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October 28, 2011

URGENT: Drug Recall for XIGRIS [drotrecogin alfa (activated)]

Dear ASD Healthcare Customer,

Eli Lilly and Company is initiating withdrawal of Xigris, 5 mg and 20 mg vials [drotrecogin alfa (activated)] following discussions with the U.S. Food and Drug Administration. Lilly is initiating this withdrawal due to new clinical trial findings of lack of efficacy, which call into question the benefit-risk profile of the product. While there were no new safety findings, the study showed no 28-day survival benefit of Xigris in septic shock patients.

The withdrawal extends to all distributed Xigris lots that remain within expiration date. This represents a distribution period from 5 May 2009 to 25 October 2011.

What You Should Do:

1. Check your inventory for the withdrawn product.

2. If the withdrawn product is found, stop dispensing or using the product.

3. Return all withdrawn product to the supplier (wholesaler/distributor) from whom you purchased. Your supplier will provide you with a credit.

4. Your supplier will return the product to Eli Lilly and Company.

5. Eli Lilly and Company will provide credit to your supplier.

Please see the formal recall letter from Eli Lilly and Company for full details.

Call ASD Healthcare at 800.746.6273 or asd-returnsmail@asdhealthcare.com to inquire about returning this product.

Sincerely,

ASD Healthcare