Drug Recall List
Last Updated: November 2023

| Drug | Recall Details | Contact | Date | Drug Recall Class* |
| :---: | :---: | :---: | :---: | :---: |
| Oxandrin (oxandrolone) | 54569587500 49884030202 00185027260 68084042511 00591354560 00245027206 68084042421 00185027101 00245027111 00591354401 49884030101 68084042411 | If you have questions about this recall, call Upsher Smith Laboratories,Inc., 1-855-899-9180 | $\begin{aligned} & \text { November } \\ & 2023 \end{aligned}$ | Class III |
| $\begin{aligned} & \text { Brexafemme (ibrexafungerp) } \\ & 75788-115-04 \end{aligned}$ | Cross contamination | If you have questions about this recall, call Scynexis, 1-888-982-7299 | $\begin{aligned} & \text { October } \\ & 2023 \end{aligned}$ | Class $1$ |
| $\begin{aligned} & \hline \text { Sucralfate } \\ & 66689-305-16 \end{aligned}$ | Bacillus cereus contamination | If you have questions about this recall, call Vista Pharm, 1-877-530-1633 | $\begin{aligned} & \text { September } \\ & 2023 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |

## * Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

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| Sandimmune (cyclosporine) <br> 00078-0110-22 | Crystal formation | If you have questions <br> about this recall, call <br> Novartis Pharmaceuticals, <br> $1-800-882-6577$ | September <br> 2023 | Class <br> I |
| :--- | :--- | :--- | :--- | :--- |
| Digoxin <br> $10135-0747-01$ <br> $10135-0748-01$ | Label mix-up | If you have questions <br> about this recall, call <br> Marlex, 1-888-582-1953 | September <br> 2023 | Class <br> I |
| CROMOLYN SOD <br> $76204-025-96$ | Manufacturing issues related to an <br> alternative source of drug | If you have questions <br> about this recall, call <br> Ritedose <br> Pharmaceuticals.,1-803- <br> $806-3300$ | August <br> 2023 | Class <br> II |
| Ydemy (drospirenone/ethinyl <br> estradiol and levomefolate) <br> 68180-904-71 <br> 68180-904-73 | Lack of sterility assurance | If you have questions <br> about this recall, call Lupin <br> Pharmaceuticals Inc.1- <br> $410-576-2000$ ext 3 | August <br> 2023 | Class <br> II |
| Sodium Chloride Injection <br> 0264-5802-00 <br> 0264-5802-10 | Lack of sterility assurance | If you have questions <br> about this recall, call B. <br> Braun Medical,1-844-247- <br> 5287 | July <br> 2023 | Class |
| ALBUTEROL <br> 69097-142-60 |  | If you have questions <br> about this recall, call Cipla <br> $844-247-5287$ | July <br> 2023 | Class <br> I |

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| Contour Next Gen Glucose Meters 00193-7917-01 00193-7383-01 | Incorrect factory-set units of measurement | If you have questions about this recall, call Ascensia Diabetes Care US, 1-862-225-2902 | $\begin{aligned} & \text { May } \\ & 2023 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \text { CEFEPIME HCL } \\ & 70594-0089-02 \\ & 66794-0209-41 \end{aligned}$ | Quality concerns at its manufacturing facility | If you have questions about this recall, call Sedgwick, 1-888-7195826 | $\begin{aligned} & \hline \text { May } \\ & 2023 \end{aligned}$ | $\begin{aligned} & \hline \text { Class } \\ & \text { II } \end{aligned}$ |
| $\begin{aligned} & \text { Fentanyl buccal } \\ & 51862-634-28 \\ & 51862-635-28 \\ & 51862-636-28 \\ & 51862-637-28 \\ & 51862-638-28 \end{aligned}$ | Safety updates were omitted | If you have questions about this recall, call Mayne Pharma, 1-7523800 | $\begin{aligned} & \text { May } \\ & 2023 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { IIII } \end{aligned}$ |
| Prednisolone $50383-0042-24$ $50383-0042-48$ $50383-0042-24$ $50383-0042-48$ | CGMP Deviations: Firm went out of business and could no longer continue stability studies. | If you have questions about this recall, call Akorn, 1-800-932-5676 | $\begin{aligned} & \hline \text { May } \\ & 2023 \end{aligned}$ | $\begin{aligned} & \hline \text { Class } \\ & \text { II } \end{aligned}$ |
| $\begin{aligned} & \text { VALPROIC ACD } \\ & 50383-0792-16 \end{aligned}$ | CGMP Deviations: Firm went out of business and could no longer continue stability studies. | If you have questions about this recall, call Akorn, 1-800-932-5676 | $\begin{aligned} & \text { May } \\ & 2023 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| Fluticasone Propionate 50383-0700-09 <br> 50383-0700-16 | CGMP Deviations: Firm went out of business and could no longer continue stability studies. | If you have questions about this recall, call Akorn, 1-800-932-5676 | $\begin{aligned} & \text { May } \\ & 2023 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |

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| SULFAMETHOXAZOLE/TRIMET <br> HOPRIM <br> $50383-0823-16$ | CGMP Deviations: Firm went out of <br> business and could no longer <br> continue stability studies. | If you have questions <br> about this recall, call <br> Akorn, 1-800-932-5676 | May <br> 2023 | Class <br> II |
| :--- | :--- | :--- | :--- | :--- |
| PROMETHAZINE <br> $50383-0801-16$ | CGMP Deviations: Firm went out of <br> business and could no longer <br> continue stability studies. | If you have questions <br> about this recall, call <br> Akorn, 1-800-932-5676 | May <br> 2023 | Class <br> II |
| LIDOCAINE SOL 2\% VISC <br> $50383-0775-04$ | CGMP Deviations: Firm went out of <br> business and could no longer <br> continue stability studies. | If you have questions <br> about this recall, call <br> Akorn, 1-800-932-5676 | May <br> 2023 | Class |
| LEVETIRACETAM <br> $50383-0241-16$ | CGMP Deviations: Firm went out of <br> business and could no longer <br> continue stability studies. | If you have questions <br> about this recall, call <br> Akorn, 1-800-932-5676 | May <br> 2023 | Class |
| LIDOCAINE <br> 17478-0711-31 | CGMP Deviations: Firm went out of <br> business and could no longer <br> continue stability studies. | If you have questions <br> about this recall, call <br> Akorn, 1-800-932-5676 | May <br> 2023 | Class <br> II |
| KETOROLAC SOL 0.5\% <br> 17478-0209-19 <br> 17478-0209-10 | CGMP Deviations: Firm went out of <br> business and could no longer <br> continue stability studies. | If you have questions <br> about this recall, call <br> Akorn, 1-800-932-5676 | May <br> 2023 | Class <br> II |
| HYDROCORTISONE/ACETIC <br> ACID <br> $50383-0901-10 ~$ | CGMP Deviations: Firm went out of <br> business and could no longer <br> continue stability studies. | If you have questions <br> about this recall, call <br> Akorn, 1-800-932-5676 | May <br> 2023 | Class <br> II |

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| $\begin{array}{\|l} \hline \text { GUAIATUSSIN AC } \\ 50383-0087-16 \\ 50383-0087-04 \\ 50383-0087-16 \end{array}$ | CGMP Deviations: Firm went out of business and could no longer continue stability studies. | If you have questions about this recall, call Akorn, 1-800-932-5676 | $\begin{aligned} & \text { May } \\ & 2023 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \text { CLOBETASOL CRE } \\ & 50383-0267-30 \\ & 50383-0267-60 \\ & 50383-0979-04 \end{aligned}$ | CGMP Deviations: Firm went out of business and could no longer continue stability studies. | If you have questions about this recall, call Akorn, 1-800-932-5676 | $\begin{aligned} & \text { May } \\ & 2023 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| $\begin{aligned} & \text { CICLOPIROX } \\ & 50383041906 \end{aligned}$ | CGMP Deviations: Firm went out of business and could no longer continue stability studies. | If you have questions about this recall, call Akorn, 1-800-932-5676 | $\begin{aligned} & \text { May } \\ & 2023 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| ATROPINE $17478-0215-02$ $17478-0215-05$ $76478-0015-05$ $17478-0215-15$ $76478-0015-02$ | CGMP Deviations: Firm went out of business and could no longer continue stability studies. | If you have questions about this recall, call Akorn, 1-800-932-5676 | $\begin{aligned} & \text { May } \\ & 2023 \end{aligned}$ | $\begin{aligned} & \hline \text { Class } \\ & \text { II } \end{aligned}$ |
| $\begin{aligned} & \text { ACETIC ACID } \\ & 50383-0889-15 \end{aligned}$ | CGMP Deviations: Firm went out of business and could no longer continue stability studies. | If you have questions about this recall, call Akorn, 1-800-932-5676 | $\begin{aligned} & \text { May } \\ & 2023 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| Acetaminophen \& Codeine Phosphate 50383-0079-16 | CGMP Deviations: Firm went out of business and could no longer continue stability studies. | If you have questions about this recall, call Akorn, 1-800-932-5676 | $\begin{aligned} & \text { May } \\ & 2023 \end{aligned}$ | Class II |

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| RIFAMPIN <br> $61748-0018-30$ | CGMP Deviations: Firm went out of <br> business and could no longer <br> continue stability studies. | If you have questions <br> about this recall, call <br> Akorn, 1-800-932-5676 | May <br> 2023 | Class <br> II |
| :--- | :--- | :--- | :--- | :--- |
| OLOPATADINE <br> $50383-0943-23$ | CGMP Deviations: Firm went out of <br> business and could no longer <br> continue stability studies. | If you have questions <br> about this recall, call <br> Akorn, 1-800-932-5676 | May <br> 2023 | Class <br> II |
| AZELASTINE <br> $50383-0942-30$ | CGMP Deviations: Firm went out of <br> business and could no longer <br> continue stability studies. | If you have questions <br> about this recall, call <br> Akorn, 1-800-932-5676 | May <br> 2023 | llass |
| PROMETH/COD <br> $50383-0804-16$ | CGMP Deviations: Firm went out of <br> business and could no longer <br> continue stability studies. | If you have questions <br> about this recall, call <br> Akorn, 1-800-932-5676 | May <br> 2023 | Class |
| LIDO/PRILOCN CRE <br> $50383-0667-30 ~$ | CGMP Deviations: Firm went out of <br> business and could no longer <br> continue stability studies. | If you have questions <br> about this recall, call <br> Akorn, 1-800-932-5676 | May <br> 2023 | Class <br> II |
| MEGESTROL AC <br> $50383-0859-24 ~$ | CGMP Deviations: Firm went out of <br> business and could no longer <br> continue stability studies. | If you have questions <br> about this recall, call <br> Akorn, 1-800-932-5676 | May <br> 2023 | Class <br> II |
| AMANTADINE <br> $50383-0807-16 ~$ | CGMP Deviations: Firm went out of <br> business and could no longer <br> continue stability studies. | If you have questions <br> about this recall, call <br> Akorn, 1-800-932-5676 | May <br> 2023 | Class <br> II |

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| $\begin{aligned} & \text { ALBUTEROL } \\ & 50383-0740-16 \end{aligned}$ | CGMP Deviations: Firm went out of business and could no longer continue stability studies. | If you have questions about this recall, call Akorn, 1-800-932-5676 | $\begin{aligned} & \text { May } \\ & 2023 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \text { OXCARBAZEPIN } \\ & 50383-0312-84 \end{aligned}$ | CGMP Deviations: Firm went out of business and could no longer continue stability studies. | If you have questions about this recall, call Akorn, 1-800-932-5676 | $\begin{aligned} & \text { May } \\ & 2023 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { LACTULOSE } \\ & 50383-0357-31 \\ & 50383-0779-16 \\ & 50383-0779-32 \\ & 50383-0795-16 \\ & 50383-0779-33 \\ & \hline \end{aligned}$ | CGMP Deviations: Firm went out of business and could no longer continue stability studies. | If you have questions about this recall, call Akorn, 1-800-932-5676 | $\begin{aligned} & \text { May } \\ & 2023 \end{aligned}$ | Class II |
| APRACLONIDIN 17478-0716-10 | CGMP Deviations: Firm went out of business and could no longer continue stability studies. | If you have questions about this recall, call Akorn, 1-800-932-5676 | $\begin{aligned} & \text { May } \\ & 2023 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { CROMOLYN SOD } \\ & 17478-0291-11 \end{aligned}$ | CGMP Deviations: Firm went out of business and could no longer continue stability studies. | If you have questions about this recall, call Akorn, 1-800-932-5676 | $\begin{aligned} & \text { May } \\ & 2023 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { TIMOLOL MAL } \\ & 17478-0288-10 \\ & 17478-0288-11 \\ & 17478-0288-12 \\ & \hline \end{aligned}$ | CGMP Deviations: Firm went out of business and could no longer continue stability studies. | If you have questions about this recall, call Akorn, 1-800-932-5676 | $\begin{aligned} & \hline \text { May } \\ & 2023 \end{aligned}$ | Class <br> II |

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| $\begin{aligned} & \hline \text { PILOCARPINE } \\ & 17478-0223-12 \\ & 17478-0223-12 \\ & 17478-0224-12 \\ & 17478-0226-12 \end{aligned}$ | CGMP Deviations: Firm went out of business and could no longer continue stability studies. | If you have questions about this recall, call Akorn, 1-800-932-5676 | $\begin{aligned} & \hline \text { May } \\ & 2023 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Fyremadel (ganirelix acetate) 55566-1010-1 | glass particulate | If you have questions about this recall, call Ferring Pharmaceuticals Inc, 1-888-337-7464 | April 2023 | Class II |
| FreeStyle Libre Readers 57599080300 <br> 57599080500 <br> 57599000200 <br> 57599000098 <br> 57599071687 <br> 57599000021 <br> 57599000024 <br> 57599000041 | Defective batteries | If you have questions about this recall, call Abott, 1-855-632-8658 | April 2023 | Class I |
| Makena (hydroxyprogesterone caproate) <br> 55150030901 <br> 66993003901 <br> 67457096701 <br> 69238179701 <br> 55150031001 <br> 71225010401 <br> 71225010501 | The benefits of Makena do not outweigh the risks | If you have questions about this recall, call AMAG Pharmaceuticals, 1-877-411-2510 | April 2023 | Class II |

## * Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
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| Dabigatran Etexilate <br> 67877-475-60 | CGMP Deviations | If you have questions <br> about this recall, call <br> Ascend Laboratories, LLC, <br> $1-877-272-7901$ | March <br> 2023 | Class II |
| :--- | :--- | :--- | :--- | :--- |
| Nurtec ODT <br> $72618-300-2$ | The tablets are in a non-child <br> resistant blister card | If you have questions <br> about this recall, call <br> Pfizer Inc., 1-800-438-1985 | March <br> 2023 | Class II |
| Atovaquone <br> $31722-629-21$ | Microbial Contamination | If you have questions <br> about this recall, call <br> Camber, 1-732-529-0430 | March <br> 2023 | Class I |
| Brimonidine Tartrate <br> $60505-0564-1$ <br> $60505-0564-2$ <br> $60505-0564-3$ | Lack of sterility assurance | If you have questions <br> about this recall, call <br> Apotex Corp, 1-800-706- <br> 5575 | March <br> 2023 | Class II |
| Heparin Sodium Injection <br> $25021-404-01$ | Incorrect labeling | If you have questions <br> about this recall, call <br> Sagent Pharmaceuticals <br> Inc, 1-866-625-1618 | March <br> 2023 | Class II |
| Dabigatran Etexilate <br> $67877-0474-60$ | In you have questions <br> about this recall, call <br> Ascend Laboratories, LLC, <br> $1-877-272-7901$ | March <br> 2023 | Class II |  |
| Epinephrine Professional EMS, <br> Epinephrine Convenience Kit <br> $24357-012-12$ <br> $24357-011-13$ | Incorrect NDC number on the outer <br> carton of lot | If you have questions <br> about this recall, call <br> Focus Health Group, 1- <br> $800-249-1972$ | March <br> 2023 | Class III |

* Drug Recall Class

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| Blue Cross Blue Care Network of Michigan |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Insulin Syringes with the BD Micro-Fine IV Needle 08290329461 <br> 08290329420 $08290329424$ | Manufacturing issues | If you have questions about this recall, call BD. 1-844-823-5433 | $\begin{aligned} & \hline \text { February } \\ & 2023 \end{aligned}$ | Class II |
| Sodium chloride injection 00264580200 <br> 00264580210 | Lack of sterility assurance | If you have questions about this recall, call B. Braun Medical 1-800-2272862 | $\begin{aligned} & \text { February } \\ & 2023 \end{aligned}$ | Class II |
| TIROSINT®-SOL (levothyroxine sodium) 71858010505 71858011005 71858011205 71858011305 71858011505 71858011705 71858012005 71858012505 71858013005 71858013505 71858014005 71858014505 71858015005 71858015505 71858016005 | Subpotent | If you have questions about this recall, call IBSA Pharma Inc. 1-800-5873513 | $\begin{aligned} & \text { February } \\ & 2023 \end{aligned}$ | Class II |

