Provider Newsflash
October 2024

FDA Recall – Staska Pharmaceuticals, Inc.

Purpose of this communication:

- We are writing to inform you the FDA has published notice of a voluntary recall by Staska Pharmaceuticals, Inc. of lot number SP2400058 of Ascorbic Acid Solution for injection 500 mg/mL, 50 mL vials with an expiration date of 12/31/24 distributed nationwide between 7/31/24 and 8/2724. The recall is due to the presence glass particulates found in the one lot of vials used in the production of this batch.

What do I need to do?

- Please review the following recall notice: Staska Pharmaceuticals, Inc. Issues Voluntary Nationwide Recall of Ascorbic Acid Solution for Injection (Preservative Free, Non-Corn) 500mg/mL Due to the Presence of Glass 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.

EDRC# 1604. 092021