FDA report “Baxter International Inc. announced it is voluntarily recalling two lots of INTRAVIA containers in the U.S. and Canada due to complaints received for particulate matter found inside the fluid path. Intravenous administration of a solution containing sterile particulate matter may lead to adverse health consequences. The extent and severity of harm depends on the size, number, and composition of the foreign material, and patient’s underlying medical condition.”

Full story click here

Voluntarily recall of two lots of INTRAVIA IV fluid containers http://ctt.ec/0QPau+ @ivteam #ivteam

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