Distributors/Pharmacies</b> - Immediately examine your inventory, quarantine and discontinue distribution of this lot.</li> <li><b>Distributors</b> - <u>Complete the enclosed Business Reply Card even if you do not have any product on hand.</u></li> <li><b>Distributors</b> – Please pass this Recall Notice on <b>ONLY</b> to pharmacies that received this product lot.</li> <li><b>Pharmacies</b> – If you have units of the affected products/lot in inventory, <b>please contact Sedgwick at (855) 215-5142</b> to receive a notification package with the Business Reply Card and return instructions.</li> <li>Business Reply Cards can be submitted by any of these methods.  <b>Fax:</b> (855) 207-2756  <b>Email:</b> <a href="mailto:ahp3655@sedgwick.com">ahp3655@sedgwick.com</a>  <b>Mail:</b> 2670 Executive Dr., Ste. A, Indianapolis, IN 46241  <b>a) Once Business Reply Cards have been received by Sedgwick, a Return Kit, which will include the required DEA 222 Form, will be mailed to your facility.</b> </li> <li><b>Distributors/Pharmacies</b> – Return recalled product to Sedgwick as instructed in recall/return packet.</li> <li><b>Pharmacies</b> - You do not need to contact any patients.</li> </ol>			
<b>OTHER</b>	This recall extends to the Retail Level only. No other lots, packages, or formulations are affected. Please reorder stock immediately. For questions about the recall process, call Sedgwick at (855) 215-5142. This recall is being conducted with the knowledge of the Food and Drug Administration. We appreciate your immediate attention and cooperation and sincerely regret any inconvenience caused by this action.		

**To receive credit, the Business Reply Card and recalled product must be returned to Sedgwick by May 31, 2026. All reimbursements will be provided once recalled product is returned to and processed by Sedgwick.**

Thank you for your support in complying with the requests in this letter. We apologize for the inconvenience that this incident may cause you or your customers.

Sincerely,

Becky Mahon (Jan 30, 2026 17:47:22 EST)

Becky A Mahon  
Sr. Manager, Regulatory Affairs and Stability  
American Health Packaging