carecentrix

Provider Newsflash March 26, 2021

FDA RECALL

Purpose of this communication:

We are writing to inform you that effective immediately the FDA has posted notice of a voluntary recall issued by Zydus Pharmaceuticals, Inc. of 4 lots of their Acyclovir Sodium Injection 50 mg/ml, 10 ml and 20 ml vials after receiving several complaints of crystallization in the vials. Administration of crystalized Acyclovir Sodium Injection may cause life-threatening adverse consequences.

What do I need to do?

- Please review the following recall notice:
- https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/zydus-pharmaceuticals-usa-inc-issues-voluntary-nationwide-recall-acyclovir-sodium-injection-50-mgml?utm_medium=email&utm_source=govdelivery
- 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.