The Board of Pharmacy has received notice of the following product withdrawal. The Board strongly encourages pharmacies to immediately review their quality assurance and recall policies and procedures to determine if any corrective action is required.

Product: Injectafer® (ferric carboxymaltose injection)

NDC No.: 0517-0650-01

Strength: 750 mg/ 15 mL (50 mg/mL) per vial

Package Size: 15 mL/ vial

Manufactured By: American Regent Inc.

Lot No: 20131L0C2

Withdrawal of the above-mentioned lot for Injectafer® (ferric carboxymaltose injection) was initiated due to a trend in Customer Product Quality Complaints related to Lot No. 20131L0C2. This withdrawal is being conducted to the **USER LEVEL.**