



Medtronic is informing patients worldwide of a voluntary recall of specific lots of infusion sets used with all models of Medtronic insulin pumps” FDA (2017).

FDA report “Medtronic is informing patients worldwide of a voluntary recall of specific lots of infusion sets used with all models of Medtronic insulin pumps. The recall is related to a certain discontinued component in these infusion sets and does not include insulin pumps or glucose sensors. The company determined, through recent field reports from patients and root cause analysis, that a component, the vent membrane, in the recalled infusion sets may be susceptible to being blocked by fluid during the process of priming/fill-tubing. This situation can lead to potential over-delivery of insulin shortly after an infusion set change, which may cause hypoglycemia.”

Full Alert

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