May 21, 2018

COM-2018-009

Dear provider of pharmaceutical services,

Recall: Metoprolol succinate extended-release tablets

At PharmPix we are committed to the health and well-being of patients. It is for this reason that we inform that on May 16, 2018 the US Food and Drug Administration (FDA) issued a statement notifying the voluntary retirement of more than 8,160 bottles of metoprolol succinate extended-release tablets, which are manufactured by Dr. Reddy's Laboratories. The recall was made after a 75mg clopidogrel tablet was found in one of the bottles and affects 100 tablets bottles. The FDA designated it a Class II recall. The affected lot is detailed on Table 1.

Table 1. Affected lot of metoprolol succinate extended-release

Product	Affected Lot	Expiration date	NDC
Metoprolol succinate extended-release	C706254	08/2019	55111-468-01

The Pharmacy must:

1. Identify if they have the product in inventory and immediately stop using and dispensing the product.
2. Contact all members that received the recalled medication in the previous 90 days.
3. Contact Dr. Reddy's Laboratories for questions regarding this recall.

For additional information visit:

Managed Care Health website at:
https://www.managedhealthcareconnect.com/content/foreign-tablet-prompts-beta-blocker-recall

Department of Clinical Pharmacy

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