



# California Drug Recall Information



## Recall Name

**Qualitest Recalls Hydrocodone Bitartrate and Acetaminophen Tablets, USP 10mg/500mg Due to the Potential for Oversized Tablets**

Recall Date	Product Description	Recalling Firm	Recall Reason
9/10/12	Hydrocodone Bitartrate and Acetaminophen Tablets, USP 10mg/500mg  NDC# 0603-3888-21	<b>Qualitest Pharmaceuticals, Inc.</b> Huntsville, AL	<i>Potential for Oversized Tablets</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	Hydrocodone Bitartrate and Acetaminophen Tablets, USP 10mg/500mg, 100 count  Suspect Lot: <ul style="list-style-type: none"> <li>C1440512A; Expiry date 12/13</li> </ul> Lot number can be found on the side of the bottle	<b>CA</b> , nationwide	Distributed between May 14 and August 3, 2012

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/Safety/Recalls/ucm318827.htm>