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| Epinephrine (L-Adrenaline) | Discoloration | If you have questions about this recall, call Spectrum Laboratory Products 1-800-772-8786 | $\begin{aligned} & \text { January } \\ & 2023 \end{aligned}$ | Class I |
| :---: | :---: | :---: | :---: | :---: |
| Sodium Bicarbonate Injection 51754500105 51754500101 72572074020 72572074001 | vial breakage | If you have questions about this recall, call Exela Pharma Sciences 1--828-341-6118 Ext1017 | $\begin{aligned} & \text { December } \\ & 2022 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { quinapril/HCTZ } \\ & 65862016290 \end{aligned}$ | Impurities | If you have questions about this recall, call Aurobindo Pharma, 1-866-850-2876 | October 2022 | Class II |
| sodium bicarbonate 8.4\% <br> 51754500105 <br> 51754500101 <br> 72572074020 <br> 72572074001 | Vial breakage | If you have questions about this recall, call Exela Pharma Sciences 1--828-341-6118 Ext1017 | $\begin{aligned} & \text { October } \\ & 2022 \end{aligned}$ | Class I |
| $\begin{aligned} & \hline \text { Clopidogrel } \\ & 05140703210 \end{aligned}$ | Mislabeling | If you have questions about this recall, call Golden State Medical Supply 1-800-284-8633, Ext 116 | $\begin{aligned} & \text { October } \\ & 2022 \end{aligned}$ | Class I |
| $\begin{array}{\|l\|} \hline \text { Atenolol } \\ 06042902710 \end{array}$ | Mislabeling | If you have questions about this recall, call Golden State Medical Supply 1-800-284-8633, Ext 116 | October 2022 | Class I |

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Blue Cross
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| Acyclovir Sodium Injection <br> 05186260450 | Product complaint/particulate inside <br> the vial | If you have questions <br> about this recall, call <br> Qualanex 1-866-850-2876 <br> Option 2 | September <br> 2022 | Class I |
| :--- | :--- | :--- | :--- | :--- |
| Insulin Glargine <br> 04950239475 | Labeling: Missing label: Label <br> missing from some prefilled pens | If you have questions <br> about this recall, call <br> Mylan 1-888-406-9305 | August <br> 2022 | Class I |
| Morphine Sulfate Extended- <br> Release 60mg <br> 63629108901 | Labeling: Label Mix-up | If you have questions <br> about this recall, <br> Bryant Ranch Prepack, <br> Inc, 1-877-885-0882 | July <br> 2022 | Class I |
| Morphine Sulfate Extended- <br> Release 30mg <br> 63629108801 | Labeling: Label Mix-up | If you have questions <br> about this recall, <br> Bryant Ranch Prepack, <br> Inc, 1-877-885-0882 | July <br> 2022 | Class I |
| Losartan Potassium <br> 06438093305 <br> 06438093308 <br> 06438093405 <br> 06438093408 <br> 06438093505 <br> 06438093508 |  | If you have questions <br> about this recall, <br> Strides Inc.1-609-773- <br> 5000 | June <br> 2022 | Class II |

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Blue Cross
Blue Shield
Blue Care Network
of Michigan

| Losartan HCTZ 70518257800 68180021709 70518256400 68180021609 33342005010 70518323100 | CGMP Deviations | If you have questions about this recall, RemedyRepack Inc., 1-866-845-3791 | $\begin{aligned} & \hline \text { June } \\ & 2022 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Losartan Potassium \& HCT 68788775809 68788775803 | CGMP Deviations | If you have questions about this recall, Preferred Pharmaceuticals, Inc., 1-714-777-3729 | $\begin{aligned} & \hline \text { June } \\ & 2022 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { Vitamin D3 } \\ & 73198007530 \end{aligned}$ | Lack of assurance of sterility | If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-6732222. | $\begin{aligned} & \hline \text { June } \\ & 2022 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Testosterone Cypionate } \\ & 73198005410 \\ & 73198005505 \end{aligned}$ | Lack of assurance of sterility | If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-6732222. | $\begin{aligned} & \text { June } \\ & 2022 \end{aligned}$ | Class II |
| Daytrana (methylphenidate transdermal system) 06896855523 | Defective Delivery Syst | If you have questions about this recall, Noven Pharmaceuticals Inc, 1-305-253-5099 | $\begin{aligned} & \text { June } \\ & 2022 \end{aligned}$ | Class II |

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| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \text { EPINEPHrine } \\ & \text { 69374-544-10 } \\ & 69374-925-10 \end{aligned}$ | Lack of Assurance of Sterility | If you have questions about this recall, Nephron Sterile Compounding Center LLC, 1-800-4434313 | $\begin{aligned} & \text { June } \\ & 2022 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { Zonisamide } \\ & 06191977590 \end{aligned}$ | CGMP Deviations | If you have questions about this recall, Direct Rx, 1-678-619-5510 | $\begin{aligned} & \text { June } \\ & 2022 \end{aligned}$ | Class II |
| Losartan Pot/HCTZ 07218929090 07218929790 07218928990 07218916730 07218916790 | CGMP Deviations | If you have questions about this recall, Direct Rx, 1-678-619-5510 | $\begin{aligned} & \text { June } \\ & 2022 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Trulicity (dulaglutide) } \\ & 00002143480 \\ & 00002143380 \end{aligned}$ | TEMPERATURE ABUSE | If you have questions about this recall, Eli Lilly \& Company, 1-800-545-5979 | $\begin{aligned} & \text { June } \\ & 2022 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Humalog KwikPen } \\ & 00002879959 \\ & 00002751659 \end{aligned}$ | TEMPERATURE ABUSE | If you have questions about this recall, Eli Lilly \& Company, 1-800-545-5979 | $\begin{aligned} & \text { June } \\ & 2022 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

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| $\begin{array}{\|l} \hline \text { Xanax XR (alprazolam) extended } \\ \text { release } \\ 00009006807 \end{array}$ | Failed Dissolution Specifications | If you have questions about this recall, Viatris Inc,1-800.796.9526 | $\begin{aligned} & \text { May } \\ & 2022 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \hline \text { Accupril (Quinapril HCI Tablets) } \\ & 00071053023 \\ & 00071053223 \\ & 00071053523 \\ & \hline \end{aligned}$ | CGMP Deviations | If you have questions about this recall, Pfizer Inc.,1-800-438-1985 | $\begin{aligned} & \text { May } \\ & 2022 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { GaviLyte -C (Polyethylene Glycol } \\ & 3350,240 \mathrm{~g}) \\ & 04338606019 \end{aligned}$ | Failed Stability Specification | If you have questions about this recall, Lupin Pharmaceuticals Inc.1-410-576-2000 ext 3 | $\begin{aligned} & \text { May } \\ & 2022 \end{aligned}$ | Class II |
| Trulicity (dulaglutide) 00002143480 <br> 00002143380 | TEMPERATURE ABUSE | If you have questions about this recall, Eli Lilly \& Company, 1-800-545-5979 | $\begin{aligned} & \text { May } \\ & 2022 \end{aligned}$ | Class II |
| Esomeprazole Magnesium Delayed-Release $04229200916$ $04229201016$ | Failed Impurities/Degradation Specifications | If you have questions about this recall, call Mylan 1-888-406-9305 | $\begin{aligned} & \text { May } \\ & 2022 \end{aligned}$ | Class II |
| Pantoprazole Sodium 01366809690 | CGMP deviation | If you have questions about this recall, Torrent Pharma Inc, 1-888-280-2040 | $\begin{aligned} & \text { May } \\ & 2022 \end{aligned}$ | Class II |
| Lidocaine 00591207072 00591207030 | cGMP Deviations | If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800 | $\begin{aligned} & \hline \text { May } \\ & 2022 \end{aligned}$ | Class II |

## * Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

|  |  |  |  |  |
| :--- | :--- | :--- | :--- | :--- |
| Vancomycin <br> 70004092444 | CGMP Deviations | If you have questions <br> about this recall, SCA <br> Pharmaceuticals1-877- <br> $550-5059$ | May <br> 2022 | Class II |
| Fentanyl <br> 70004023132 | CGMP Deviations | If you have questions <br> about this recall, SCA <br> Pharmaceuticals1-877- <br> $550-5059$ | May <br> 2022 | Class II |
| Norepinephrine <br> 7000407840 | CGMP Deviations | If you have questions <br> about this recall, SCA <br> Pharmaceuticals1-877- <br> $550-5059$ | May <br> 2022 | Class II |
| Losartan Potassium <br> 7051832821 <br> 7051832820 | CGMP Deviations | If you have questions <br> about this recall, <br> RemedyRepack Inc., 1- <br> $866-845-3791$ | May <br> 2022 | Class II |
| Erythromycin Topical <br> 06373905368 | If you have questions <br> about this recall, call <br> McKesson Drug <br> Company, 1-330-487- <br> 0740. | May <br> 2022 | Class II |  |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Betamethasone Dipropionate 06373999665 | CGMP Deviations | If you have questions about this recall, call McKesson Drug Company, 1-330-4870740. | $\begin{aligned} & \text { May } \\ & 2022 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Lidocaine Prilocaine 06373905466 | CGMP Deviations | If you have questions about this recall, call McKesson Drug Company, 1-330-4870740. | $\begin{aligned} & \hline \text { May } \\ & 2022 \end{aligned}$ | Class II |
| Lidocaine Hydrochloride 06373997764 | CGMP Deviations | If you have questions about this recall, call McKesson Drug Company, 1-330-4870740. | $\begin{aligned} & \text { May } \\ & 2022 \end{aligned}$ | Class II |
| Halobetasol Propionate 06373997764 | CGMP Deviations | If you have questions about this recall, call McKesson Drug Company, 1-330-4870740. | $\begin{aligned} & \text { May } \\ & 2022 \end{aligned}$ | Class II |
| Losartan Potassium \& Hydrochlorothiazide 03334205007 <br> 03334205010 <br> 03334205044 | CGMP Deviations | If you have questions about this recall, Macleods Pharma Usa Inc, 1-888-943-3210 | $\begin{aligned} & \hline \text { May } \\ & 2022 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

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| 03334205207 <br> 03334205210 <br> 03334205244 <br> 03334205107 <br> 03334205110 <br> 03334205144 |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Losartan Potassium 03334204410 03334204444 03334204507 03334204510 03334204544 03334204607 03334204610 03334204644 | CGMP Deviations | If you have questions about this recall, Macleods Pharma Usa Inc, 1-888-943-3210 | $\begin{aligned} & \text { May } \\ & 2022 \end{aligned}$ | Class II |
| Losartan Potassium <br> 06818037603 <br> 06818037609 <br> 06818037703 <br> 06818037709 <br> 06818037803 <br> 06818037809 <br> 06818037803 <br> 06818037809 <br> 06818021706 <br> 06818021709 <br> 06818021606 <br> 06818021609 | CGMP Deviations | If you have questions about this recall, Lupin Pharmaceuticals Inc.1-410-576-2000 ext 3 | $\begin{aligned} & \text { May } \\ & 2022 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Blue Cross Blue Shield Blue Care Network of Michigan |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Zonisamide 06846212801 06846212901 06846213001 06846213005 | cGMP deviations | If you have questions about this recall, Glenmark Pharmaceuticals Inc, 1-888-721-7115 | $\begin{aligned} & \text { May } \\ & 2022 \end{aligned}$ | Class II |
| Lansoprazole 04359856178 | Failed Dissolution Specifications | If you have questions about this recall, Dr. Reddy's Laboratories, Inc., 1-888-375-3784 | $\begin{aligned} & \text { May } \\ & 2022 \end{aligned}$ | Class II |
| Hydroxocobalamin 73198008030 | CGMP Deviations | If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-6732222. | $\begin{aligned} & \text { May } \\ & 2022 \end{aligned}$ | Class II |
| Ultratest, Testosterone Cypionate 73198005810 | CGMP Deviations | If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-6732222. | $\begin{aligned} & \text { May } \\ & 2022 \end{aligned}$ | Class II |
| Testosterone Cypionate 73198005405 73198005510 73198005505 | CGMP Deviations | If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-6732222. | $\begin{aligned} & \text { May } \\ & 2022 \end{aligned}$ | Class II |

[^7]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

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| Formula F2, Papaverine <br> 73198000210 | CGMP Deviations | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 2022 | Class II |
| :--- | :--- | :--- | :--- | :--- |
| T-50, Papaverine <br> 73198002210 | CGMP Deviations | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 2022 | Class II |
| PGE-3, Alprostadil <br> 73198003010 | CGMP Deviations | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 2022 | Class II |
| PGE-2, Alprostadil <br> 73198002910 | CGMP Deviations | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 2022 | Class II |
| PGE-1, Alprostadil |  | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 73198-0028-10. | CGMP Deviations |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

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| :--- | :--- | :--- | :--- | :--- |
| FA, Papaverine <br> 73198000610 | CGMP Deviations | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 2022 | Class II |
| BIMIX-3, Papaverine <br> 73198002710 | CGMP Deviations | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 2022 | Class II |
| RE-2, Papaverine <br> 73198001610 | CGMP Deviations | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 2022 | Class II |
| RE-1, Papaverine <br> 73198001510 <br> 73198001503 | CGMP Deviations | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 2022 | Class II |

* Drug Recall Class

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Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
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| QM-4, Papaverine <br> 73198002010 | CGMP Deviations | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 2022 | Class II |
| :--- | :--- | :--- | :--- | :--- |
| QM-3, Papaverine <br> 73198001910 | CGMP Deviations | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 2022 | Class II |
| ST-2, Papaverine <br> 73198001210 | CGMP Deviations | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 2022 | Class II |
| ST-1, Papaverine <br> 73198001110 | CGMP Deviations | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 2022 | Class II |
| SB-6, Papaverine <br> 73198002510 | CGMP Deviations | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 2022 | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

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| :--- | :--- | :--- | :--- | :--- |
| SB-5, Papaverine <br> 73198002410 | CGMP Deviations | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 2022 | Class II |
| SB-4, Papaverine <br> 73198002310 | CGMP Deviations | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 2022 | Class II |
| T-101, Papaverine <br> 73198001410 <br> 73198001405 | CGMP Deviations | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 2022 | Class II |
| T-106, Papaverine |  | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 73198001310 <br> 73198001305 | CGMP Deviations |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
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| NB-243, Papaverine <br> $731980009-0$ | CGMP Deviations | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 2022 | Class II |
| :--- | :--- | :--- | :--- | :--- |
| Formula F9, Papaverine <br> 73198000410 | CGMP Deviations | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 2022 | Class II |
| T-105, Papaverine <br> 73198000510 <br> 73198000505 <br> 73198000503 | CGMP Deviations | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 2022 | Class II |
| QM-2 Papaverine <br> 073198001810 | CGMP Deviations | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 2022 | Class II |
| Betamethasone Dipropionate <br> 06373999665 | CGMP Deviations | If you have questions <br> about this recall, call <br> McKesson Drug <br> Company, 1-330-487- <br> 0740. | May <br> 2022 | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

