carecentrix

Provider Newsflash January 17, 2020

FDA RECALL

Purpose of this communication:

We are writing to inform you that effective immediately the FDA has issued notice of a voluntary recall of all unused sterile drug products within expiration date by AXIA Pharmaceutical due to a lack of sterility assurance.

What do I need to do?

- Please review the following recall notice:
- http://s2027422842.t.en25.com/e/es?s=2027422842&e=294226&elqTrackId=376c7bc788024cd5a73d955f2e3dcbdc&elq=f214831d030c4e5b852f0574f6b6009c&elqaid=10935&elqat=1
- Notify impacted patients and facilitate the repair, 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 for your prompt attention and cooperation in this matter.