ICU Medical – Recall of Lactated Ringer's injection

- On May 8, 2020, ICU Medical announced a voluntary, consumer-level recall of one lot of Lactated Ringer's injection due to the presence of particulate matter identified as iron oxide.

- Lactated ringers injection was manufactured by Hospira for ICU Medical. It was distributed nationwide within the U.S. between September and October 2019.

Product Description	NDC#	Lot# (Expiration Date)
Lactated Ringer's injection, 1000 mL flexible container	0409-7953-09	07-514-FW2 (7/1/2010)

- Lactated Ringer's injection is indicated for parenteral replacement of extracellular losses of fluid and electrolytes, with or without minimal carbohydrate calories, as required by the clinical condition of the patient.

- Administration of a drug product that contains metal particulate matter could result in adverse events ranging from inflammation at the site of injection to more serious events that could include the formation of a blood clot obstructing the flow of blood which could lead to end-organ damage or death.

- To date, ICU Medical has not received any adverse event reports related to this recall.

- Patients who have the recalled Lactated Ringer's injection should stop use and contact their physician or healthcare provider if they have experienced any problems that may be related to using the recalled Lactated Ringer's injection.

- Anyone with an existing inventory of the recalled product should stop use and distribution and quarantine the product immediately.

- Contact ICU Medical by phone at **1-844-654-7780** for further information regarding this recall.