

DATE: May 17, 2012

RC: 12-092

RECEIVED MAY 21 2012



****AUTO**MIXED AADC 750 4 501
00002540 12-092 ATTN: PHARMACY
HEMOPHILIA OF GEORGIA
8800 ROSWELL RD
ATLANTA, GA 30350-1826

IMPORTANT NOTICE

Urgent Recall

URGENT!!!

DRUG RECALL!!!

URGENT!!!

FDA/SUPPLIER CLASS OF RECALL: Not Yet Classified
LEVEL OF NOTIFICATION: McKesson Customer
SUPPLIER: Vygon - #00432

Product Description	Lot #	Exp Date	NDC	UPC	Econo #
SYR HEP 10UN 3ML FILL 10ML 240	87-133-9D 03/2012, 00-015-9D 02/2013		63807056031	36380756031	1942655
SYR HEP 10UN 5ML FILL 10ML 240	97-053-9D 01/2013, 94-044-9D 09/2012, 93-063-9D 09/2012, 91-059-9D 07/2012, 90-048-9D 06/2012, 97-071-9D 01/2013		63807056051	36380756051	1943026
SYR HEP 100UN 3ML FIL 10ML 240	01-062-9D 05/2013, 99-130-9D 03/2013, 01-005-9D 05/2013, 97-040-9D 01/2013, 91-036-9D 07/2012, 10-010-9D 02/2014, 07-046-9D 11/2013, 08-060-9D 12/2013		63807066031	36380766031	1941111
SYR HEP 100UN 5ML FIL 10ML 240	01-055-9D 05/2013, 09-009-9D 01/2014, 07-020-9D 11/2013, 07-033-9D 11/2013, 08-037-9D 12/2013, 09-007-9D 01/2014, 09-022-9D 01/2014, 10-001-9D 02/2014, 94-045-9D 01/2012, 95-014-9D 05/2012, 95-109-9D 06/2012, 08-048-9D 12/2013		63807066051	36380766051	1941756

Vygon is conducting a voluntary recall for the above products/lots following testing of retention samples for potency. Some lots failed Anti-factor IIa assay for potency. Affected product started shipping July 2010. This recall is to the McKesson Customer Level.

McKesson Customers should immediately examine your inventory for the affected items/lots completely and stop dispensing immediately. If you have the affected product, using your Telxon, McKesson Connect, SMO, Econolink, Closed Loop Distribution, Mobile Manager or Telestock, please transmit a request for return authorization as unsaleable, manufacturer recall, and return within 30 days to your McKesson servicing distribution center for credit.

PLEASE NOTE: Customers currently participating in a McKesson administered Return to Vendor program should return this product to their designated returns processor. All others should follow the instructions provided by the manufacturer.

This recall is being conducted with the knowledge of the FDA.

Information contained in this document was provided by Vygon.