PRODUCT RECALLS

LifeScan, Inc., OneTouch Verio IQ Blood Glucose Meter – Recall Class I, 3/11/13

Meter will shut off and revert to "set up mode" at glucose values above 1023 mg/dL instead of displaying EXTREME HIGH GLUCOSE. Because diagnosis and treatment of extreme hyperglycemia may be delayed or incorrect treatment may be given, serious adverse health consequences, including death may occur. For complete information on the recall go to: http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm349127.htm

Abbott FreeStyle lnsulinx® Blood Glucose Meters – Voluntary Recall, 4/15/13.

The company determined that at extremely high blood glucose levels of 1024 mg/dL and above, the FreeStyle lnsulinx Meter will display and store in memory an incorrect test result that is 1024 mg/dL below the measured result. For example, at a blood glucose value of 1066 mg/dL, the meter will display and store a value of 42 mg/dL (1066 mg/dL - 1024 mg/dL = 42 mg/dL). For complete information go to: http://www.fda.gov/Safety/Recalls/ucm348391.htm

Fresenius Kabi USA notified health professionals of a voluntary recall of one lot – Lot 6103882 – of Magnesium Sulfate Injection, USP due to the potential presence of glass particles in the vials. The recalled product is labeled with Product Code 6450 and packaged as 500mg/mL strength in 50mL glass vials (25 vials per tray). The product was shipped in the United States between May 30, 2012 and June 6, 2012 and has an expiration date of October 31, 2014. For additional information go to: http://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm354329.htm?source=govdelivery