Teva Pharmaceuticals USA, Inc.

URGENT DRUG RECALL – RETAIL LEVEL - INITIATED 5/11/15

Zebeta® (bisoprolol fumarate), 10mg, Tablets

RECALLED BY:

Teva Pharmaceuticals USA, Inc. Horsham, PA 19044

Lot#	Exp. Date	Strength	Bottle Size	NDC
34019710A	5/2017	10mg	30 count	51285-061-01

Dear Customer:

Teva Pharmaceuticals USA, Inc. is voluntarily recalling the above lot of Zebeta® (bisoprolol fumarate), 10mg, Tablets distributed under the Duramed Pharmaceuticals, Inc. label. This recall is being carried out to the RETAIL LEVEL due to an out of specification dissolution test result during stability testing. The use of or exposure to the product may cause temporary or medically reversible adverse health consequences, or the probability of serious adverse health consequences is remote.

Wholesalers / Distributors / Retailers - Please perform the following activities:

- Examine your inventory immediately for the specified lot of Zebeta® (bisoprolol fumarate), 10mg, Tablets.
- Our records indicate we shipped this product between September 20, 2014 and October 29, 2014.
- Immediately discontinue distribution of the specific lot being recalled.
- Please perform a SUB-RECALL to your RETAIL accounts using this Recall Notification and Stock Response Form.
- Promptly complete the attached recall stock response and reply via fax number 817-868-5362 or mail, even if you have **no** product to return.

Completed Recall Stock Response forms can be mailed, emailed, or sent via FAX to Inmar Attn: Recall Coordinator, 4332 Empire Road Suite 200, Fort Worth, TX 76155. Inmar Email address: recallnotice@inmar.com. Inmar FAX: 817-868-5362. Inmar will send a Return Goods Authorization label and shipping label. Appropriate credit for product returns, plus handling and shipping expenses, will be issued upon receipt of said product with the Return Goods Authorization form. All recalled product returned without a Return Goods Authorization label may delay the issuance of your credit.

This recall is being made with the knowledge of the Food & Drug Administration. Your cooperation and prompt response to this notice is appreciated. If you have Customer Service related questions, please contact Teva Customer Service at 800-545-8800. For medical-related questions please contact Medical Information at 888-838-2872, option 3, then option 4. If you need a Recall Stock Response form, contact Inmar at 800-967-5952 or acquire it from clsnetlink.com.

Sincerely,

James S. Young

Director, Quality Systems & Regulatory Compliance

Teva Pharmaceuticals USA, Inc.

Teva Pharmaceuticals USA, Inc.

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STOCK RESPONSE FORM

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RECT CUSTO	MERS ONLY: D	oes this respons	e include <u>all</u> DC loca	ations? YES	NO	
stomer/Store Na	me:			DEA	#:	
EA # is required	; if not provided i	the processing of	f your form will be de	elayed		
dress:						
			State:Zip:			
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Lot #	Exp. Date	Strength	Bottle Size	NDC	Quantity to Return (count partial as 1)	
34019710A	5/2017	10mg	30 count	51285-061-01		
ON DIRECT CU		LY: Please com	plete the following:	DEA #:_		
				directed to the follow		
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Recall S	_	-		d between 7-10 busines do not resubmit respon	ss days contact Inmar at nse form.	
		•		Customer Service at 80 on 888-838-2872, option		
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1	_	this form to: 81	7-868-5362 or E-ma	ail at: <u>recallnotice@</u> in	mar.com	

Store

Kit

D.B

Inmar/MedTurn Use Only:

Labels

Scan