



2020 Drug recalls

December 28 - [Sunstar Americas Inc. Expands Voluntary Nationwide Recall of Paroex® Chlorhexidine Gluconate Oral Rinse USP, 0.12% Due to Microbial Contamination | FDA](#)

October 28 - [Sunstar Americas Inc. Issues Voluntary Nationwide Recall of Paroex® Chlorhexidine Gluconate Oral Rinse USP, 0.12% Due to Microbial Contamination | FDA](#)

October 5 - [Marksans Pharma Limited Issues Expansion of Voluntary Nationwide Recall of Metformin Hydrochloride Extended-Release Tablets, USP 500mg & 750mg, Due to the Detection of N-Nitrosodimethylamine \(NDMA\) | FDA](#)

September 21 - [FDA Alerts of Perrigo's voluntary albuterol inhaler recall | FDA](#)

June 5 - [Apotex Corp. Issues Voluntary Nationwide Recall of Metformin Hydrochloride Extended-Release Tablets 500mg Due to the Detection of N-nitrosodimethylamine \(NDMA\) | FDA](#)

June 5 - [Teva Pharmaceuticals USA, Inc. Initiates Voluntary Nationwide Recall of Metformin Hydrochloride Extended-Release Tablets USP 500 mg and 750 mg Due to Detection of N-Nitrosodimethylamine \(NDMA\) | FDA](#)

June 1 - [Amneal Pharmaceuticals LLC Issues Voluntary Nationwide Recall of Metformin Hydrochloride Extended Release Tablets, USP, 500 mg and 750 mg, Due to Detection of N-Nitrosodimethylamine \(NDMA\) Impurity | FDA](#)

February 27 - [American Health Packaging Issues Voluntary Nationwide Recall of Ranitidine Tablets, USP 150 mg, 100 Count Unit Dose Blisters Due to the Detection of N-nitrosodimethylamine \(NDMA\) Impurity | FDA](#)

January 10 - [Taro Pharmaceuticals U.S.A., Inc. Issues Voluntary Nationwide Recall of Lamotrigine Tablets USP, 100 mg, 100 Count Bottles | FDA](#)

January 8 - [Mylan Initiates Voluntary Nationwide Recall of Three Lots of Nizatidine Capsules, USP, Due to the Detection of Trace Amounts of NDMA \(N-Nitrosodimethylamine\) Impurity Found in the Active Pharmaceutical Ingredient Manufactured by Solara Active Pharma Sciences](#)

January 7 - [Appco Pharma LLC Issues Voluntary Nationwide Recall of Ranitidine Hydrochloride Capsules 150 mg and 300 mg Due to an Elevated Amount of Unexpected Impurity, N-Nitrosodimethylamine \(NDMA\) | FDA](#)



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U.S. Department of Health and Human Services

200 Independence Avenue, SW
Room 509F, HHH Building
Washington, D.C. 20201
1-800-368-1019
(TDD/TTY: **1-800-537-7697**)

Complaint forms are available at:
hhs.gov/ocr/office/file/index.html.

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