Provider Newsflash

October, 2021

Company Announcement/Recall – MiniMed 600 Insulin Pumps

Purpose of this communication:

We are writing to inform you that the FDA has posted notice of an update to the Class I recall previously issued by Medtronic on November 21, 2019 of their MiniMed 600 Series Insulin Pumps. Medtronic has now expanded the recall to replace all of the MiniMed 600 series insulin pumps that contain the clear retainer rings. Retainer rings lock the insulin cartridge into place in the pump's reservoir compartment. If the ring is loose, damaged or missing, and the reservoir does not lock firmly into place, under or over delivery of insulin may occur. Medtronic is now replacing these pumps with a pump that has the updated black retainer ring even if the clear ring is not damaged and regardless of the warranty status of the pump.

What do I need to do?

- Please review the following recall notice: https://www.fda.gov/medical-devices/medical-device-recalls/medtronic-recalls-minimed-insulin-pumps-incorrect-insulin-dosing?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 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.