Provider Newsflash
June, 2021

FDA Recall

Purpose of this communication:

We are writing to inform you that effective immediately the FDA has issued notice of a Class 1 recall by Smiths Medical of specific models and lots of Jelco Hypodermic Needle-Pro Fixed Needle Insulin Syringes due to identification of skewed odd number line graduation markings on the syringe barrels. As a result of this issue, there is a potential for administration of an incorrect dose of insulin which could result in hyperglycemia or hypoglycemia and may lead to serious harm or death.

What do I need to do?

- Please review the following recall notice:
- https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/smiths-medical-issues-worldwide-notification-regarding-recall-jelcor-hypodermic-needle-pror-fixed?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 in advance for your cooperation and continued partnership.