April 17, 2020 **COM-2020-031**

RECALL NOTIFICATION

DATE:
April 15, 2020

DRUG NAME:
Clopidogrel Tablets USP, 75 mg

COMPANY:
International Laboratories, LLC

REASON:
Mislabeling (May contain Clopidogrel 75mg or Simvastatin 10mg)

Dear provider of pharmaceutical services,

At PharmPix we are committed to the health and well-being of patients. The clinical team wants to communicate you with the latest up-to-date information on the Food and Drug Administration (FDA) drug recalls.

Affected product:

NDC	Product description	Lot #
54458-0888-16	**Clopidogrel Tablets, USP 75 mg 30 count**	117099A

Pharmacy required action:

- Identify if they have the product in inventory and immediately stop using and dispensing it.
- Contact all members that in the previous 90 days received the recalled medication and advise them to stop using, return the drug, and contact their physician or healthcare provider if they have experienced any problems that may be related to taking or using the recalled drug product.

Contact information:

- Consumers: Contact Inmar at 855-258-7280 or via email internationallabs@inmar.com or by using mailing address Recall Coordinator 635 Vine St. Winston Salem, NC 27101 Inmar's business hours are (Monday – Friday 9 AM – 5 PM EST).

Note:
Remember that any adverse event related to this or any other pharmaceutical product can be reported to the FDA's MedWatch

Adverse Event Reporting program:

- Online: www.fda.gov/medwatch/report.htm

- Regular Mail or Fax: Download form www.fda.gov/MedWatch/getforms.htm or Call 1-800-332-1088 to request a reporting form, then complete and return to the address on the pre-addressed form, or submit by fax to 1-800-FDA-0178

Additional information:

- **https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/international-laboratories-llc-issues-voluntary-nationwide-recall-one-1-lot-clopidogrel-tablets-usp**

Best regards,

PharmPix Clinical Department

Reference:

- Center for Drug Evaluation and Research. (2020, April 15). International Laboratories, LLC Issues Voluntary Nationwide Recall of One (1) Lot of Clopidogrel Tablets USP, 75 mg Packaged in Bottles of 30 Tablets Due to Mislabeling NDC # 54458-888-16; Lot # 117099A. Retrieved April 17, 2020, from https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/international-laboratories-llc-issues-voluntary-nationwide-recall-one-1-lot-clopidogrel-tablets-usp

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