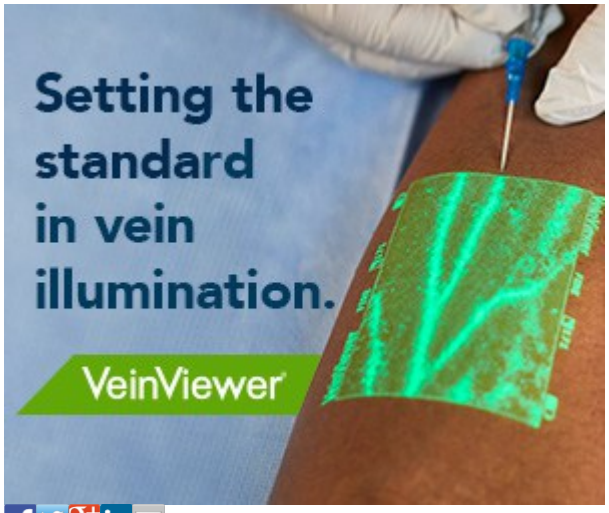

The FDA report “Baxter International Inc. initiated a recall in the United States of two lots of 0.9% Sodium Chloride Injection USP in 100 mL MINI-BAG PLUS Container to the hospital/user level. The recall is being initiated as a result of two complaints (one per lot) of particulate matter that was identified as a fragment of the frangible from the vial adapter. The issue was identified upon standard visual inspection prior to patient administration.”

Recall in the United States of two lots of 0.9% Sodium Chloride Injection http://ctt.ec/Va841+ @ivteam #ivteam

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