

FDA Recall – Cardinal Health

Purpose of this communication:

- We are writing to inform you that effective immediately the FDA has published a voluntary recall issued by Cardinal Health of their Covidien and Cardinal Health Urology and OR room specific kits and trays that contain Nurse Assist Products (sterile normal saline and sterile water) which were previously recalled for lack of sterility assurance.

What do I need to do?

- Please review the following recall notice: https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/cardinal-health-issues-medical-device-recall-nurse-assist-products-contained-within-kitstrays?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.