June 7, 2018
Web Announcement 1612

Hospira Issues Voluntary Nationwide Recall for Two Lots of Naloxone Hydrochloride Injection

Hospira, Inc., a Pfizer company, is voluntarily recalling lots **72680LL and 76510LL of Naloxone Hydrochloride Injection, USP, 0.4 mg/mL, 1 mL in 2.5 mL, Carpuject™ Single-use cartridge syringe system (NDC 0409-1782-69)**, to the hospital/institution level due to the potential presence of embedded and loose particulate matter on the syringe plunger.

Hospira, Inc. has notified wholesalers/distributors/hospitals to arrange for return of any recalled product. Distributors or retailers with an existing inventory of the lots, which are being recalled, should stop use and distribution, and quarantine immediately.

For additional assistance, call Stericycle at 1-800-805-3093 between the hours of 8 a.m. to 5 p.m. Eastern Time, Monday through Friday.

Healthcare professionals with questions regarding this recall can contact Pfizer using the following information.

Contact	Contact Information	Areas of Support
Pfizer Medical Information	1-800-438-1985, option 3 (8 a.m. to 7 p.m. Eastern Time Monday through Friday)	Medical inquiries
Pfizer Safety	1-800-438-1985, option 1 (24 hours a day 7 days per week)	To report adverse events or product complaints