carecentrix

Provider Newsflash
May 2023

FDA Recall – Akorn Drug Products

Purpose of this communication:

- We are writing to inform you that effective immediately the FDA has posted notice of a voluntary nationwide recall by Akorn Operating Company, LLC of various human and animal drug products within expiration date due to the company shut down. As a result of a bankruptcy, the firm will no longer have a quality program to support or guarantee that products will all intended specifications through the labeled shelf life of the product.

What do I need to do?

- Please review the following recall notice: https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/update-akorn-issues-voluntary-nationwide-recall-various-human-and-animal-drug-products-within-expiry?utm_medium=email&utm_source=govdelivery
- 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.