



Innovative Pharmaceuticals Offering Therapeutic Excellence

February 3, 2009

URGENT PRODUCT RECALL

Dear Recall Coordinator:

Product:

Ther-Rx Corporation is voluntarily recalling all lots of Chromagen[®], Chromagen[®] FA, Chromagen[®] Forte, Encora[®], Niferex[®] Gold, Niferex[®] 150 Forte Capsule, PrimaCare[®], PrimaCare[®] ONE, PrimaCare[®] Advantage[™], PreCare Premier[®], PreCare[®] Chewable, PremesisRx[®], PreCare Conceive[®] and Repliva 21/7[®] because they may potentially have been manufactured under conditions that did not sufficiently comply with cGMP's. This recall involves the following NDC's only:

Chromagen[®] caplet, NDC 64011-0198-26

Chromagen[®] FA caplet, NDC 64011-0199-26

Chromagen[®] Forte caplet, NDC 64011-0197-26

Encora[®] capsule/tablet, NDC 64011-0166-36

Niferex[®] Gold tablet, NDC 64011-0162-26

Niferex[®] 150 Forte Capsule, NDC 64011-0164-26

PrimaCare[®] capsule/tablet, NDC 64011-0204-28

PrimaCare[®] ONE capsule, NDC 64011-0200-19

PrimaCare[®] ONE capsule, NDC 64011-0218-19

PrimaCare[®] Advantage[™] capsule/tablet, NDC 64011-0230-28

PreCare Premier[®] tablet, NDC 64011-0195-19

PreCare[®] Chewable tablet, NDC 64011-0024-19

PremesisRx[®] tablet, NDC 64011-0019-19

PreCare Conceive[®] tablet, NDC 64011-0014-19

Repliva 21/7[®] tablet, NDC 64011-0207-34



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Level:

This recall is being carried out to the wholesale level.

This recall is being made with the knowledge of the Food and Drug Administration.

Action for Distributor:

Distributor

- Stop distributing the affected lots. **Quarantine product immediately.**
- Please perform a physical count and record this data on the response letter which is included.
- Send back the response letter even if you do not have any product to:
Ther-Rx Corporation
Email: info@ther-rx.com or fax to **(314) 646-3701**
- Notifications of this recall are being sent to direct wholesale customers of Ther-Rx Corporation. For customers who redistribute the products to other wholesale customers, please notify them to contact Capital Returns 1-888-851-3502 to receive recall product return packet and instructions.
- Recalled product must be returned to:

**Capital Returns
6101 North 64th Street
Milwaukee, WI 53218
1-888-851-3502**

Please mark all returned containers "RECALL" and a numbered debit memo must be included.

A copy of the debit memo must be sent to the following address:

Ther-Rx Corporation
Attn: Tammy Butler
One Corporate Woods Drive
Bridgeton, MO 63044

Do not return product until you have contacted Capital Returns for a recall product return packet instructions.

You will be credited for the returned product, the appropriate processing fees and shipping costs associated with this recall.

Any customer inquiries related to this action should be addressed to Capital Returns Customer Service at Phone 1-888-851-3502, with representative's available Monday through Friday, 8 am to 6 pm EST.

Sincerely,

A handwritten signature in black ink that reads "David Kaiser".

David Kaiser
Senior QA Manager, Compliance
Recall Coordinator



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PRODUCT RECALL – Response Letter

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We have reviewed our inventory (check one):

- No units were detected in our inventory
- Yes, one or more units were detected in our inventory

Name: _____ Title: _____

Company: _____ DEA License: _____

Address: _____ City, State: _____

Phone #: _____ Date: _____

Please fax to Ther-Rx 314-646-3701 or e-mail to Ther-Rx Customer Service at info@ther-rx.com.