carecentrix

Provider Newsflash
March 2024

FDA Recall – Eugia US, LLC

Purpose of this communication:

- We are writing to inform you that effective immediately the FDA has issued notice of a voluntary recall initiated by Eugia US, LLC (f/k/a AuroMedics Pharma, LLC) of lot number 3MC23011, Exp. date November, 2026 of Methocarbamol Injection, USP 1000 mg/10 mL (100mg/mL) single dose 10 mg vial due to the presence of white particles floating inside of the vial. The entire lot was shipped to wholesalers nationwide from January 12, 2024 through January 16, 2024. If the particulate matter reaches the blood vessels or is injected intravascularly it can travel to various organs and block blood vessels in the heart, lungs or brain which can cause stroke and even lead to death.

What do I need to do?

- Please review the following recall notice: Eugia US LLC (f/k/a AuroMedics Pharma LLC) Issues Voluntary Nationwide Recall of Methocarbamol Injection, USP 1000 mg/10 mL (100mg/mL) (Single Dose Vial) Due to Presence of White Particles | FDA
- Notify impacted patients and facilitate the replacement and/or resolution of the recall, according to the guidelines issued by the manufacturer in the FDA notification.
- Confirm if you have any CareCentrix patients affected by this recall and notify the CareCentrix Quality Department by faxing information to: 919-792-6718 Attn: Quality Department.

Thank you in advance for your cooperation and continued partnership.