Voluntary Recall: Diocto Liquid and Diocto Syrup by Rubby Laboratories: Possible Product Contamination

At PharmPix we are committed to the health and well-being of patients. It is for this reason that we inform you that on August 3, 2017 the FDA issued a statement notifying the voluntary retirement of several batches of the Product Diocto Liquid and Diocto Syrup, (docusate sodium solutions) manufactured by PharmaTech, LLC due to a risk of product contamination with Burkholderia cepacia. The use of this contaminated product could result in serious infection or even life threatening, especially in patient with compromised immune system and in patients with chronic lung condition such a cystic fibrosis. The affected lots are detailed in the following table.

Product	Affected Lot	NDC
Diocto Liquid and Diocto Syrup (473ml)	All lots	0536-0590-85
Diocto Liquid and Diocto Syrup) (473ml)	All lots	0536-1001-85

The Pharmacy must:

1. Identify if they have the product in inventory and immediately stop using and dispensing the product.
2. For questions regarding this recall can contact Rugby laboratories.

For additional information visit:

https://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm570014.htm

Department of Clinical Pharmacy

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