



February 4, 2009

Medication Recall Alert

MEDICATIONS:	Multiple Medications Manufactured by Ther-Rx Corporation
DOSAGE FORMS:	Multiple, See Below
NDC/LOT NUMBERS:	All NDC and Lot Numbers
ISSUE:	Ther-Rx Corporation's nationwide recall has been expanded to include various prescription prenatal vitamins and iron supplement products.

The following was posted on the FDA Med Watch Web site on February 4, 2009:

"Ther-Rx Corporation, a subsidiary of KV Pharmaceutical Company is issuing a voluntary nationwide recall of prescription prenatal and iron supplement nutritionals as identified below (all lots within their expiration dates) at a wholesale level. The Company is taking this action as a precautionary measure, because the products may have been manufactured under conditions that did not sufficiently comply with current Good Manufacturing Practice (cGMP).

Patients who may have these products in their possession should continue to take them in accordance with their prescriptions, as the risk of suddenly stopping needed medications may place patients at risk. Patients should contact their physician or healthcare provider if they have experienced any problems that may be related to taking or using these products, or to obtain replacement medications or prescriptions.

Products Recalled to the Wholesale Level:

Prescription Iron Supplement Products:

Chromagen® Caplet
Chromagen® FA Caplet
Chromagen® Forte Caplet
Encora® Capsule
Niferex® Gold Tablet
Niferex® 150 Forte Capsule
Repliva 21/7® Tablet

Prescription Prenatal Vitamin Products:

PreCare® Chewable Tablet
PreCare® Conceive Tablet
PreCare Premier® Tablet
PremesisRx® Tablet
PrimaCare® Capsule/Tablet
PrimaCare® Advantage™ Capsule/Tablet
PrimaCare® ONE Capsule

This recall is being conducted with the knowledge of the U.S. Food and Drug Administration (FDA). At this time, the Company is unable to determine when distribution of these products will resume.

Patients with questions about the recall should call the telephone number above, or contact their healthcare providers.”

A complete copy of the accompanying “Recall—Firm Press Release” dated February 3, 2009 from the FDA and Ther-Rx Corporation may be viewed at:

http://www.fda.gov/oc/po/firmrecalls/therrx02_09.html