



February 4, 2009

Medication Recall Alert

MEDICATIONS: Multiple Medications Manufactured by Ethex Corporation

DOSAGE FORMS: Multiple, See Below

NDC/LOT NUMBERS: All NDC and Lot Numbers

ISSUE: Ethex 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:

"ETHEX Corporation, a subsidiary of KV Pharmaceutical Company (NYSE: KVa/KVb), is issuing a voluntary nationwide recall of the products 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/Distributor Level:

Prescription Prenatal Vitamin Products:

Advanced NatalCare® Tablets
Advanced-RF NatalCare® Tablets
Cal-Nate™ Tablets
CareNatal™ DHA Tablets
ComBgen Tablets
ComBiRx™ Tablets
NataCaps Capsules
NatalCare Gloss Tablets
NatalCare PIC Tablets
NatalCare PIC Forte Tablets
NatalCare Plus Tablets
NatalCare Rx Tablets
NatalCare Three Tablets
NataTab FA Tablets
NataTab RX Tablets
NutriNate® Chewable Tablets
NutriSpire™ Tablets
Prenatal MR 90 FE Tablets
Prenatal MTR w/Selenium Tablets

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Prenatal Rx 1 Tablets
Prenatal Z, Advanced Formula Tablets
Ultra NatalCare Tablets

Prescription Iron Supplement Products:

Anemagen Caplets
Anemagen Forte Caplets
Conison™ Capsules
Fe-Tinic™ 150 Forte Capsules

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 Ethex Corporation may be viewed at:

http://www.fda.gov/oc/po/firmrecalls/ethex02_09.html