September 4,2018

COM-2018-018

Dear provider of pharmaceutical services,

Recall: Montelukast tablets by Camber Pharmaceuticals

At PharmPix we are committed to the health and well-being of patients. It is for this reason that we inform that on August 31, 2018 the US Food and Drug Administration (FDA) issued a statement notifying voluntary retirement of Montelukast Sodium Tablets manufactured by Camber Pharmaceuticals, Inc. The recall was made due incorrect drug in bottles. Sealed bottles labeled as Montelukast Sodium Tablets, 10 milligrams, 30 count bottles from Camber were found to instead contain 90 tablets of Losartan Potassium Tablets, 50 mg. The affected lot is detailed on Table 1.

Table 1. Affected lot of Montelukast Sodium Tablets 10mg

Product	Affected Lot	Expiration date	NDC
Montelukast Sodium Tablets 10mg	MON17384	12/31/2019	31722-726-30

The Pharmacy must:

1. Identify if they have the product in inventory and immediately stop using and dispensing the product.
2. It is important that pharmacy contact all members that in the previous 90 days received the recalled medication.
3. For questions regarding this recall can access www.fda.gov.

For additional information visit:
https://www.fda.gov/NewsEvents/Newsroom/PressAnnouncements/ucm619174.htm

Department of Clinical Pharmacy

CHANGING THE WAY PBM's WORK, NOW AND FOREVER