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| :--- | :--- | :--- | :--- | :--- |
| AT-6, Papaverine <br> 73198004010 | CGMP Deviations | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 2022 | Class II |
| AT-1, Papaverine <br> 073198003910 | CGMP Deviations | If you have questions <br> about this recall, call <br> Olympia Compounding <br> Pharmacy, 1-407-673- <br> 2222. | May <br> 2022 | Class II |
| alprazolam XR <br> 05976200681 | Viatris Inc | If you have questions <br> about this recall, call <br> Viatris Inc, 1-724-514- <br> 1800 | May <br> 2022 | Class II |
| Insulin Glargine (insulin glargine- <br> yfgn) <br> 04950239380 | Labeling: Missing label on the vial | If you have questions <br> about this recall, call <br> Mylan 1-888-406-9305 | April <br> 2022 | Class II |
| Travoprost <br> 00378965132 |  | If you have questions <br> about this recall, call <br> Mylan 1-888-406-9305 | April <br> 2022 | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
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| TETRACAINE 1\% Tetracaine HCl <br> 04249443710 | Lack of Assurance of Sterility | If you have questions about this recall, Vitae Enim Vitae Scientific, Inc, Inc, 1-303-236-3010 | $\begin{aligned} & \hline \text { April } \\ & 2022 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \text { Sucralfate Oral } \\ & 06933914817 \\ & 06933914819 \end{aligned}$ | Labeling: Label Mix-Up | If you have questions about this recall, Vitae Enim Vitae Scientific, Inc, Inc, 1-303-236-3010 | $\begin{aligned} & \hline \text { April } \\ & 2022 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Rifampin } \\ & 06745744560 \end{aligned}$ |  | If you have questions about this recall, call Mylan 1-888-406-9305 | $\begin{aligned} & \text { April } \\ & 2022 \end{aligned}$ | Class II |
| quinapril $\mathrm{HCl} /$ hydrochlorothiazide | CGMP Deviations | If you have questions about this recall, Pfizer Inc.,1-800-438-1985 | $\begin{aligned} & \hline \text { April } \\ & 2022 \end{aligned}$ | Class II |
| PHENOBARBITAL Sodium 04249441525 <br> 04249441503 <br> 04249441625 <br> 04249441603 | Lack of Assurance of Sterility | If you have questions about this recall, Vitae Enim Vitae Scientific, Inc, Inc, 1-303-236-3010 | $\begin{aligned} & \hline \text { April } \\ & 2022 \end{aligned}$ | Class II |
| PAPAVERINE <br> HYDROCHLORIDE <br> 072516-02425 <br> 007251602410 | Lack of Assurance of Sterility | If you have questions about this recall, Vitae Enim Vitae Scientific, Inc, Inc, 1-303-236-3010 | $\begin{aligned} & \text { April } \\ & 2022 \end{aligned}$ | Class II |
| Orphenadrine Citrate ExtendedRelease 00185002201 | CGMP Deviations | If you have questions about this recall, Sandoz, Inc, 1-609-627-8500 | $\begin{aligned} & \text { April } \\ & 2022 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class $\mathbf{2}$ Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Lansoprazole Delayed-Release 068788639009 | Out of specification | If you have questions about this recall, Preferred Pharmaceuticals, Inc., 1-714-777-3729 | $\begin{aligned} & \text { April } \\ & 2022 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \hline \text { Janumet } \\ & 00006057502 \\ & 00006057503 \end{aligned}$ | Presence of Foreign Substance | If you have questions about this recall, <br> MERCK SHARP \& DOHME CORP, 1-(908) 423-1000 | $\begin{aligned} & \hline \text { April } \\ & 2022 \end{aligned}$ | Class II |
| Glycopyrrolate 01310701401 | Presence of Foreign Substance | If you have questions about this recall, Aurolife Pharma, LLC, 1-732-839-9400 option 2 | $\begin{aligned} & \text { April } \\ & 2022 \end{aligned}$ | Class II |
| Econazole Nitrate 068788740603 | CGMP Deviations | If you have questions about this recall, Preferred Pharmaceuticals, Inc., 1-714-777-3729 | $\begin{aligned} & \text { April } \\ & 2022 \end{aligned}$ | Class II |
| Diclofenac Sodium 68788791801 | CGMP Deviations | If you have questions about this recall, Preferred Pharmaceuticals, Inc., 1-714-777-3729 | $\begin{aligned} & \hline \text { April } \\ & 2022 \end{aligned}$ | Class II |
| Clobetasol Propionate 068788776801 | CGMP Deviations | If you have questions about this recall, Preferred Pharmaceuticals, Inc., 1-714-777-3729 | $\begin{aligned} & \text { April } \\ & 2022 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Accuretic (quinapril <br> HCI/hydrochlorothiazide) <br> 00071311223 <br> 0007102223 | CGMP Deviations | If you have questions <br> about this recall, Pfizer <br> Inc., 1-800-438-1985 | April <br> 00071022023 <br> 00071022323 |  |
| :--- | :--- | :--- | :--- | :--- |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| $\begin{aligned} & 05256501480 \\ & 05256501426 \end{aligned}$ |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Nystatin and Triamcinolone Acetonide $\begin{aligned} & 052565004215 \\ & 005256504230 \\ & 05256504260 \end{aligned}$ | cGMP deviations | If you have questions about this recall, Teligent Pharma, 1-856-697-1441 | $\begin{aligned} & \text { March } \\ & 2022 \end{aligned}$ | Class II |
| Lidocaine 05256512215 05256512230 05256512207 05256500814 00536128128 05038334135 | cGMP deviations | If you have questions about this recall, Teligent Pharma, 1-856-697-1441 | $\begin{aligned} & \hline \text { March } \\ & 2022 \end{aligned}$ | Class II |
| Hydrocortisone Butyrate 05256508702 05256508704 05186215904 05256508702 05256508704 | cGMP deviations | If you have questions about this recall, Teligent Pharma, 1-856-697-1441 | $\begin{aligned} & \hline \text { March } \\ & 2022 \end{aligned}$ | Class II |
| Halobetasol Propionate 07051203350 05256507315 05256507351 06373999867 | cGMP deviations | If you have questions about this recall, Teligent Pharma, 1-856-697-1441 | $\begin{aligned} & \hline \text { March } \\ & 2022 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Gentamicin Sulfate <br> 05256508515 <br> 05256508530 <br> 05256509015 <br> 05256509030 <br> 07051203630 | cGMP deviations | If you have questions about this recall, Teligent Pharma, 1-856-697-1441 | $\begin{aligned} & \text { March } \\ & 2022 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \text { Fluocinonide } \\ & 05256505415 \\ & 05256505460 \\ & 05256505430 \\ & 05256507911 \\ & 05256502520 \\ & 05256502559 \\ & \hline \end{aligned}$ | cGMP deviations | If you have questions about this recall, Teligent Pharma, 1-856-697-1441 | $\begin{aligned} & \text { March } \\ & 2022 \end{aligned}$ | Class II |
| Econazole Nitrate 05256502215 05256502285 05256502230 | cGMP deviations | If you have questions about this recall, Teligent Pharma, 1-856-697-1441 | $\begin{aligned} & \text { March } \\ & 2022 \end{aligned}$ | Class II |
| Diflorasone Diacetate 07051203160 <br> 05256506360 | cGMP deviations | If you have questions about this recall, Teligent Pharma, 1-856-697-1441 | $\begin{aligned} & \text { March } \\ & 2022 \end{aligned}$ | Class II |
| Diclofenac Sodium 06191967505 | Defective Container | If you have questions about this recall, Direct $R x, 1-678-619-5510$ | $\begin{aligned} & \text { March } \\ & 2022 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Diclofenac Sodium } \\ & 05256500205 \\ & 07051202505 \end{aligned}$ | cGMP deviations | If you have questions about this recall, Teligent Pharma, 1-856-697-1441 | $\begin{aligned} & \text { March } \\ & 2022 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class $\mathbf{2}$ Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

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| Desoximetasone 05256504599 05256504560 07051203710 05256503099 05256503060 05256503015 | cGMP deviations | If you have questions about this recall, Teligent Pharma, 1-856-697-1441 | $\begin{aligned} & \text { March } \\ & 2022 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Desonide 05256503815 05256503860 | cGMP deviations | If you have questions about this recall, Teligent Pharma, 1-856-697-1441 | $\begin{aligned} & \text { March } \\ & 2022 \end{aligned}$ | Class II |
| Clobetasol Propionate 05256509415 05256509430 05256509445 05256505502 05256505504 05256503915 05256503930 05256503945 05256503960 05256508215 05256508230 05256508260 07051202860 | cGMP deviation | If you have questions about this recall, Teligent Pharma, 1-856-697-1441 | $\begin{aligned} & \text { March } \\ & 2022 \end{aligned}$ | Class II |
| Betamethasone Dipropionate 05256502329 <br> 05256502359 | cGMP deviation | If you have questions about this recall, Teligent Pharma, 1-856-697-1441 | $\begin{aligned} & \text { March } \\ & 2022 \end{aligned}$ | Class II |

## * Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Alprazolam <br> 06042950418 | CGMP Deviation | If you have questions <br> about this recall, Golden <br> State Medical Supply Inc., <br> $1-805-477-9866$ Ext 4. | March <br> 2022 | Class II |
| :--- | :--- | :--- | :--- | :--- |
| hydrALAZINE <br> 00904644061 | Failed Impurities/Degradation <br> Specifications | If you have questions <br> about this recall, The <br> Harvard Drug Group, 1- <br> $800-875-0123$ | March <br> 2022 | Class II |
| Alprazolam C-IV <br> 06191983660 <br> 72189005860 | CGMP Deviations | If you have questions <br> about this recall, Direct <br> Rx, 1-678-619-5510 | March <br> 2022 | Class II |
| PALIPERIDONE EXTENDED- <br> RELEASE <br> 00904693761 | Failed Dissolution Specifications | If you have questions <br> about this recall, The <br> Harvard Drug Group, 1- <br> $800-875-0123$. | March <br> 2022 | Class II |
| Oxycodone Hydrochloride <br> 00360687406 <br> 06068-40677 <br> 06068740640 | Impurity failure at 0-time of the <br> repackaged lot | If you have questions <br> about this recall, American <br> Health Packaging, 1-614- <br> $492-8177$. | March <br> 2022 | Class II |
| Methylphenidate Hydrochloride <br> 6498022101 | Failed Tablet Specifications | If you have questions <br> about this recall, <br> RISING <br> PHARMACEUTICALS, <br> $1-866-562-4597$ | March <br> 2022 | Class II |

* Drug Recall Class

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NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Diazepam <br> 00527176836 | Failed Impurities/Degradation <br> Specifications | If you have questions <br> about this recall, Lannett <br> Company, Inc.1-215-333- <br> 9000 | February <br> 2022 | Class II |
| :--- | :--- | :--- | :--- | :--- |
| Morphine Sulfate <br> 73177010504 | Labeling; label mix-up | If you have questions <br> about this recall, <br> STAQ Pharma, Inc., 1- <br> $833-397-0106$. | February <br> 2022 | Class I |
| Hydromorphone HCL <br> 73177010405 | Labeling; label mix-up | If you have questions <br> about this recall, <br> STAQ Pharma, Inc., 1- <br> $833-397-0106$. | February <br> 2022 | Class I |
| Doxylamine Succinate and <br> Pyridoxine Hydrochloride <br> 0591-2132-01 | Failed Dissolution Specifications | If you have questions <br> about this recall, <br> Teva Pharmaceuticals <br> USA, 1-800-545-8800 | February <br> 2022 | Class II |
| Tretinoin <br> 00555080802 | Failed Dissolution Specifications | If you have questions <br> about this recall, <br> Teva Pharmaceuticals <br> USA, 1-800-545-8800 | February <br> 2022 | Class II |
| Proctofoam <br> 00037682210 |  | If you have questions <br> about this recall, call <br> Mylan 1-888-406-9305 | February <br> 2022 | Class II |

* Drug Recall Class

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Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Metformin Hydrochloride 07257803601 | CGMP Deviations | If you have questions about this recall, <br> VIONA <br> PHARMACEUTICALS <br> INC, 1-888-304-5022 | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \text { Proctofoam HC } \\ & 00037682210 \end{aligned}$ | cGMP deficiencies | If you have questions about this recall, call Mylan 1-888-406-9305 | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Diazepam } \\ & 00527176836 \end{aligned}$ | Failed Impurities/Degradation Specifications | If you have questions about this recall, Lannett Company, Inc.1-215-3339000. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| Trypan Blue 05446134801 | CGMP Deviations | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { Vancomycin HCl } \\ & 05446134801 \end{aligned}$ | CGMP Deviations | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Tetracaine HCl } \\ & 05 \Delta 46119503 \end{aligned}$ | CGMP Deviations | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |

[^8]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Promethazine HCl Topical <br> 05446134101 | CGMP Deviations | If you have questions <br> about this recall, <br> Edge Pharma, LLC. 1- <br> $802-992-1178$. | February <br> 2022 | Class II |
| :--- | :--- | :--- | :--- | :--- |
| Vitamin K Oral <br> 05446113203 | CGMP Deviations | If you have questions <br> about this recall, <br> Edge Pharma, LLC. $1-$ <br> $802-992-1178$. | February <br> 2022 | Class II |
| Lidocaine HCI/Phenylephrine HCI <br> 05446104503 | CGMP Deviations | If you have questions <br> about this recall, <br> Edge Pharma, LLC. $1-$ <br> $802-992-1178$. | February <br> 2022 | Class II |
| Phenol, Topical <br> 05446121103 |  | If you have questions <br> about this recall, | February <br> Edge Pharma, LLC. $1-$ <br> $802-992-1178$. | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class $\mathbf{2}$ Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Lidocaine $\mathrm{HCl} /$ Oxymetazoline HCl 05446125601 | CGMP Deviations | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| LET Topical Gel 05446060701 | CGMP Deviations | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| LT Topical 05446164701 | CGMP Deviations | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| Dibutyl Squaric Acid 05446104703 05446115603 | CGMP Deviations | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| Dexamethasone sodium phosphate 05446062201 | CGMP Deviations | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Cantharidin PLUS } \\ & 05446097003 \end{aligned}$ | CGMP Deviations | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| Cantharidin Gel 05446057203 | CGMP Deviations | If you have questions about this recall, Edge Pharma, LLC. 1- | February $2022$ | Class II |

* Drug Recall Class

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NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

|  |  | 802-992-1178. |  |  |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \text { Vancomycin HCl } \\ & 05446073601 \end{aligned}$ | Lack of Assurance of Sterility | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | February 2022 | Class II |
| Betadine (povidone-iodine) 05446168001 | Lack of Assurance of Sterility | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \hline \text { February } \\ & 2022 \end{aligned}$ | Class II |
| Phenylephrine $\mathrm{HCl} /$ Tropicamide/Cyclopentolate HCl/ Ketorolac 05446085903 | Lack of Assurance of Sterility | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| Phenylephrine $\mathrm{HCl} /$ Tropicamide/Cyclopentolate HCI/Ketorolac 05446099301 | Lack of Assurance of Sterility | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| Phenylephrine $\mathrm{HCl} /$ Tropicamide 05446081501 | Lack of Assurance of Sterility | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |

* Drug Recall Class

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| Phenylephrine HCI/Lidocaine 05446111801 | Lack of Assurance of Sterility | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \hline \text { Phenylephrine } \mathrm{HCl} \\ & 05446127001 \\ & 05446154505 \\ & 05446154410 \\ & 05446165201 \end{aligned}$ | Lack of Assurance of Sterility | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { MVASI } \\ & 05446166113 \end{aligned}$ | Lack of Assurance of Sterility | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | February 2022 | Class II |
| $\begin{array}{\|l\|} \hline \text { Neostigmine } \\ 05446154905 \\ \hline \end{array}$ | Lack of Assurance of Sterility | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \hline \text { February } \\ & 2022 \end{aligned}$ | Class II |
| $\begin{array}{\|l\|} \hline \text { Moxifloxacin } \\ 05446105001 \\ \hline \end{array}$ | Lack of Assurance of Sterility | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { Mitomycin-C } \\ & 05446141601 \\ & 05446100901 \\ & 05446101101 \end{aligned}$ | Lack of Assurance of Sterility | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |

[^9]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| $\begin{aligned} & \hline \text { Methotrexate } \\ & 05446150505 \end{aligned}$ | Lack of Assurance of Sterility | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Methacholine 05446160005 05446124601 05446124701 05446124901 | Lack of Assurance of Sterility | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| Lidocaine HCL / Bupivacaine HCL <br> 05446154818 | Lack of Assurance of Sterility | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| Gemcitabine 05446156650 | Lack of Assurance of Sterility | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| Epinephrine/Lidocaine 05446086301 | Lack of Assurance of Sterility | If you have questions about this recall, <br> Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Edetate Disodium (EDTA) } \\ & 00544614271 \\ & 05446142810 \end{aligned}$ | Lack of Assurance of Sterility | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |

[^10]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Dexamethasone sodium phosphate $05446084801$ | Lack of Assurance of Sterility | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{array}{\|l\|} \hline \text { Cefuroxime } \\ 05446100301 \\ \hline \end{array}$ | Lack of Assurance of Sterility | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| Ceftazidime 05446073301 | Lack of Assurance of Sterility | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| Lidocaine $\mathrm{HCl} /$ Epinephrine 05446126801 | Lack of Assurance of Sterility | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | February $2022$ | Class II |
| $\begin{aligned} & \hline \text { Lidocaine HCl } \\ & 05446085010 \end{aligned}$ | Lack of Assurance of Sterility | If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { Polymyxin B } \\ & 05515023410 \end{aligned}$ | Presence of Particulate Matter | If you have questions about this recall, AuroMedics Pharma LLC, 1-732-839-9400 ext 2. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| $\begin{array}{\|l\|} \hline \text { Moxifloxacin } \\ 06586284003 \end{array}$ | Failed impurities/degradation specifications | If you have questions about this recall, Aurobindo Pharma, 1-866-850-2876 | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |

* Drug Recall Class

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| Blue Cross Blue Care Network of Michigan |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Pioglitazone 05723722105 | Superpotent and Failed Tablet/Capsule Specifications | If you have questions about this recall, Aurobindo Pharma, 1-866-850-2876 | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| Metoprolol Succinate ExtendedRelease 06787759001 | Failed Dissolution Specifications | If you have questions about this recall, Ascend Laboratories LLC, 1-201-476-1977. | $\begin{array}{\|l} \hline \text { February } \\ 2022 \end{array}$ | Class II |
| Pyrazinamide 06725366010 | cGMP Deviations | If you have questions about this recall, ANI Pharmaceuticals, Inc., 1-800-308-6755. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| Alprazolam 06725390110 06725390150 06725390111 06725390210 06725390250 06725390211 06725390310 06725390350 | cGMP Deviations | If you have questions about this recall, ANI Pharmaceuticals, Inc., 1-800-308-6755. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |
| Metoprolol Succinate ExtendedRelease 06800150103 | Failed Dissolution Specifications | If you have questions about this recall, American Health Packaging, 1-614-492-8177. | $\begin{aligned} & \text { February } \\ & 2022 \end{aligned}$ | Class II |

