State of California—Health and Human Services Agency
California Department of Public Health

Food and Drug Branch – Drug Recalls

CALIFORNIA DRUG RECALL INFORMATION SHEET

Hospira Issues A Voluntary Nationwide Recall For One Lot of Propofol Injectable Emulsion (Containing Benzyl Alcohol)

Recall Date	Product Description	Recalling Firm	Recall Reason
08/22/2022	Propofol Injectable Emulsion	Hospira, Inc., a Pfizer Company	Potential presence of visible particulates

Recall Class	Product Identification	Distribution	Affected Dates
N/A	Single Patient Use Injectable Emulsion Lot # : EA7470 NDC # for Vial: 0409-4699-54 NDC # for Tray: 0409-4699-24 **Size/Packaging:** Glass Fliptop Vial 100 ml vials Glass Fliptop Vial	Nationwide including California	N/A

FOR ADDITIONAL INFORMATION, PLEASE VISIT THE FDA WEBSITE

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