FDA report “Hospira, Inc. announced it is initiating a voluntary nationwide user-level recall of one lot of Lactated Ringers and 5% Dextrose Injection, USP, 1000 mL, Flexible Container, NDC 0409-7929-09, Lot 35-118-JT, Expiry 1NOV2015. This action is due to one confirmed customer report where particulate was identified within the solution of the primary container. The particulate was identified as a filamentous-like structured particulate indicative of mold. Analysis of the primary container and overwrap indicated a puncture in the same physical location, causing the primary container to leak.”

Click here for the full story.

Recall of Hospira Lactated Ringer’s and 5% Dextrose 1000L injection http://ctt.ec/bvcHk+
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Other intravenous and vascular access resources that may be of interest (External links – IVTEAM has no responsibility for content).