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|  |  |  |  |  |
| :--- | :--- | :--- | :--- | :--- |

## * Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
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| 8.4\% Sodium Bicarbonate 05175450011 | Lack of Assurance of Sterility | If you have questions about this recall, Exela Pharma Sciences LLC, 1-888-451-4321 | $\begin{aligned} & \text { January } \\ & 2022 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \hline \text { Metformin } \\ & 70518292000 \end{aligned}$ | CGMP Deviations | If you have questions about this recall, RemedyRepack Inc., 1-866-845-3791 | $\begin{aligned} & \text { January } \\ & 2022 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| Norepinephrine Bitartrate 00703115303 | Lack of Assurance of Sterility | If you have questions about this recall, Teva Pharmaceuticals USA, $1-800-545-8800$ | $\begin{aligned} & \text { January } \\ & 2022 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| MethyIPREDNISolone Acetate 01671409001 | Lack of Assurance of Sterility | If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800 | $\begin{aligned} & \text { January } \\ & 2022 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| Carbamazepine 01366826801 | Failed Dissolution Specifications | If you have questions about this recall, Torrent Pharma Inc, 1-888-280-2040. | $\begin{aligned} & \text { January } \\ & 2022 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| Clobetasol Propionate 05167212593 | Microbial Contamination of NonSterile Products | If you have questions about this recall, Taro Pharmaceuticals U.S.A., Inc.,1-800-544-1449 ext 6066. | $\begin{aligned} & \text { January } \\ & 2022 \end{aligned}$ | Class |

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NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Pregabalin <br> 04733568788 | Failed Tablet/Capsule Specifications | If you have questions <br> about this recall, <br> SUN PHARMACEUTICAL <br> INDUSTRIES INC, 1-800- <br> $818-4555$ | January <br> 2022 | Class <br> II |
| :--- | :--- | :--- | :--- | :--- |
| Lidocaine Hydrochloride <br> 06373999764 | Superpotent Drug | If you have questions <br> about this recall, Teligent <br> Pharma, 1-856-697-1441 | January <br> 2022 | Class <br> II |
| Lidocaine Hydrochloride <br> 06373999764 | Superpotent Drug | If you have questions <br> about this recall, Teligent <br> Pharma, 1-856-697-1441 | January <br> 2022 | Class <br> I |
| Betamethasone Dipropionate <br> 05256502329 | Failed Stability Specifications | If you have questions <br> about this recall, Teligent <br> Pharma, 1-856-697-1441 | January <br> 2022 | Class <br> II |
| Penicillin V Potassium <br> 00093412573 <br> 00093412574 | Subpotent | If you have questions <br> about this recall, <br> Teva Pharmaceuticals <br> USA, <br> $1-800-545-8800$ | December <br> 2021 | Class <br> II |
| Fexofenadine Hydrochloride <br> 00904697940 | Failed Impurities/Degradation <br> Specifications | If you have questions <br> about this recall, The <br> Harvard Drug Group, 1- <br> $800-875-0123 . ~$ | December <br> 2021 | Class <br> II |

[^11]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

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| Clindamycin and Benzoyl <br> Peroxide Gel <br> 00781726368 | Superpotent Drug | If you have questions about this recall, TOLMAR, Inc., 1-970-2124500 | $\begin{aligned} & \hline \text { December } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| :---: | :---: | :---: | :---: | :---: |
| Lidocaine Hydrochloride 05256500950 | Superpotent Drug | If you have questions about this recall, Teligent Pharma, 1-856-697-1441 | $\begin{aligned} & \text { December } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \hline \end{aligned}$ |
| Lidocaine Hydrochloride 06373999764 <br> 05256500950 | CGMP Deviations | If you have questions about this recall, Teligent Pharma, 1-856-697-1441 | $\begin{aligned} & \text { December } \\ & 2021 \end{aligned}$ | $\begin{array}{\|l} \hline \text { Class } \\ \text { II } \\ \hline \end{array}$ |
| $\begin{aligned} & \hline \text { Cefixime } \\ & 70518274902 \end{aligned}$ | Failed Impurities/Degradation Specifications | If you have questions about this recall, RemedyRepack Inc., 1-866-845-3791 | $\begin{aligned} & \text { December } \\ & 2021 \end{aligned}$ | $\begin{array}{\|l\|} \hline \text { Class } \\ \text { II } \end{array}$ |
| Diclofenac Sodium 68788770701 | Defective container | If you have questions about this recall, Preferred Pharmaceuticals, Inc., 1-714-777-3729 | $\begin{aligned} & \text { December } \\ & 2021 \end{aligned}$ | $\begin{array}{\|l} \hline \text { Class } \\ \text { II } \end{array}$ |
| Hydrocodone Bitartrate and Acetaminophen <br> 03172299701 | Product Mix-up | If you have questions about this recall, Ascent Pharmaceuticals, Inc., 1-631-851-0550 | $\begin{aligned} & \hline \text { December } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |

[^12]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

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| $\begin{aligned} & \hline \text { Cefixime } \\ & 06787758450 \end{aligned}$ | Failed impurities/degradation specifications | If you have questions about this recall, <br> Ascend Laboratories, LLC, 1-201-476-1977 | $\begin{aligned} & \text { December } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \hline \text { Class } \\ & \text { II } \end{aligned}$ |
| :---: | :---: | :---: | :---: | :---: |
| Ascorbic Acid | Lack of Assurance of Sterility | If you have questions about this recall, ASP Cares, 1-210-615-7400 | $\begin{aligned} & \text { December } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \hline \text { Class } \\ & \text { II } \end{aligned}$ |
| $\begin{aligned} & \text { Diclofenac Sodium } \\ & 05266500205 \\ & 05266502505 \end{aligned}$ | Defective Container | If you have questions about this recall, Teligent Pharma, 1-856-697-1441 | $\begin{aligned} & \text { December } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \hline \text { Class } \\ & \text { II } \end{aligned}$ |
| Flocinolone Acetonide 06516270486 <br> 06516270386 | Subpotent Drug | If you have questions about this recall, Amneal Pharmaceuticals., 1-833-582-0812. | $\begin{aligned} & \text { December } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \hline \text { Class } \\ & \text { II } \end{aligned}$ |
| $\begin{aligned} & \text { Cefixime } \\ & 06787758450 \end{aligned}$ | Failed Impurities/Degradation Specifications | If you have questions about this recall, Ascend Laboratories LLC, 1-201-476-1977 | November 2021 | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| Tadalafil 01671407501 01671407701 | Incorrect Product Formulation | If you have questions about this recall, <br> SUN PHARMACEUTICAL INDUSTRIES INC, 1-800-818-4555 | November 2021 | $\begin{aligned} & \hline \text { Class } \\ & \text { II } \end{aligned}$ |

[^13]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Ezetimibe and Simvastatin 05140719305 | Failed Excipient Specification | If you have questions about this recall, Dr. Reddy's Laboratories, Inc., 1-888-375-3784 | November 2021 | $\begin{aligned} & \hline \text { Class } \\ & \text { II } \end{aligned}$ |
| :---: | :---: | :---: | :---: | :---: |
| Ezetimibe and Simvastatin 05140719290 <br> 05140719205 | cGMP Deviations | If you have questions about this recall, Dr. Reddy's Laboratories, Inc., 1-888-375-3784 | November 2021 | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| Methocarbamol 07133517952 07133517954 07133517957 | Labeling: Label Error on Declared Strength | If you have questions about this recall, 1-877-885-0882. | November 2021 | Class $1$ |
| Irbesartan and Hydrochlorothiazide 06818041306 06818041309 06818041406 06818041409 | CGMP Deviations | If you have questions about this recall, Lupin Pharmaceuticals Inc.1-410-576-2000 ext 3 | November 2021 | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| Irbesartan 06818041006 06818041009 06818041106 06818041109 06818041206 06818041209 | CGMP Deviations | If you have questions about this recall, Lupin Pharmaceuticals Inc.1-410-576-2000 ext 3 | November 2021 | $\begin{aligned} & \hline \text { Class } \\ & \text { II } \end{aligned}$ |

## * Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Ezetimibe and Simvastatin <br> 04359874290 <br> 04359874210 <br> 0435987430 <br> 04359874490 <br> 04359874530 <br> 04359874590 <br> 04359874505 <br> 04359874330 <br> 04359874390 <br> 04359874305 | Failed Excipient Specifications | If you have questions <br> about this recall, Dr. <br> Reddy's Laboratories, <br> Inc., 1-888-375-3784 | November <br> 2021 | Class <br> II |
| :--- | :--- | :--- | :--- | :--- |
|  |  |  |  |  |
| Rocuronium Bromide <br> 6679402841 |  |  |  |  |

[^14]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| AirDuo Digihaler 232/14 <br> (fluticasone propionate 113 mcg <br> and salmeterol 14 mcg ) <br> 05931013606 | Subpotent drug | If you have questions <br> about this recall, <br> Teva Pharmaceuticals <br> USA, <br> $1-800-545-8800$ | October <br> 2021 | Class <br> II |
| :--- | :--- | :--- | :--- | :--- |
| AirDuo Digihaler 113/14 <br> (fluticasone propionate 113 mcg <br> and salmeterol 14 mcg ) <br> 05931012906 | Subpotent drug | If you have questions <br> about this recall, <br> Teva Pharmaceuticals <br> USA, <br> $1-800-545-8800$ | October <br> 2021 | Class <br> II |
| AirDuo Digihaler 55/14 <br> (fluticasone propionate 55 mcg <br> and salmeterol 14 mcg ) <br> 05931011106 | Subpotent drug | If you have questions <br> about this recall, <br> Teva Pharmaceuticals <br> USA, | October <br> 2021 | Class |
| II |  |  |  |  |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

|  |  |  |  |  |
| :--- | :--- | :--- | :--- | :--- |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class $\mathbf{2}$ Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| $\begin{aligned} & 07360700102 \\ & 07360700110 \end{aligned}$ |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \text { Meclizine HCl Tablets, } 25 \mathrm{mg} \\ & 01657175201 \end{aligned}$ | Labeling: Incorrect Instructions | If you have questions about this recall, RISING PHARMACEUTICALS, 1-866-562-4597 | October $2021$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| Firvanq (vancomycin hydrochloride for oral solution) 06562820605 | Product Mix-up | If you have questions about this recall, Azurity Pharmaceuticals, Inc., 1-800-461-7449 | $\begin{aligned} & \hline \text { October } \\ & 2021 \end{aligned}$ | Class $1$ |
| Morphine Sulfate 06332345201 | Defective container | If you have questions about this recall, Fresenius Kabi USA, 1-800-551-7176. | $\begin{aligned} & \hline \text { October } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| Valproic Acid 06068726256 06068726242 | CGMP Deviations | If you have questions about this recall, American Health Packaging, 1-800-707-4621. | $\begin{aligned} & \text { October } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| Lyrica CR (pregabalin) 00071102901 | Failed Dissolution Specifications | If you have questions about this recall, Pfizer Inc.,1-800-438-1985. | $\begin{array}{\|l} \hline \text { October } \\ 2021 \end{array}$ | $\begin{aligned} & \hline \text { Class } \\ & \text { II } \end{aligned}$ |

[^15]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Zonisamide <br> 61919077530 | CGMP deviations | If you have questions <br> about this recall, Direct <br> Rx, 1-678-619-5510 | September <br> 2021 | Class <br> II |
| :--- | :--- | :--- | :--- | :--- |
| Metoprolol Tartrate <br> 06586206499 | Presence of Foreign Substance | If you have questions <br> about this recall, <br> Aurobindo Pharma USA <br> Inc., 1-866-850-2876 | September <br> 2021 | Class <br> II |
| Entacapone <br> 00904682204 | Failed Dissolution Specifications | If you have questions <br> about this recall, The <br> Harvard Drug Group, 1- <br> $800-875-0123$. | September <br> 2021 | Class <br> II |
| Valproic Acid <br> 06043262116 |  | If you have questions <br> about this recall, <br> Morton Grove <br> Pharmaceuticals, Inc., 1- <br> $888-721-7115$ | September <br> 2021 | Class |
| II |  |  |  |  |

[^16]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| $\begin{array}{\|l\|} \hline \text { Arformoterol Tartrate } \\ 06846283365 \\ 06846283335 \end{array}$ | Lack of Assurance of Sterility | If you have questions about this recall, Glenmark Pharmaceuticals Inc, 1-888-721-7115 | $\begin{aligned} & \text { September } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| :---: | :---: | :---: | :---: | :---: |
| Zonisamide 06846212901 06846213001 06846213005 | CGMP Deviations | If you have questions about this recall, Glenmark Pharmaceuticals Inc, 1-888-721-7115 | $\begin{aligned} & \text { September } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| $\begin{aligned} & \hline \text { Chlorzoxazone } \\ & 06846272401 \\ & 06846272501 \end{aligned}$ | CGMP Deviations | If you have questions about this recall, Glenmark Pharmaceuticals Inc, 1-888-721-7115 | $\begin{aligned} & \text { September } \\ & 2021 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Naproxen Sodium } \\ & 06846217801 \\ & 06846217901 \\ & 06846217905 \end{aligned}$ | CGMP Deviations | If you have questions about this recall, Glenmark Pharmaceuticals Inc, 1-888-721-7115 | $\begin{aligned} & \text { September } \\ & 2021 \end{aligned}$ | Class II |
| Fulvestrant 06846231732 | Lack of Assurance of Sterility | If you have questions about this recall, Glenmark Pharmaceuticals Inc, 1-888-721-7115 | $\begin{aligned} & \text { September } \\ & 2021 \end{aligned}$ | Class II |

## * Drug Recall Class

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Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Carbamazepine <br> 00904617261 | Failed Dissolution Specifications | If you have questions <br> about this recall, The <br> Harvard Drug Group, 1- <br> $800-875-0123$. | September <br> 2021 | Class <br> II |
| :--- | :--- | :--- | :--- | :--- |
| Betadine Solution Swabstick <br> 06761815301 <br> 06761815303 | Subpotent Drug | If you have questions <br> about this recall, <br> AVRIO HEALTH L.P, 1- <br> $888-726-7535$ | September <br> 2021 | Class <br> II |
| Chantix (varenicline) <br> 00069047103 <br> 00069046856 <br> 00069046956 <br> 00069046903 | CGMP Deviations | If you have questions <br> about this recall, Pfizer <br> Inc., 1-800-438-1985. | September <br> 2021 | Class <br> II |
| Oxycodone Hydrochloride <br> 04285800201 | Presence of Foreign <br> Tablets/Capsules; A single foreign <br> tablet Hydrochlorothiazide/Lisinopril <br> $25 / 20$ was found in one bottle | If you have questions <br> about this recall, Akorn, <br> Inc, Rhodes <br> Pharmaceuticals, L.P., <br> $1-401-262-9400$, Prompt 2 | September <br> 2021 | Class <br> II |
| Betamethasone Dipropionate <br> 06174848030 | Failed impurities/degradation <br> specification: Out of Specification for <br> an unknown impurity observed in <br> topical product. | If you have questions <br> about this recall, Akorn, <br> Inc, 1-800-932-5676, | September <br> Prompt 2 | Class |
| II |  |  |  |  |

* Drug Recall Class

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NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Clopidogrel <br> 03334206015 | Presence of foreign matter | If you have questions <br> about this recall, <br> Macleods Pharma Usa <br> Inc, $1-888-943-3210$ | September <br> 2021 | II |
| :--- | :--- | :--- | :--- | :--- |
| Cyclobenzaprine Hydrochloride <br> 00591333001 <br> 07019901401 <br> 05723726601 | CGMP Deviations | If you have questions <br> about this recall, <br> Teva Pharmaceuticals <br> USA, 1-800-545-8800 | September <br> 2021 | II |
| FLUDARABINE PHOSPHATE <br> FOR INJECTION <br> 02420123701 | Lack of Assurance of Sterility: the <br> manufacturing firm had microbial <br> recoveries during environmental <br> monitoring in aseptic areas of <br> manufacturing. | If you have questions <br> about this recall, <br> Custopharm, Inc., 1760- <br> $683-0901$ | September <br> 2021 | II |
| Erythromycin Topical Solution <br> 05256502759 | Defective container: possibility for <br> lack of seal integrity. | If you have questions <br> about this recall, Teligent <br> Pharma, 1-856-697-1441 | September <br> 2021 | II |
| Atovaquone Oral Suspension <br> 01070222321 | Temperature abuse: the firm received <br> customer complaints of unusual <br> grittiness in the product. | If you have questions <br> about this recall, <br> KVK-Tech, Inc., 1-215- <br> $579-1842$ | September <br> 2020 | II |
| Carvedilol <br> 06838209505 <br> 70518182601 | Presence of Foreign <br> Tablets/Capsules; report of two <br> Paroxetine tablets were found in the <br> bottle | If you have questions <br> about this recall, <br> RemedyRepack Inc., 1- <br> $866-845-3791$ | September <br> 2021 | II |

