FDA report “Hospira, Inc. announced a voluntary nationwide user-level recall of one lot of Heparin Sodium, 1,000 USP Heparin Units/500 mL (2 USP Heparin Units/mL), in 0.9% Sodium Chloride Injection, 500 mL, NDC 0409-7620-03 Lot 41-046-JT with expiration date of 01NOV 2015. This action is due to one confirmed customer report of particulate in a single unit. The foreign particle was confirmed by Hospira as human hair, sealed between the tube and the film at the round seal of the unused Administrative Port on the non-print side of the container.”

Hospira announce voluntary recall of one lot of Heparin Sodium http://ctt.ec/AdV3f+ @ivteam #ivteam

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