

Agency News Flash

August 3, 2015

Product Recall

Dear Clients:

Purpose of this communication

We are writing to inform you that effective immediately, the FDA has issued a voluntary recall of 0.9% Sodium Chloride Injection, USP (AUTO-C) by Baxter due to the potential for leaking containers, particulate matter and missing port protectors.

Why are we doing this?

- Product Recall – For additional information, please visit the following website:
- http://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm456793.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery
- EFFECTIVE IMMEDIATELY

What do I need to do?

- ❖ It is our responsibility as a provider to notify our clients and the public to facilitate the repair, replacement and/or resolution of the recall according to the guidelines issued by the manufacturer in the FDA notification
- ❖ Please notify our agency's Quality Assurance Department if you are affected by this recall by faxing information to: 678-964-2217
Attn: Hilreth Brown, RN, BSN

Thank you for your prompt attention and cooperation.

CONYERS OFFICE: 1804 Overlake Drive, Conyers, GA 30013

MACON OFFICE: 2012 Riverside Dr. Macon, GA 31204

Phone: 678-964-2026 Fax: 678-964-2217

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