VistaPharm

January 18, 2017

URGENT VOLUNTARY DRUG RECALL

Dear Valued Customer:

VistaPharm, Inc. is informing you of a voluntary recall for the products listed. The list of products is included in Attachment 1 of this letter.

See enclosed product labeling (Attachment 2) for each of the affected products for ease in identifying the product at the retail level.

These lots are being recalled because the purified water used to manufacture the drug products may have been contaminated with the bacteria, Burkholderia cepacia. Adverse events following the ingestion of a contaminated product could range from no symptoms to potentially life-threatening symptoms. A "normal" healthy person is at less risk of developing any adverse health consequences than is a person with an underlying health condition, such as a patient whose immune system is compromised, a cystic fibrosis patient, or a child with an immature immune system. The products were distributed between 03/2015 and 06/2016. To date, there have been no illnesses associated with the use of these drug products and the testing of the drug products complies with all VistaPharm specifications and during finished goods release testing, the bacteria was not detected.

Please immediately examine your inventory and quarantine product subject to the recall. In addition, if you may have further distributed this product, please identify your customers and notify them at once of this product recall. Your notification to your customers may be enhanced by including a copy of this recall notification letter.

This recall is being carried out to the retail level. Your assistance is appreciated and necessary to prevent consumer illness.

In our best efforts to coordinate the prompt and secure return of the affected product, VistaPharm, Inc. has contracted with PharmLink Inc. to provide reverse distribution support for this recall.

Shipping Address:
8285 Bryan Dairy Road, #160
Largo, FL 33777-1350

SHIPPING DOCK: #21

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