



The affected products include all lots distributed February 16, 2017, to July 19, 2017, remaining within expiry, and they would be packaged in a syringe or IV bag. Administration of a drug product intended to be sterile that is not sterile could result in serious infections that may be life-threatening” FDA (2017).

FDA report “Cantrell Drug Company is voluntarily recalling all lots of unexpired sterile drug products to the hospital and user level due to lack of sterility assurance. The recalled products were distributed to health care facilities nationwide, except to the states of Connecticut, Hawaii, South Carolina and Vermont. Administration of a drug product intended to be sterile that is not sterile could result in serious infections that may be life-threatening.”

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