PROVIDER ALERT

Levothyroxine and Liothyronine (Thyroid Tablets, USP) Drug Recall Notification

To:	Alliance Contracted Pharmacies	**Recall Class:**	*Not yet classified*
From:	Alliance Pharmacy Services	**Recall Issue Date:**	08/09/2018
Date:	10/08/2018	**Recall #:**	*Not yet classified*
Subject:	**DRUG RECALL NOTICE**	**Manufacturer**:	Westminster Pharmaceuticals, LLC.
Products:	Medi-Cal, Group Care	**Reason:**	Risk of Adulteration

DRUG AFFECTED	NDC	LOT #	Action to take
Levothyroxine and Liothyronine (Thyroid Tablets, USP) 15mg X 100ct	69367-159-04	15918VP03 15918VP02 15918VP01 15918007 15918006 15918005 15918004 15918003 15918002 15918001 15917VP03 15917VP02 15917VP01	Remove from Distribution and return to manufacturer
Levothyroxine and Liothyronine (Thyroid Tablets, USP) 30mg X 100ct	69367-155-04	15517VP01 15517VP02 15517VP03 15518001 15518002	Remove from Distribution and return to manufacturer
Levothyroxine and Liothyronine (Thyroid Tablets, USP) 60mg X 100ct	69367-156-04	15617VP05 15617VP04 15617VP03 15617VP01 15617VP01	Remove from Distribution and return to manufacturer

Questions? Please call the Alliance Pharmacy Services Department
Monday – Friday, 8 am – 5 pm
Phone Number: **510.747.4541**
www.alamedaalliance.org

Levothyroxine and Liothyronine (Thyroid Tablets, USP) 90mg X 100ct	69367-157-04	15717VP-01 15717VP-02 15717VP-03 15718004 15717002	Remove from Distribution and return to manufacturer
Levothyroxine and Liothyronine (Thyroid Tablets, USP) 120mg X 100ct	69367-158-04	15817VP-01 15817VP-02 15817VP-03	Remove from Distribution and return to manufacturer

Source: **www.fda.gov/Safety/Recalls/ucm616601.htm**

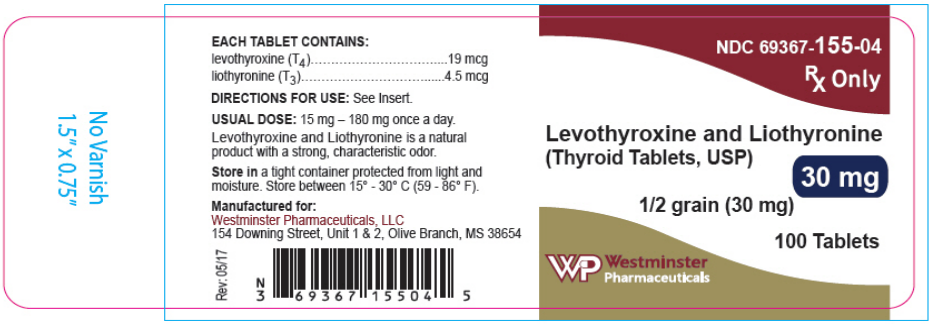

Image above reflects example of recalled drug.

Westminster is notifying its direct accounts by email and by phone to immediately discontinue distribution of the product being recalled and to notify their sub-wholesale accounts of this product recall and make arrangements for impacted product to be returned to Westminster. Instructions for returning recalled products are given in the Recall Notice Letter and Recall Response Form. **Consumers that have these products which are being recalled should not discontinue use before contacting their physician for further guidance.**

Customers and patients with medical-related questions, information about an adverse event or other questions about the Westminster's product's being recalled should contact Westminster's Regulatory Affairs Department.

Monday-Friday, 9 am – 5 pm EST
Voicemail is available 24 hours/day, 7 days/week
Toll-Free: **888.354.9939**
Email: **recalls@wprx.com**

Adverse reactions or quality problems experienced with the use of this product may be reported to the FDA's MedWatch Adverse Event Reporting Program online, by regular mail or by fax.

- **Online**: Complete and submit the report **www.fda.gov/medwatch/report.htm**
- **Regular Mail or Fax**:
  - Download form **www.fda.gov/MedWatch/getforms.htm**
  - To request a reporting form, please call toll-free at **1.800.332.1088**

  Please complete and return to the address on the pre-addressed form, or submit by fax to **1.800.FDA.0178** (**1.800.332.0178**)