



B. Braun recently identified an adverse quality trend in customer complaints reporting that some containers in lot J5J706 exhibited leakage and, in a few instances, visible particulate matter identified to be microbial growth” FDA (2016).

FDA report “B. Braun Medical Inc. is recalling one lot of 5% Dextrose Injection USP 100/150mL container (Lot #J5J706, catalog #S5104-5264, NDC 0264-1510-32) to the consumer level. B. Braun recently identified an adverse quality trend in customer complaints reporting that some containers in lot J5J706 exhibited leakage and, in a few instances, visible particulate matter identified to be microbial growth.

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A compromise of container integrity has the potential for leakage of the solution, usually identified prior to the use of the product. Leaking containers allow contamination of the solution, which can and has led to microbial contamination. Intravenous administration of a non-sterile product can result in serious infections that may be life-threatening.”

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