* Drug Recall Class

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NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| $\begin{aligned} & \hline \text { Adenosine } \\ & 01671418001 \end{aligned}$ | Lack of Assurance of Sterility | If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800 | $\begin{aligned} & \text { August } \\ & 2021 \end{aligned}$ | II |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \text { Leucovorin Calcium } \\ & 00703514001 \\ & 00703514591 \end{aligned}$ | Lack of Assurance of Sterility | If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800 | $\begin{aligned} & \text { August } \\ & 2021 \end{aligned}$ | II |
| Octreotide Acetate 00703331101 <br> 00703330101 | Lack of Assurance of Sterility | If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800 | $\begin{aligned} & \text { August } \\ & 2021 \end{aligned}$ | II |
| Methylprednisolone Acetate 00703005101 <br> 00703006301 | Lack of Assurance of Sterility | If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800 | $\begin{aligned} & \text { August } \\ & 2021 \end{aligned}$ | II |
| $\begin{aligned} & \text { Alprostadil } \\ & 00703150101 \end{aligned}$ | Lack of Assurance of Sterility | If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800 | $\begin{array}{\|l\|} \hline \text { August } \\ 2021 \end{array}$ | II |
| Metoclopramide 00703450201 | Lack of Assurance of Sterility | If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800 | $\begin{aligned} & \text { August } \\ & 2021 \end{aligned}$ | II |
| Adenosine Injection 00703877601 | Lack of Assurance of Sterility | If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800 | $\begin{aligned} & \text { August } \\ & 2021 \end{aligned}$ | II |

## * Drug Recall Class

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NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Norepinephrine Bitartrate <br> 00703115301 | Lack of Assurance of Sterility | If you have questions <br> about this recall, <br> Teva Pharmaceuticals <br> USA, 1-800-545-8800 | August <br> 2021 | II |
| :--- | :--- | :--- | :--- | :--- |
| Epoprostenol Sodium <br> 00703199501 | Lack of Assurance of Sterility | If you have questions <br> about this recall, <br> Teva Pharmaceuticals <br> USA, 1-800-545-8800 | August <br> 2021 | II |
| Leucovorin Calcium <br> 00703514501 | Lack of Assurance of Sterility | If you have questions <br> about this recall, <br> Teva Pharmaceuticals <br> USA, 1-800-545-8800 | August <br> 2021 | II |
| Octreotide Acetate <br> 00703333301 | Lack of Assurance of Sterility | If you have questions <br> about this recall, <br> Teva Pharmaceuticals <br> USA, 1-800-545-8800 | August <br> 2021 | II |
| Vecuronium Bromide <br> 00703291401 | Lack of Assurance of Sterility | If you have questions <br> about this recall, <br> Teva Pharmaceuticals <br> USA, 1-800-545-8800 | August <br> 2021 | II |
| Idarubicin Hydrochloride <br> 00703415611 | Lack of Assurance of Sterility | If you have questions <br> about this recall, <br> Teva Pharmaceuticals <br> USA, 1-800-545-8800 | August <br> 2021 | II |
| Amikacin Sulfate <br> 00703904001 | If you have questions <br> about this recall, <br> Teva Pharmaceuticals <br> USA, 1-800-545-8800 | August <br> 2021 | II |  |

* Drug Recall Class

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Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

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| Haloperidol Decanoate <br> 00703713101 <br> 00703713301 | Lack of Assurance of Sterility | If you have questions <br> about this recall, <br> Teva Pharmaceuticals <br> USA, 1-800-545-8800 | August <br> 2021 | II |
| :--- | :--- | :--- | :--- | :--- |
| Methylprednisolone Acetate <br> 00703003101 <br> 00703004301 <br> 00703004501 | Lack of Assurance of Sterility | If you have questions <br> about this recall, <br> Teva Pharmaceuticals <br> USA, 1-800-545-8800 | August <br> 2021 | II |
| DAUNOrubicin Hydrochloride <br> 00703523313 | Lack of Assurance of Sterility | If you have questions <br> about this recall, <br> Teva Pharmaceuticals <br> USA, 1-800-545-8800 | August <br> 2021 | II |
| GaviLyteTM <br> 04338606019 | Failed Stability Specification; Out of <br> specification for Osmolarity | If you have questions <br> about this recall, Lupin <br> Pharmaceuticals Inc.1- <br> $410-576-2000$ ext 3 | August <br> 2021 | II |
| Combipatch <br> 06896805148 <br> 06896805258 | Failed Stability Specifications; out of <br> specification for shear. | If you have questions <br> about this recall, <br> Noven Pharmaceuticals <br> Inc, 1-305-253-5099 | August <br> 2021 | II |
| Tizanidine HCI <br> 05511118015 | Failed Tablet/Capsule Specification: <br> Some tablets are shaved | If you have questions <br> about this recall, <br> Dr. Reddy's Laboratories, <br> Inc., 1-888-375-3784 | August <br> 2021 | II |

[^17]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Triamcinolone Acetonide <br> 05256501480 | Correct Labeled Product Mispack | If you have questions <br> about this recall, Teligent <br> Pharma, 1-856-697-1441 | August <br> 2021 | II |
| :--- | :--- | :--- | :--- | :--- |
| Econazole Nitrate <br> 05256501480 | Correct Labeled Product Mispack | If you have questions <br> about this recall, Teligent <br> Pharma, 1-856-697-1441 | August <br> 2021 | II |
| Ethosuximide <br> 00121067016 | Voluntary: Firm initiated | If you have questions <br> about this recall, PAI <br> Holdings, LLC. dba <br> Pharmaceutical <br> Associates Inc1-864-277- <br> 7282 ext 0 | August <br> 2021 | II |
| Cimetidine Hydrochloride <br> 00121064908 | Voluntary: Firm initiated | If you have questions <br> about this recall, PAI <br> Holdings, LLC. dba | August <br> Pharmaceutical <br> Associates Inc1-864-277- <br> 7282 ext 0 | II |
| Nystatin Oral <br> 00121086816 <br> 00121086802 |  | If you have questions <br> about this recall, PAI <br> Holdings, LLC. dba <br> Pharmaceutical | August <br> Associates Inc1-864-277- <br> 7282 ext 0 | II |

* Drug Recall Class

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Blue Cross Blue Shield
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| Venlafaxine <br> 01671465701 | Voluntary: Firm initiated |  | August <br> 2021 | II |
| :--- | :--- | :--- | :--- | :--- |
| Sulfamethoxazole and <br> Trimethoprim <br> 06586242005 | Voluntary: Firm initiated | If you have questions <br> about this recall, <br> Aurobindo Pharma USA <br> Inc., 1-866-850-2876 | August <br> 2021 | II |
| Zyprexa Intramuscular, <br> Olanzapine <br> 00002759701 | Voluntary: Firm initiated | If you have questions <br>  <br> Company. 1-317-276- <br> 2000 | August <br> 2021 | II |
| Chantix <br> 00069046856 <br> 00069047103 <br> 00069046956 |  | If you have questions <br> about this recall, Pfizer <br> Inc., 1-800-438-1985. | August <br> 2021 | II |
| Estriol <br> 04614430001 | CGMP Deviations | If you have questions <br> about this recall, <br> API Solutions Inc., <br> $1-855-878-1489 ~$ | August <br> 2021 | II |
| NIFEdipine EXTENDED- <br> RELEASE <br> 00904708061 <br> 00904708006 | Failed Dissolution Specification | If you have questions <br> about this recall, Ingenus <br> Pharmaceutical. 1-877- <br> $748-1970$ | July <br> 2021 | II |

* Drug Recall Class

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Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Solifenacin Succinate 06909726102 | CGMP Deviations | If you have questions about this recall, <br> CIPLA, 1-866-604-3268. | $\begin{aligned} & \hline \text { July } \\ & 2021 \end{aligned}$ | II |
| :---: | :---: | :---: | :---: | :---: |
| Xylocaine-MPF with Epinephrine 06332348737 06332348707 | Low out of specification results for epinephrine assay | If you have questions about this recall, Genentech Inc, 1-888-835-2555. | $\begin{aligned} & \text { July } \\ & 2021 \end{aligned}$ | II |
| Xolair (omalizumab) 05024221501 | Failed Stability Specifications | If you have questions about this recall, Fresenius Kabi USA, 1-800-551-7176. | $\begin{aligned} & \hline \text { July } \\ & 2021 \end{aligned}$ | II |
| Buprenorphine and Naloxone Sublingual Film $04778135503$ | Subpotent drug | If you have questions about this recall, Alvogen, Inc. 1-866-770-3024 | $\begin{aligned} & \text { July } \\ & 2021 \end{aligned}$ | II |
| Topotecan Injection 00703471471 | Presence of Particulate Matter | If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800 | $\begin{aligned} & \text { July } \\ & 2021 \end{aligned}$ | I |
| Metformin Hydrochloride Extended-Release 07257803601 | CGMP Deviations | If you have questions about this recall, VIONA PHARMACEUTICALS INC, 1-888-304-5022 | $\begin{aligned} & \text { June } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| DermOtic Oil (fluocinolone acetonide oil) <br> 68791-103-20 | Presence of Foreign Substance | If you have questions about this recall, Hill Dermaceuticals, Inc.,1-407-323-1998 | $\begin{aligned} & \text { June } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |

## * Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

|  |  |  |  |  |
| :--- | :--- | :--- | :--- | :--- |

* Drug Recall Class

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NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| BD ChloraPrep Clear 05436540032 | Non-sterility | If you have questions about this recall, CareFusion 213, LLC, 1-201-847-6800. | $\begin{aligned} & \hline \text { May } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { I } \end{aligned}$ |
| :---: | :---: | :---: | :---: | :---: |
| Estradiol Transdermal System 06896834378 | Defective Delivery System | If you have questions about this recall, Noven Pharmaceuticals Inc, 1-305-253-5099 | $\begin{aligned} & \text { April } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \hline \text { Class } \\ & \text { II } \end{aligned}$ |
| Minivelle 06896866758 | Defective Delivery System | If you have questions about this recall, Noven Pharmaceuticals Inc, 1-305-253-5099 | $\begin{aligned} & \text { April } \\ & 2021 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { Itraconazole } \\ & 05974628230 \end{aligned}$ | Failed Dissolution Specifications | If you have questions about this recall, Jubilant Cadista Pharmaceuticals, Inc., 1-410-860-2836 | $\begin{aligned} & \text { April } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \hline \text { Class } \\ & \text { II } \end{aligned}$ |
| Riomet (metformin hydrochloride oral solution) $01063120602$ | Microbial Contamination of NonSterile Product | If you have questions about this recall, SUN PHARMACEUTICAL INDUSTRIES INC, 1-800-818-4555 | $\begin{aligned} & \text { April } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \hline \text { Class } \\ & \text { II } \end{aligned}$ |
| Cefprozil 06818040201 06818040202 06818040203 | Superpotent Drug | If you have questions about this recall, Lupin Pharmaceuticals Inc., 1-866-587-4617 | $\begin{aligned} & \text { April } \\ & 2021 \end{aligned}$ | Class II |

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NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Guanfacine Extended-Release 06050539281 36050539281 | Cross Contamination | If you have questions about this recall, Apotex Corp., 1866.390-4411. | $\begin{aligned} & \text { April } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| :---: | :---: | :---: | :---: | :---: |
| Mometasone Furoate 00713070185 00713070153 | CGMP Deviaitons | If you have questions about this recall, Cosette Pharmaceuticals, Inc., 1 866.390-4411. | $\begin{aligned} & \text { April } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| Neomycin Sulfate 03982203105 | Failed Stability | If you have questions about this recall, X-Gen Pharmaceuticals Inc., 1-866-390-4411. | $\begin{aligned} & \text { April } \\ & 2021 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { Telmisartan } \\ & 06233208730 \end{aligned}$ | Labeling: Label-mixup | If you have questions about this recall, Alembic Pharmaceuticals Limited, 1-908-393-9604 | $\begin{aligned} & \text { April } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { I } \end{aligned}$ |
| $\begin{aligned} & \text { Progesterone Capsules } \\ & 04359835001 \end{aligned}$ | Failed Dissolution Specifications | If you have questions about this recall, Dr. Reddy's Laboratories, Inc., 1-888-375-3784 | $\begin{aligned} & \text { March } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| Metoclopramide Injection 0703450204 | Chemical contamination | If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800 | $\begin{aligned} & \text { March } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| Famotidine Tablets 06586286099 | Presence of foreign tablets/capsules | If you have questions about this recall, Aurobindo Pharma USA Inc., 1-866-850-2876 | $\begin{aligned} & \text { March } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |

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Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Phenylephrine HCI <br> 02502131501 | Lack of Assurance of Sterility | If you have questions <br> about this recall, Sagent <br> Pharmaceuticals Inc, 1- <br> $866-625-1618$ | March <br> 2021 | Class <br> II |
| :--- | :--- | :--- | :--- | :--- |
| Gabapentin <br> 05038331107 | Failed Impurities/Degradation <br> Specifications | If you have questions <br> about this recall, Akorn, <br> Inc, 1-800-932-5676, <br> Prompt 2 | March <br> 2021 | Class <br> II |
| Nortriptyline HCL <br> 06191985330 | Failed Impurities/Degradation <br> Specifications | If you have questions <br> about this recall, Direct <br> Rx, 1-678-619-5510 | March <br> 2021 | Class II |
| Spironolactone Tablets <br> 63629106701 | Labeling: Label Mix-Up | If you have questions <br> about this recall, <br> BRP Pharmaceuticals, 1- <br> $877-885-0882$ | March <br> 2021 | Class |
| II |  |  |  |  |

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Blue Cross Blue Shield
Blue Care Network
of Michigan

| Omeprazole Delayed Release <br> Capsules <br> 05140712910 | Failed Impurities/Degradation <br> Specifications | If you have questions <br> about this recall, <br> Golden State Medical <br> Supply Inc., 1-805-477- <br> 9866 | March <br> 2021 | Class <br> II |
| :--- | :--- | :--- | :--- | :--- |
| Fludeoxyglucose F 18 <br> 07631833450 | Lack of Assurance of Sterility | If you have questions <br> about this recall, <br> Massachusetts General <br> Hospital PET Center, 1- <br> 617-726-2000 | March <br> 2021 | Class <br> II |
| Irinotecan Hydrochloride <br> 05992371402 | CGMP Deviations | If you have questions <br> about this recall, <br> Areva Pharmaceuticals <br> Inc, 1-855-853-4760 | March <br> 2021 | Class |
| II |  |  |  |  |

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| $\begin{aligned} & \text { Toposar } \\ & 00703565701 \end{aligned}$ | Lack of Assurance of Sterility | If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800 | $\begin{aligned} & \text { March } \\ & 2021 \end{aligned}$ | $\begin{array}{\|l\|} \hline \text { Class } \\ \text { II } \end{array}$ |
| :---: | :---: | :---: | :---: | :---: |
| Metoclopramide Injection 00703450201 <br> 00703450204 | Lack of Assurance of Sterility | If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800. | $\begin{aligned} & \text { March } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \hline \text { Class } \\ & \text { II } \end{aligned}$ |
| $\begin{aligned} & \text { Leucovorin Calcium } \\ & 00703514001 \\ & 00703514501 \end{aligned}$ | Lack of Assurance of Sterility | If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800. | $\begin{aligned} & \text { March } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| MethyIPREDNISolone Acetate 00703003101 00703005101 00703005104 00703004501 00703004301 00703006301 | Lack of Assurance of Sterility | If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800. | $\begin{aligned} & \hline \text { March } \\ & 2021 \end{aligned}$ | Class II |
| Epoprostenol Sodium 00703198501 | Lack of Assurance of Sterility | If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800. | $\begin{aligned} & \text { March } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \hline \text { Class } \\ & \text { II } \end{aligned}$ |
| Sterile Diluent for Epoprostenol Sodium 0703925801 | Lack of Assurance of Sterility | If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800. | $\begin{aligned} & \text { March } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \hline \text { Class } \\ & \text { II } \end{aligned}$ |

## * Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
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| $\begin{aligned} & \text { Desmopressin Acetate } \\ & 00703505101 \\ & 00703505103 \end{aligned}$ | Lack of Assurance of Sterility | If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800. | $\begin{aligned} & \text { March } \\ & 2021 \end{aligned}$ | $\begin{array}{\|l\|} \hline \text { Class } \\ \text { II } \end{array}$ |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \hline \text { Dacarbazine } \\ & 00703507501 \\ & 00703507503 \end{aligned}$ | Lack of Assurance of Sterility | If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800. | $\begin{aligned} & \text { March } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \hline \text { Class } \\ & \text { II } \end{aligned}$ |
| Cisatracurium Besylate 07128871206 | Labeling: Label mix-up | If you have questions about this recall, Meitheal Pharmaceuticals Inc, 1-224-443-4617. | $\begin{aligned} & \text { March } \\ & 2021 \end{aligned}$ | Class I |
| BD ChloraPrep Hi-Lite Orange 2\% w/v chlorhexidine gluconate (CHG) and $70 \% \mathrm{v} / \mathrm{v}$ isopropyl alcohol (IPA), Sterile Solution, 0.01 fl . oz 05436540033 | Non-sterility | If you have questions about this recall, CareFusion 213, LLC, 1-201-847-6800. | $\begin{aligned} & \hline \text { March } \\ & 2021 \end{aligned}$ | Class I |
| ChlroraPrep One-Step 2\% w/v chlorhexidine gluconate (CHG) and $70 \% \mathrm{v} / \mathrm{v}$ isopropyl alcohol (IPA) Non-Sterile Solution-Clear, 0.10 fl . Oz 05436540001 | Non-sterility | If you have questions about this recall, CareFusion 213, LLC, 1-201-847-6800. | $\begin{aligned} & \text { March } \\ & 2021 \end{aligned}$ | Class I |
| ChloraPrep With Tint 2\% w/v chlorhexidine gluconate (CHG) and $70 \% \mathrm{v} / \mathrm{v}$ isopropyl alcohol (IPA) Non-Sterile Solution - HiLite Orange, 0.10 fl . Oz | Microbial Contamination |  |  | Class I |

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NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| 05436540011 |  |  |  |  |
| :--- | :--- | :--- | :--- | :--- |
| BD ChloraPrep Clear, 2\% w/v <br> chlorhexidine gluconate (CHG) <br> and 70\% v/v isopropyl alcohol <br> (IPA) Sterile Solution, 0.10 fl <br> 05436540032 | Non-sterility | If you have questions <br> about this recall, <br> CareFusion 213, LLC, 1- <br> $201-847-6800$. | March <br> 2021 | Class I |
| Meclizine HCI Tablets <br> 05253612901 <br> 05253613301 |  | Failed Dissolution Specifications | If you have questions <br> about this recall, Wilshire <br> Pharmaceuticals, Inc, Ltd, <br> $1-877-495-6856$. | February <br> 2021 |
| Cephalexin for Oral Suspension <br> 06787754568 | CGMP Deviations | If you have questions <br> about this recall, Alkem <br> Laboratories, Ltd, 1-636- <br> $343-5664$. | February <br> 2021 | II |

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| Acetaminophen Injection 05515030701 | Discoloration and failed pH specifications | If you have questions about this recall, AuroMedics Pharma LLC, 1-732-839-9400 ext 2. | $\begin{aligned} & \hline \text { February } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \hline \text { Class } \\ & \text { II } \end{aligned}$ |
| :---: | :---: | :---: | :---: | :---: |
| Ketorolac Tromethamine Injection 06332316201 | Presence of Particulate Matter | If you have questions about this recall, Fresenius Kabi USA, 1-800-551-7176. | $\begin{aligned} & \text { February } \\ & 2021 \end{aligned}$ | Class |
| Nitrofurantoin Capsules 06808444601 06808444611 | Failed Dissolution Specifications | If you have questions about this recall, American Health Packaging, 1-800-707-4621. | $\begin{aligned} & \text { January } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| Chlorhexidine Gluconate 06809402861 | cGMP Deviations | If you have questions about this recall, Precision Dose Inc., 1-800-3979228. | $\begin{aligned} & \hline \text { January } \\ & 2021 \end{aligned}$ | $\begin{array}{\|l} \hline \text { Class } \\ \text { II } \end{array}$ |
| Metformin Hydrochloride 02903305601 | cGMP Deviations | If you have questions about this recall, Nostrum Laboratories Inc, 1-816-308-4941. | $\begin{array}{\|l} \hline \text { January } \\ 2021 \end{array}$ | $\begin{array}{\|l} \hline \text { Class } \\ \text { II } \end{array}$ |
| Paroex (Chlorhexidine Gluconate) $05237602102$ $05237602104$ | cGMP Deviations | If you have questions about this recall, Sunstar Americas, Inc., 1-800-5288527 | $\begin{aligned} & \text { January } \\ & 2021 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Levetiracetam } \\ & 05038324116 \end{aligned}$ | Defective container | If you have questions about this recall, Akorn, Inc, 1-800-932-5676, Prompt 2 | $\begin{aligned} & \text { January } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |

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NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Cephalexin 06787754488 06787754468 | Failed Impurity/Degradation Specifications | If you have questions about this recall, Ascend Laboratories LLC, 1-877-272-7901. | $\begin{aligned} & \text { January } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \text { Esomeprazole Magnesium } \\ & 06909752734 \\ & 06909752834 \\ & 06909752934 \\ & \hline \end{aligned}$ | Cross- contamination with other products | If you have questions about this recall, CIPLA, 1-866-604-3268. | $\begin{aligned} & \text { January } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| $\begin{aligned} & \text { Cephalexin } \\ & 06787754568 \\ & 06909752734 \end{aligned}$ | Failed Impurity/Degradation Specifications | If you have questions about this recall, Ascend Laboratories LLC, 1-877-272-7901. | $\begin{aligned} & \text { January } \\ & 2021 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| Auryxia (ferric citrate) 05992263101 | CGMP Deviations | If you have questions about this recall, Akebia Therapeutics dba Keryx Biopharmaceuticals, Inc, 1-617-871-2098. | $\begin{aligned} & \text { December } \\ & 2020 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| Hydroxyzine Hydrochloride 06043215004 <br> 06043215016 | Failed Impurities/Degradation Specification | If you have questions about this recall, Morton Grove Pharmaceuticals, Inc., 1-847-967-5600, Prompt 4. | $\begin{aligned} & \text { December } \\ & 2020 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| Vumerity (diroximel fumarate) 06440602001 | Failed dissolution specifications | If you have questions about this recall, Biogen MA Inc., 1-844-477-4672. | $\begin{aligned} & \text { December } \\ & 2020 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |

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| Lansoprazole Delayed-Release 06838277177 | Failed Dissolution Specifications | If you have questions about this recall, Zydus Pharmaceuticals (USA) Inc, 1-877-9938779, Prompt 2. | $\begin{aligned} & \text { December } \\ & 2020 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| :---: | :---: | :---: | :---: | :---: |
| Lansoprazole Delayed-Release 06838277277 | Failed Dissolution Specifications | If you have questions about this recall, Zydus Pharmaceuticals (USA) Inc, 1-877-9938779, Prompt 2. | $\begin{aligned} & \text { December } \\ & 2020 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| Sildenafil $04229174801$ | Product mix-up | If you have questions about this recall, AVKARE Inc., 1-855-3613993. | $\begin{aligned} & \text { December } \\ & 2020 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| TraZODONE Hydrochloride 04229183410 | Product mix-up | If you have questions about this recall, Shilpa Medicare Limited, 1-732-637-1971. | $\begin{aligned} & \text { December } \\ & 2020 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| Imatinib Mesylate 07248520330 | GMP Deviations | If you have questions about this recall, Shilpa Medicare Limited, 1-732-637-1971. | $\begin{aligned} & \text { December } \\ & 2020 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |

[^20]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Imatinib Mesylate <br> 07248520290d | GMP Deviations | If you have questions <br> about this recall, <br> Shilpa Medicare Limited, <br> $1-732-637-1971$. | December <br> 2020 | Class <br> II |
| :--- | :--- | :--- | :--- | :--- |
| Anagrelide <br> 01366846201 | Failed Dissolution Specifications | If you have questions <br> about this recall, <br> Torrent Pharma Inc, 1- <br> $888-280-2040$. | December <br> 2020 | Class I |
| Lidocaine HCL 2\%) Topical <br> Anesthetic Hydrogel <br> 06697710703 | Microbial Contamination of Non- <br> Sterile Drug Product | If you have questions <br> about this recall, <br> MPM Medical LLC, 1-800- <br> $232-5512$. | December <br> 2020 | Class I |
| Chlorhexidine Gluconate <br> 007016602715 | Microbial contamination of non-sterile <br> products | If you have questions <br> about this recall, <br> Lohxa LLC, 1-800-641- <br> 5564. | December <br> 2020 | Class I |
| Tizanidine <br> 06787761415 | Failed Dissolution Specifications | If you have questions <br> about this recall, Ascend <br> Laboratories LLC, 1-877- <br> $272-7901$. | December <br> 2020 | Class I |
| Aripiprazole <br> 00904651204 | Failed Dissolution Specifications | If you have questions <br> about this recall, The <br> Harvard Drug Group, 1- <br> $800-875-0123$. | December <br> 2020 | Class II |
| ARIPIPRAZOLE <br> 06042944930 | If you have questions <br> about this recall, Golden <br> State Medical Supply Inc., | December <br> 2020 | Class II |  |

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|  |  | $1-805-477-9866$. |  |  |
| :--- | :--- | :--- | :--- | :--- |
| Paroex (chlorhexidine gluconate) <br> 05237602102 <br> 05237602104 | Microbial Contamination of Non- <br> sterile Product | If you have questions <br> about this recall, Sunstar <br> Americas, Inc., 1-609-751- <br> 9600. | December <br> 2020 | Class I |
| Levetiracetam <br> 07193006352 | Presence of foreign tablet | If you have questions <br> about this recall, Jubilant <br> EYWA PHARMA INC, 1- <br> 609-751-9600. | December <br> 2020 | Class II |
| clomiPRAMINE Hydrochloride <br> 05974671130 | Failed Tablet | If you have questions <br> about this recall, Jubilant <br> Cadista Pharmaceuticals, <br> Inc., 1-800-308-3985. | December <br> 2020 | Class II |
| Lansoprazole Delayed-Release <br> Orally Disintegrating Tablets <br> 06838277177 | Failed Dissolution Specifications | If you have questions <br> about this recall, Zydus <br> Pharmaceuticals, 1-877- <br> $993-8779$. | November <br> 2020 | Class II |
| Lansoprazole Delayed-Release <br> Orally Disintegrating Tablets <br> 06838277277 | Failed Dissolution Specifications | If you have questions <br> about this recall, Zydus <br> Pharmaceuticals, 1-877- <br> $993-8779$. | November <br> 2020 | Class II |
| Mesalamine Delayed-Release <br> Tablets <br> 00591224522 | Failed Dissolution Specifications | If you have questions <br> about this recall, Teva <br> Pharmaceuticals USA,1- | November <br> 2020 | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class $\mathbf{2}$ Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| metformin HCL ER 750 mg 04306390230 <br> 04306390260 | CGMP Deviations | If you have questions about this recall, PD-Rx Pharmaceuticals, Inc., 1-800-299-7379 | $\begin{aligned} & \text { November } \\ & 2020 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| metformin HCL ER 500 mg tablets <br> 07278900930 <br> 07278900960 <br> 07278900993 <br> 07278900990 | CGMP Deviations | If you have questions about this recall, PD-Rx Pharmaceuticals, Inc., 1-800-299-7379 | $\begin{aligned} & \text { November } \\ & 2020 \end{aligned}$ | Class II |
| Metformin Hydrochloride Extended-Release Tablets 02903305501 | CGMP Deviations | If you have questions about this recall, Nostrum Laboratories Inc, 1-816-841-4636. | November $2020$ | Class II |
| Metformin Hydrochloride Extended-Release Tablets 02903305601 | CGMP Deviations | If you have questions about this recall, Nostrum Laboratories Inc, 1-816-841-4636. | $\begin{aligned} & \text { November } \\ & 2020 \end{aligned}$ | Class II |
| Chlorhexidine Gluconate 20\% 05329600120 | Discoloration | If you have questions about this recall, Medichem S.A., 1-201-420-1800 | November $2020$ | Class II |
| Metformin Hydrochloride Extended Release Tablets 04948362401 | CGMP Deviations | If you have questions about this recall, Medichem S.A., 631-7539090; ext. 160 | November $2020$ | Class II |

[^21]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Metformin Hydrochloride - <br> Extended-Release Tablets <br> 04948362309 <br> 04948362301 <br> 04948362350 <br> 04948362310 <br> 04948362350 | CGMP Deviations | If you have questions <br> about this recall, <br> Marksans Pharma <br> Limited, 631-753-9090; <br> ext. 160 | November <br> 2020 | Class II |
| :--- | :--- | :--- | :--- | :--- |
| Metformin HCL E/R 500 mg <br> 07218906490 |  |  |  |  |
| Metformin Hydrochloride <br> Extended-Release Tablets, USP <br> 500 mg <br> 07093430930 <br> 07093430960 <br> 07093430990 <br> 07093430998 | CGMP Deviations | If you have questions <br> about this recall, Direct <br> Rx, 1-678-619-5510 | November <br> 2020 | Class II |
| Metformin Hydrochloride <br> Extended-Release Tablets <br> 07093433430 | CGMP Deviations | If you have questions <br> about this recall, Denton <br> Pharma, Inc., 1-800-722- <br> 0772. | November <br> 2020 | II |
| Mesalamine Delayed-Release <br> Tablets <br> 04229156412 | Failed Dissolution Specifications | If |  |  |

[^22]Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Metformin Hydrochloride Extended Release 750 mg 70518248000 | CGMP Deviations | If you have questions about this recall, RemedyRepack Inc., 1-866-845-3791 | $\begin{aligned} & \text { November } \\ & 2020 \end{aligned}$ | II |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \text { Hydrocortisone butyrate Cream, } \\ & 0.1 \% \\ & 06868227015 \end{aligned}$ | SUBPOTENT DRUG | If you have questions about this recall, <br> Boehringer Ingelheim Pharmaceuticals, Inc., 1-908-927-1400 | $\begin{aligned} & \text { November } \\ & 2020 \end{aligned}$ | II |
| Catapres (clonidine hydrochloride, USP) 0.3 mg 00597001101 | An extraneous peak was observed for dissolution testing. | If you have questions about this recall, Boehringer Ingelheim Pharmaceuticals, Inc., 1-800-243-0127 | $\begin{aligned} & \text { November } \\ & 2020 \end{aligned}$ | II |
| Catapres (clonidine hydrochloride, USP) 0.2 mg 00597000701 | An extraneous peak was observed for dissolution testing. | If you have questions about this recall, Boehringer Ingelheim Pharmaceuticals, Inc., 1-800-243-0127 | $\begin{aligned} & \text { November } \\ & 2020 \end{aligned}$ | II |
| Catapres (clonidine hydrochloride,USP) 0.1 mg 0597000601 | An extraneous peak was observed for dissolution testing. | If you have questions about this recall, Boehringer Ingelheim Pharmaceuticals, Inc., 1-800-243-0127 | $\begin{aligned} & \text { November } \\ & 2020 \end{aligned}$ | II |
| NP Thyroid 120, Thyroid Tablets, USP 2 grain ( 120 mg ) 04219232701 | Subpotent Drug | If you have questions about this recall, Acella Pharmaceuticals, LLC1-800-541-4802 ext 1, | $\begin{aligned} & \text { October } \\ & 2020 \end{aligned}$ | I |

## * Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| NP Thyroid 120, Thyroid Tablets, <br> USP 2 grain (120 mg) <br> 04219232801 | Subpotent Drug | If you have questions <br> about this recall, Acella <br> Pharmaceuticals, LLC1- <br> $800-541-4802$ ext 1, | October <br> 2020 | I |
| :--- | :--- | :--- | :--- | :--- |
| Losartan Pot/HCTZ 50/12.5 mg <br> 06191904090 | CGMP Deviations | If you have questions <br> about this recall, Torrent <br> Pharma Inc 1-800-912- <br> 9561 | October <br> 2020 | II |
| RIOMET ER (metformin <br> hydrochloride for extended- <br> release oral suspension) <br> 01063101917 | CGMP Deviations | If you have questions <br> about this recall, Sun <br> Pharmaceutical Industries, <br> Inc., 1-800-818-4555 | October <br> 2020 | II |
| Eye Itch Relief, Ketotifen <br> Fumarate Ophthalmic Solution <br> 0.035\% <br> 05977992001 | CGMP Deviations | If you have questions <br> about this recall, Akorn, <br> Inc, 1-800-932-5676. | October <br> 2020 | II |
| Potassium Chloride Extended- <br> Release Tablets <br> 06438086006 | Failed Dissolution Specifications | If you have questions <br> about this recall, <br> Strides Inc.1-609-773- <br> 5000. | October <br> 2020 | II |
| Nature-Throid, 1.5 Grain, (97.5 <br> mg), (Thyroid U.S.P. 1.5 gr. <br> (97.5mg)/Liothyronine (T3) <br> 13.5mcg/Levothyroxine (T4) <br> $57 m c g$ | Subpotent Drug |  | If you have questions <br> about this recall, <br> Preferred <br> Pharmaceuticals, Inc. 1- <br> 68788760509 |  |

## * Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Nature-Throid, 1 Grain, 65 mg (Thyroid U.S.P. 1 gr. <br> (65mg)/Liothyronine (T3) <br> $9 \mathrm{mcg} /$ Levothyroxine (T4) 38mcg <br> 68788760409 <br> 68788760401 | Subpotent Drug | If you have questions about this recall, Preferred Pharmaceuticals, Inc. 1-714-777-3729. | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |
| :---: | :---: | :---: | :---: | :---: |
| Nature-Throid, 3/4 Grain (48.75 mg ) Thyroid U.S.P. $3 / 4 \mathrm{gr}$. (48.75 mg /Liothyronine (T3) <br> $6.75 \mathrm{mcg} /$ Levothyroxine (T4) <br> 28.5 mcg <br> 68788686009 | Subpotent Drug | If you have questions about this recall, Preferred Pharmaceuticals, Inc. 1-714-777-3729. | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |
| Nature-Throid, 1/2 Grain, 32.5 mg (Thyroid U.S.P. $1 / 2 \mathrm{gr}$. (32.5 mg )/Liothyronine (T3) $4.5 \mathrm{mcg} / \mathrm{Levothyroxine}$ (T4) 19 mch 68788928301 | Subpotent Drug | If you have questions about this recall, Preferred Pharmaceuticals, Inc. 1-714-777-3729. | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |
| WP Thyroid, Westhroid Pure, 2 Grain (130 mg) Thyroid USP 06472759502 06472759504 06472759505 06472759506 06472759501 | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |
| WP Thyroid, Westhroid Pure, 1.75 Grain ( 113.75 mg ) Thyroid USP $06472761502$ | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class $\mathbf{2}$ Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| 06472761504 06472761505 06472761506 06472761501 |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| WP Thyroid, Westhroid Pure, 1.5 Grain ( 97.5 mg ) Thyroid USP 06472758502 <br> 06472758504 <br> 06472758505 <br> 06472758506 <br> 06472758501 | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |
| WP Thyroid, Westhroid Pure, 1.25 Grain ( 81.25 mg ) Thyroid USP <br> 06472760502 <br> 06472760504 <br> 06472760505 <br> 06472760506 <br> 06472760501 | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | September $2020$ | II |
| WP Thyroid, Westhroid Pure, 1 Grain (65 mg) Thyroid USP 06472757502 <br> 06472757504 <br> 06472757505 <br> 06472757506 <br> 06472757501 | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |

## * Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| WP Thyroid, Westhroid Pure, 1/2 Grain ( 32.5 mg ) Thyroid USP 06472755502 <br> 06472755504 <br> 06472755505 <br> 06472755506 <br> 06472755501 | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |
| :---: | :---: | :---: | :---: | :---: |
| WP Thyroid, Westhroid Pure, Thyroid USP, [liothyronine (T3) 2.25 mcg and levothyroxine (T4) 9.5 mcg 06472754504 <br> 06472754505 <br> 06472754506 <br> 06472754501 <br> 06472754502 | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | $\begin{aligned} & \hline \text { September } \\ & 2020 \end{aligned}$ | II |
| Nature-Throid, Thyroid USP [liothyronine (T3) 27 mcg and levothyroxine (T4) 114 mcg 06472733124 <br> 06472733125 <br> 06472733126 <br> 06472733121 <br> 06472733122 | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |
| Nature-Throid, Thyroid USP [liothyronine (T3) 22.5 mcg and levothyroxine (T4) 95 mcg $06472733104$ $06472733105$ | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

Blue Cross
Blue Shield
Blue Care Network
of Michigan

| $\begin{aligned} & 06472733106 \\ & 06472733101 \\ & 06472733102 \end{aligned}$ |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Nature-Throid, Thyroid USP [liothyronine (T3) 20.25 mcg and levothyroxine (T4) 85.5 mcg 06472733094 <br> 06472733095 <br> 06472733096 <br> 06472733091 <br> 06472733092 | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | September $2020$ | II |
| Nature-Throid, Thyroid USP [liothyronine (T3) 18 mcg and levothyroxine (T4) 76 mcg 06472733084 <br> 06472733085 <br> 06472733086 <br> 06472733081 <br> 06472733082 | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |
| Nature-Throid, Thyroid USP [liothyronine (T3) 15.75 mcg and levothyroxine (T4) 66.5 mcg 06472733074 <br> 06472733075 <br> 06472733076 <br> 06472733071 <br> 06472733072 | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Nature-Throid, Thyroid USP [liothyronine (T3) 13.5 mcg and levothyroxine (T4) 57 mcg 06472733054 <br> 06472733055 <br> 06472733056 <br> 00472733051 <br> 06472733052 | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |
| :---: | :---: | :---: | :---: | :---: |
| Nature-Throid, Thyroid USP [liothyronine (T3) 11.25 mcg and levothyroxine (T4) 47.5 mcg , 1.25 Grain ( 81.25 mg ) <br> 06472733034 <br> 06472733035 <br> 06472733036 <br> 06472733031 <br> 06472733032 | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |
| Nature-Throid, Thyroid USP [liothyronine (T3) 4.5 mcg and levothyroxine (T4) 19 mcg 06472732994 <br> 06472732995 <br> 06472732996 <br> 06472732991 <br> 06472732992 | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |

## * Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class $\mathbf{2}$ Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Nature-Throid, Thyroid USP [liothyronine (T3) 6.75 mcg 06472733024 06472733025 06472733026 06472733021 06472733022 | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |
| :---: | :---: | :---: | :---: | :---: |
| Nature-Throid, Thyroid USP [liothyronine (T3) 9 mcg and levothyroxine (T4) 38 mcg 06472733004 <br> 06472733005 <br> 06472733006 <br> 06472733001 <br> 06472733002 | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |
| Nature Throid, 1.5 Grain (97.5 mg ) Thyroid USP [Liothyronine <br> (T3) 13.5 mcg , Levothyroxine <br> (T4) 57 mcg <br> 06472733054 <br> 06472733055 <br> 06472733056 <br> 06472733051 <br> 06472733051 | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | I |
| Nature Throid, 2 Grain (130 mg) Thyroid USP [Liothyronine (T3) 18 mcg , Levothyroxine (T4) 76 mcg 06472733084 | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | I |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class $\mathbf{2}$ Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| 06472733081 <br> Nature Throid 125 Grain (81 25 |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Nature Throid, 1.25 Grain (81.25 mg ) Thyroid USP [Liothyronine (T3) 11.25 mcg , Levothyroxine (T4) 47.5 mcg 06472733031 | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | I |
| Nature Throid, 3/4 Grain (48.75 mg ) Thyroid USP [Liothyronine (T3) 6.75 mcg , Levothyroxine (T4) 28.5 mcg 06472733021 06472733022 | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | I |
| Nature Throid, 1/2 Grain (32.5 mg ) Thyroid USP [Liothyronine (T3) 4.5 mcg , Levothyroxine (T4) 19 mcg 06472732996 06472732991 | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | 1 |
| WP Thyroid, Westhroid Pure, 1/2 Grain ( 32.5 mg ) Thyroid USP [Liothyronine (T3) 4.5 mcg , Levothyroxine (T4) 19 mcg 06472755504 <br> 06472755506 <br> 06472755501 | Subpotent Drug | If you have questions about this recall, RLC Labs Inc., 1-877-797-7997 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | I |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

Blue Cross
Blue Cross
Blue Care Network
of Michigan

| Sulfamethoxazole and Trimethoprim Tablets | Presence of Foreign Substance: product complaints were received by the firm for the presence of metal wire in the tablet(s). | If you have questions about this recall, Aurobindo Pharma USA Inc., 1-866-850-2876 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |
| :---: | :---: | :---: | :---: | :---: |
| BusPIRone Hydrochloride Tablets | Failed Impurity /Degradation Specifications | If you have questions about this recall, Par Pharmaceutical Inc., 1-845-573-5500 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |
| Buprenorphine HCl Injection | Sub-potent Drug: Out-of-Specification assay results found at 3-month stability testing. | If you have questions about this recall, West-Ward Pharmaceuticals , 1-877-845-0689. | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |
| Metformin Hydrochloride Extended-Release Tablets 07638512810 | CGMP Deviations | If you have questions about this recall, <br> BAYSHORE PHARMACEUTICALS, 1- <br> 877-372-6093 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |
| Metformin Hydrochloride Extended-Release Tablets 07638512901 | CGMP Deviations | If you have questions about this recall, <br> BAYSHORE <br> PHARMACEUTICALS, 1- <br> 877-372-6093 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |
| Cephalexin for Oral Suspension 00093417773 | Labeling: Label Error on Declared Strength | If you have questions about this recall, Teva Pharmaceuticals USA,1-888-838-2872 ext 5 | $\begin{aligned} & \text { September } \\ & 2020 \end{aligned}$ | II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Lidocaine Patch 5\% <br> 00591352530 | Labeling: Incorrect or Missing Lot <br> and/or Exp date on the individual <br> transdermal pouches but not in the <br> carton | If you have questions <br> about this recall, Teva <br> Pharmaceuticals USA,1- <br> $888-838-2872$ ext 5 | August <br> 2020 | II |
| :--- | :--- | :--- | :--- | :--- |
| Mibelas 24 Fe <br> 06818091113 | Failed Impurities/Degradation <br> Specifications: Out of specification <br> result observed in related substance <br> test | If you have questions <br> about this recall, Lupin <br> Pharmaceuticals Inc.1- <br> $410-576-2000$ ext 3 | August <br> 2020 | II |
| Nystatin Oral Suspension <br> 00121061016 | Subpotent drug: Out of specification <br> for assay at the 15-month test <br> interval | If you have questions <br> about this recall, PAI <br> Holdings, LLC. dba <br> Pharmaceutical <br> Associates Inc1-864-277- <br> 7282 ext 0 | August <br> 2020 | II |
| Ear Pain MD Pain Relief Drops <br> 07242900708 | cGMP Deviations. | If you have questions <br> about this recall, Eosera, <br> Inc.1-844-732-7929 | August <br> 2020 | II |
| Ear Pain MD Pain Relief Drops <br> 07242900722 | cGMP Deviations. | If you have questions <br> about this recall, Eosera, <br> Inc.1-844-732-7929 | August <br> 2020 | II |

[^23]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| $\text { (i) (\$) } \begin{aligned} & \text { Blue Cross } \\ & \text { Bue Shield } \\ & \text { Biue Care } \\ & \text { of Michigan Network } \end{aligned}$ |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Heparin Sodium 70004065046 | Subpotent Drug: Out-of-Specification potency results at the 30-day stability timepoint. | If you have questions about this recall, SCA Pharmaceuticals1-877-550-5059 | August 2020 | II |
| BD ChloraPrep Clear 5436540032213 | Non-Sterility: Product is being recalled due to presence of Aspergillus Penicilloides | If you have questions about this recall, CareFusion LLC1-201-847-6800. | $\begin{aligned} & \text { August } \\ & 2020 \end{aligned}$ | I |
| ChloraPrep With Tint 05436540011 | Non-Sterility: Product is being recalled due to presence of Aspergillus Penicilloides | If you have questions about this recall, CareFusion LLC1-201-847-6800. | $\begin{aligned} & \text { August } \\ & 2020 \end{aligned}$ | I |
| ChlroraPrep One-Step 05436540001 | Non-Sterility: Product is being recalled due to presence of Aspergillus Penicilloides | If you have questions about this recall, CareFusion LLC1-201-847-6800. | $\begin{aligned} & \text { August } \\ & 2020 \end{aligned}$ | I |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

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| DDAVP Nasal <br> 05556625000 | Superpotent Drug | If you have questions <br> about this recall, Ferring <br> Pharmaceuticals Inc1- <br> $888-337-7464$ ext2 | August <br> 2020 |  |
| :--- | :--- | :--- | :--- | :--- |
|  |  | Superpotent Drug | I |  |
| Desmopressin Acetate <br> 06991850105 | If you have questions <br> about this recall, Ferring <br> Pharmaceuticals Inc1- <br> $888-337-7464$ ext2 | August <br> 2020 | I |  |
| STIMATE (desmopressin <br> acetate) <br> 00053687100 | Superpotent Drug | If you have questions <br> about this recall, Ferring <br> Pharmaceuticals Inc1- <br> $888-337-7464$ ext2 | August <br> 2020 | I |
| Hydrochlorothiazide <br> 16729018317 | CGMP Deviations: Light sensitive <br> drug products repackaged in <br> transparent/partially transparent <br> pouches | If you have questions <br> about this recall, Calvin <br> Scott \& Company, Inc.1- <br> $800-545-6545$. | August <br> 2020 | II |

## * Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| $\begin{aligned} & \text { Hydrochlorothiazide (HCTZ) } \\ & 29300012810 \end{aligned}$ | CGMP Deviations: Light sensitive drug products repackaged in transparent/partially transparent pouches | If you have questions about this recall, Calvin Scott \& Company, Inc.1-800-545-6545. | $\begin{aligned} & \text { August } \\ & 2020 \end{aligned}$ | II |
| :---: | :---: | :---: | :---: | :---: |
| Hydrochlorothiazide 16729018417 | CGMP Deviations: Light sensitive drug products repackaged in transparent/partially transparent pouches | If you have questions about this recall, Calvin Scott \& Company, Inc.1-800-545-6545. | $\begin{aligned} & \text { August } \\ & 2020 \end{aligned}$ | II |
| Thyroid 1 Grain 64727330002 | CGMP Deviations: Light sensitive drug products repackaged in transparent/partially transparent pouches | If you have questions about this recall, Calvin Scott \& Company, Inc.1-800-545-6545. | $\begin{aligned} & \text { August } \\ & 2020 \end{aligned}$ | II |
| Thyroid 1 Gr 64727330001 | CGMP Deviations: Light sensitive drug products repackaged in transparent/partially transparent pouches | If you have questions about this recall, Calvin Scott \& Company, Inc.1-800-545-6545. | $\begin{aligned} & \text { August } \\ & 2020 \end{aligned}$ | II |
| Thyroid Neutral 64727330802 | CGMP Deviations: Light sensitive drug products repackaged in transparent/partially transparent pouches | If you have questions about this recall, Calvin Scott \& Company, Inc.1-800-545-6545. | $\begin{aligned} & \text { August } \\ & 2020 \end{aligned}$ | II |
| Topiramate 47335070713 | CGMP Deviations: Light sensitive drug products repackaged in transparent/partially transparent pouches | If you have questions about this recall, Calvin Scott \& Company, Inc.1-800-545-6545. | $\begin{aligned} & \text { August } \\ & 2020 \end{aligned}$ | II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
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NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

|  |  |  |  |  |
| :--- | :--- | :--- | :--- | :--- |
| Phendimetrazine <br> 00702007710 | CGMP Deviations: Light sensitive <br> drug products repackaged in <br> transparent/partially transparent <br> pouches | If you have questions <br> about this recall, Calvin <br> Scott \& Company, Inc.1- <br> $800-545-6545$. | August <br> 2020 | II |
|  | CGMP Deviations: Light sensitive <br> drug products repackaged in <br> transparent/partially transparent <br> pouches | If you have questions <br> about this recall, Calvin <br> Scott \& Company, Inc.1- <br> $800-545-6545$. | August <br> 2020 | II |
| Phendimetrazine <br> 69543041011 | CGMP Deviations: Light sensitive <br> drug products repackaged in <br> transparent/partially transparent <br> pouches | If you have questions <br> about this recall, Calvin <br> Scott \& Company, Inc.1- <br> $800-545-6545$. | August <br> 2020 | Class <br> II |
| Phendimetrazine <br> 69543040911 | Labeling: Label Mix Up: bottle labeled <br> to contain Prednisone Tablets | If you have questions <br> about this recall, Lannett <br> Company, Inc.1-215-333- <br> 9000 | August <br> 2020 | Class <br> II |
| Prednisone <br> 00527293137 | Failed Stability Specifications: Out of <br> Specification result for enzyme <br> activity levels noted during routine <br> stability testing | If you have questions <br> about this recall, Sanofi- <br> Aventis U.S. LLC1-800- <br> $981-2491 ~ e x t ~ 2 ~$ | August <br> 2020 | Class II |
| Elitek (rasburicase) <br> 00024515175 |  |  |  |  |

## * Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.
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| $\begin{aligned} & \hline \text { Mibelas } 24 \text { Fe } \\ & 06818091111 \end{aligned}$ | Failed Impurities/Degradation Specifications: Out of specification result observed in related substance test | If you have questions about this recall, Lupin Pharmaceuticals Inc.1-410-576-2000 ext 3 | $\begin{aligned} & \text { August } \\ & 2019 \end{aligned}$ | $\begin{aligned} & \text { Class } \\ & \text { II } \end{aligned}$ |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \hline \text { Daptomycin } \\ & 06745781350 \end{aligned}$ | Presence of Particulate Matter | If you have questions about this recall, Mylan Institutional LLC, 1-800-796-9526 ext 1. | $\begin{aligned} & \text { August } \\ & 2020 \end{aligned}$ | Class II |
| Fentanyl Citrate 00409909412 | Lack of Assurance of Sterility | If you have questions about this recall, Pfizer Inc.,1-800-438-1985. | $\begin{aligned} & \text { August } \\ & 2020 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { Lisinopril } \\ & 06800133408 \end{aligned}$ | Presence of Foreign Tablets/Capsules | If you have questions about this recall, Lupin Pharmaceuticals Inc., 1-866-587-4617 ext 2. | $\begin{aligned} & \text { August } \\ & 2020 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Cefdinir } \\ & 06818072320 \end{aligned}$ | Superpotent Drug | If you have questions about this recall, Lupin Pharmaceuticals Inc., 1-866-587-4617 ext 2. | $\begin{aligned} & \text { July } \\ & 2020 \end{aligned}$ | Class II |
| Metformin HCl Extended Release Tablets | CGMP Deviation | If you have questions about this recall, Preferred Pharmaceuticals, Inc, 1-714-777-3729. | $\begin{aligned} & \text { July } \\ & 2020 \end{aligned}$ | Class II |
| Metformin Hydrochloride Extended-Release 00904579461 | CGMP Deviation | If you have questions about this recall, The Time-Cap, Labs Inc, 1-631-753-9090 ext 160 | $\begin{aligned} & \hline \text { July } \\ & 2020 \end{aligned}$ | Class II |

* Drug Recall Class

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| Clozapine Tablets 06586284605 | Presence of foreign tablet | If you have questions about this recall, Aurobindo Pharma USA, Inc, 1-866-850-2876 | $\begin{aligned} & \text { July } \\ & 2020 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Metformin Hydrochloride Extended-release Tablets 06818033607 | CGMP Deviations | If you have questions about this recall, Lupin Pharmaceuticals Inc., 1-866-587-4617. | $\begin{aligned} & \hline \text { July } \\ & 2020 \end{aligned}$ | Class II |
| Aripiprazole Tablets 06233209730 | Labeling: Label mix up | If you have questions about this recall, Alembic Pharmaceuticals Limited, 1-908-393-9604. | $\begin{aligned} & \text { July } \\ & 2020 \end{aligned}$ | Class II |
| Nystatin Cream 00316022130 00316022115 | Subpotent Drug | If you have questions about this recall, Crown Laboratories, Inc., 1-800-877-8869. | $\begin{aligned} & \text { July } \\ & 2020 \end{aligned}$ | Class II |
| metFORMIN HCL ER 07278900930 07278900960 07278900990 07278900993 49483062301 | CGMP Deviations | If you have questions about this recall, Amneal Pharmaceuticals., 1-833-582-0812. | $\begin{aligned} & \text { July } \\ & 2020 \end{aligned}$ | Class II |
| Metformin Hydrochloride Extended-Release Tablets 04948362301 | CGMP Deviations | If you have questions about this recall, TimeCap Labs,1-631-753-9090 ext 160 | $\begin{aligned} & \hline \text { July } \\ & 2020 \end{aligned}$ | Class II |

[^24]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Metformin Hydrochloride <br> Extended - Release Tablets <br> 06203757101 <br> 06203757110 <br> 06203757701 <br> 06203757710 | CGMP Deviations | If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800. | $\begin{array}{\|l\|} \hline \text { July } \\ 2020 \end{array}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Metformin Hydrochloride Extended-Release Tablets 53746017801 06516217809 06516217850 06516217810 53746017901 06516217910 00537460178 53746017805 53746017810 65162017811 00537460179 65162017901 | CGMP Deviations | If you have questions about this recall, Amneal Pharmaceuticals, 1-833-582-0812 | $\begin{aligned} & \hline \text { July } \\ & 2020 \end{aligned}$ | Class II |
| Heparin Sodium 70004065046 | Subpotent Drug | If you have questions about this recall, SCA Pharmaceuticals, 1-877-550-5059. | $\begin{aligned} & \text { June } \\ & 2020 \end{aligned}$ | Class II |
| Irinotecan HCL Injection 00143958301 | Defective Container | If you have questions about this recall, WestWard Pharmaceutical Corp, 1-877-845-0689. | $\begin{aligned} & \text { June } \\ & 2020 \end{aligned}$ | Class II |

* Drug Recall Class

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| Irinotecan HCL Injection 00143970101 | Defective Container | If you have questions about this recall, WestWard Pharmaceutical Corp, 1-877-845-0689. | $\begin{array}{\|l\|} \hline \text { June } \\ 2020 \end{array}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| metFORMIN HCL ER 04306342830 04306342860 04306342890 04306342898 04306342893 53746017805 | CGMP Deviations | If you have questions about this recall, Amneal Pharmaceuticals, 1-833-582-0812 | $\begin{aligned} & \hline \text { June } \\ & 2020 \end{aligned}$ | Class II |
| Metformin Hydrochloride Extended-Release Tablets 05026853115 04229161090 04229161018 04229161036 04229161010 | CGMP Deviations | If you have questions about this recall, AVKARE Inc., 1-931-292-6222. | $\begin{aligned} & \text { June } \\ & 2020 \end{aligned}$ | Class II |
| Metformin Hydrochloride Extended-Release Tablets 04229161190 <br> 04229161118 <br> 04229161150 | CGMP Deviations | If you have questions about this recall, AVKARE Inc., 1-931-292-6222. | $\begin{aligned} & \text { June } \\ & 2020 \end{aligned}$ | Class II |
| Metformin Hydrochloride Extended-Release Tablets 06050502601 | CGMP Deviations | If you have questions about this recall, Apotex Inc., 1-800-268-4623. | $\begin{aligned} & \text { June } \\ & 2020 \end{aligned}$ | Class II |

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| Gaviscon Liquid Antacid Extra <br> Strength, Cool Mint <br> 00135009541 | Labeling: Label lacks warning | If you have questions <br> about this recall, <br> Glaxosmithkline <br> Consumer Healthcare <br> Holdings Inc., 1-800-743- <br> 4014 | June <br> 2020 | Class II |
| :--- | :--- | :--- | :--- | :--- |
| Gaviscon Extra Strength Liquid <br> Antacid Extra Strength Cherry <br> 00135057441 | Labeling: Label lacks warning | If you have questions <br> about this recall, <br> Glaxosmithkline <br> Consumer Healthcare <br> Holdings Inc., 1-800-743- <br> 4014 | June <br> 2020 | Class II |
| Gaviscon Regular Strength <br> Liquid Antacid Cool Mint <br> 00135009441 | Labeling: Label lacks warning | If you have questions <br> about this recall, <br> Glaxosmithkline <br> Consumer Healthcare <br> Holdings Inc., 1-800-743- <br> 4014 | June <br> 2020 | Class II |
| Gaviscon Regular Strength <br> Liquid Antacid Cool Mint <br> 00135009442 | Labeling: Label lacks warning | If you have questions <br> about this recall, <br> Glaxosmithkline <br> Consumer Healthcare <br> Holdings Inc., 1-800-743- <br> 4014 | June <br> 2020 | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
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Class $\mathbf{3}$ Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| $\begin{aligned} & \text { Lisinopril } \\ & 06818051303 \end{aligned}$ | Product Mix Up | If you have questions about this recall, Lupin Pharmaceuticals Inc., 1-866-587-4617. | $\begin{aligned} & \hline \text { June } \\ & 2020 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Unasyn (ampicillin sodium/sulbacatam) $00049001381$ $00049001383$ | Presence of Particulate Matter | If you have questions about this recall, Pfizer Inc., 1-800-879-3477. | $\begin{aligned} & \text { June } \\ & 2020 \end{aligned}$ | Class II |
| Doxycycline Hyclate 06258469311 | Failed dissolution specification | If you have questions about this recall, American Health Packaging, 1-614-492-8177. | $\begin{aligned} & \text { June } \\ & 2020 \end{aligned}$ | Class II |
| Ketorolac Tromethamine 06332316200 <br> 06332316203 | Presence of Particulate Matter | If you have questions about this recall, Fresenius Kabi USA, LLC, 1-847-550-2300. | $\begin{aligned} & \hline \text { June } \\ & 2020 \end{aligned}$ | Class II |
| Doxycycline Hyclate 00904043004 00904043006 | Failed Dissolution Specification | If you have questions about this recall, The Harvard Drug Group, 1-732-542-1191. | $\begin{aligned} & \hline \text { June } \\ & 2020 \end{aligned}$ | Class II |
| Dextroamphetamine Saccharate, Amphetamine Aspartate, Dextroamphetamine Sulfate and Amphetamine Sulfate 00555097102 00555097302 00555077702 | Some bottles may contain mixed strengths of the product. | If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800. | $\begin{aligned} & \text { June } \\ & 2020 \end{aligned}$ | Class II |

[^26]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| $\begin{aligned} & \text { Estriol USP Micronized } \\ & 71092997702 \end{aligned}$ | Failed impurities/ degradation specifications | If you have questions about this recall, TRIOVA PHARMACEUTICALS LLC, 1-539-777-0720. | $\begin{aligned} & \text { June } \\ & 2020 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \text { NP Thyroid } 90 \\ & 04219233101 \end{aligned}$ | Superpotent Drug | If you have questions about this recall, Acella Pharmaceuticals, LLC 1-800-541-4802. | $\begin{aligned} & \text { June } \\ & 2020 \end{aligned}$ | Class I |
| $\begin{aligned} & \text { NP Thyroid } 60 \\ & 04219233001 \end{aligned}$ | Superpotent Drug | If you have questions about this recall, Acella Pharmaceuticals, LLC 1-800-541-4802. | $\begin{aligned} & \text { June } \\ & 2020 \end{aligned}$ | Class I |
| $\begin{aligned} & \hline \text { NP Thyroid } 30 \\ & 04219232901 \end{aligned}$ | Superpotent Drug | If you have questions about this recall, Acella Pharmaceuticals, LLC 1-800-541-4802. | $\begin{aligned} & \text { June } \\ & 2020 \end{aligned}$ | Class I |
| Doxycycline Hyclate tablets 05528986606 05528986610 05528986614 05528986620 05528986628 05528986630 05528986660 05528986698 05528986671 05528986687 05528986674 | Failed dissolution specifications | If you have questions about this recall, PD-Rx Pharmaceuticals, Inc., 1-800-299-7379. | $\begin{aligned} & \text { June } \\ & 2020 \end{aligned}$ | Class II |

* Drug Recall Class

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| Blue Cross Blue Shield Blue Care Network of Michigan |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Doxycycline Hyclate Tablets 00143211250 <br> 00143211205 | Failed dissolution specification | If you have questions about this recall, WestWard Columbus Inc, 1-877-845-0689. | $\begin{aligned} & \hline \text { June } \\ & 2020 \end{aligned}$ | Class II |
| Lactated Ringers 0409795309 | Presence of Particulate Matter | If you have questions about this recall, Hospira Inc., 1-877-946-7747. | $\begin{aligned} & \text { May } \\ & 2020 \end{aligned}$ | Class II |
| Aloprim (allopurinol sodium) 06745718750 | Discoloration | If you have questions about this recall, Mylan Pharmaceuticals Inc., 1-551-233-2700. | $\begin{aligned} & \text { May } \\ & 2020 \end{aligned}$ | Class II |
| Infuvite PEDiatric Pharmacy 05464356470 | Defective container | If you have questions about this recall, Sandoz, Inc, 1-609-627-8500 | $\begin{aligned} & \text { May } \\ & 2020 \end{aligned}$ | Class II |
| Epinephrine Injection 00093598627 | CGMP Deviations | If you have questions about this recall, Teva Pharmaceuticals USA, 1-888-838-2872. | $\begin{aligned} & \text { May } \\ & 2020 \end{aligned}$ | Class II |
| $\begin{array}{\|l\|} \hline \text { Nizatidine } \\ 06084630115 \end{array}$ | CGMP Deviations | If you have questions about this recall, Amneal Pharmaceuticals of New York, 1-631-952-0214 | $\begin{aligned} & \text { May } \\ & 2020 \end{aligned}$ | Class II |
| Ceftazidime 00264314511 | Failed Stability Specifications | If you have questions about this recall, Braun Medical Inc., 1-949-6602000 | $\begin{aligned} & \text { May } \\ & 2020 \end{aligned}$ | Class II |

* Drug Recall Class

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| Lisinopril Tablets <br> 68180098103 | Product Mix-Up | If you have questions <br> about this recall, Lupin <br> Pharmaceuticals Inc., 1- <br> $866-587-4617$ | April <br> 2020 | Class II |
| :--- | :--- | :--- | :--- | :--- |
| Cefixime for Oral Suspension <br> 06818040501 | Subpotent Drug | If you have questions <br> about this recall, Lupin <br> Pharmaceuticals Inc., 1- <br> $866-587-4617$ | April <br> 2020 | Class II |
| Daytrana <br> 06896855553 | Defective Delivery System | If you have questions <br> about this recall, Noven <br> Therapeutics, LLC, 1-305- <br> $253-5099$. | April <br> 2020 | Class II |
| Daytrana <br> 06896855543 | Defective Delivery System | If you have questions <br> about this recall, Noven <br> Therapeutics, LLC, 1-305- <br> $253-5099$. | April <br> 2020 | Class II |
| Daytrana <br> 06896855523 | If you have questions <br> about this recall, Noven <br> Therapeutics, LLC, 1-305- <br> $253-5099$. | April <br> 2020 | Class II |  |
| Acetaminophen and Codeine <br> Phosphate <br> 01310705801 | CGMP Deviations | If you have questions <br> about this recall, <br> Aurobindo Pharma USA, <br> Inc, 1-866-850-2876. | April <br> 2020 | Class II |
| Acetaminophen and Codeine <br> Phosphate <br> 01310705999 | CGMP Deviations | If you have questions <br> about this recall, <br> Aurobindo Pharma USA, <br> Inc, 1-866-850-2876. | April <br> 2020 | Class II |

## * Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
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| Acetaminophen and Codeine Phosphate 01310706001 | CGMP Deviations | If you have questions about this recall, Aurobindo Pharma USA, Inc, 1-866-850-2876. | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Gabapentin Capsules 06586219899 | CGMP Deviations | If you have questions about this recall, Aurobindo Pharma USA, Inc, 1-866-850-2876. | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |
| Levetiracetam Tablet 06586224708 | CGMP Deviations | If you have questions about this recall, Aurobindo Pharma USA, Inc, 1-866-850-2876. | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Simvastatin } \\ & 06586205390 \end{aligned}$ | CGMP Deviations | If you have questions about this recall, Aurobindo Pharma USA, Inc, 1-866-850-2876. | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { Mirtazapine } \\ & 01310703134 \\ & 06586219899 \end{aligned}$ | CGMP Deviations | If you have questions about this recall, Aurobindo Pharma USA, Inc, 1-866-850-2876. | $\begin{aligned} & \hline \text { April } \\ & 2020 \end{aligned}$ | Class II |
| Phentermine Hydrochloride 06586205390 <br> 06586219899 | CGMP Deviations | If you have questions about this recall, Aurobindo Pharma USA, Inc, 1-866-850-2876. | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |
| Oxycodone and Acetaminophen 01310704601 <br> 6586219899 | CGMP Deviations | If you have questions about this recall, Aurobindo Pharma USA, Inc, 1-866-850-2876. | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |

## * Drug Recall Class

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| Losartan Potassium 06042931630 06042931690 06042931610 | CGMP Deviations | If you have questions about this recall, Golden State Medical Supply Inc., 1-805-477-9866 Ext 4. | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Estradiol 06327599008 06327599005 06327599004 | CGMP Deviations | If you have questions about this recall, B \& B Pharmaceuticals, Inc., 1-800-499-3100. | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |
| Acetaminophen and Codeine Phosphate 300/30mg $52959000310$ <br> 52959000312 <br> 52959000314 <br> 52959000315 <br> 52959000316 <br> 52959000320 <br> 52959000330 <br> 52959000360 | CGMP Deviations | If you have questions about this recall, H.J. Harkins Co, 1-805-9291333 | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |
| succinylcholine Chloride 07101934104 | lack of sterility assurance. | If you have questions about this recall, PharMEDium Services, 1-800-523-7749 Option 1 | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |
| rocuronium Bromide 07101932110 07101932105 | lack of sterility assurance. | If you have questions about this recall, PharMEDium Services, 1-800-523-7749 Option 1 | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |

[^27]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

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| Phytonadione Injectable Emulsion $04359840511$ | Defective Container | If you have questions about this recall, Dr. Reddy's Laboratories Inc., 1-888-375-3784 | $\begin{aligned} & \hline \text { April } \\ & 2020 \end{aligned}$ | Class I |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \hline \text { nICARdipine HCI } \\ & 07101920601 \end{aligned}$ | lack of sterility assurance. | If you have questions about this recall, PharMEDium Services, 1-800-523-7749 Option 1 | $\begin{array}{l\|} \hline \text { April } \\ 2020 \end{array}$ | Class II |
| Losartan Potassium tablets 00591374500 <br> 00591374600 <br> 00591374700 | CGMP Deviations | If you have questions Teva Pharmaceuticals USA, 1-888-483-8279 | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |
| Losartan Potassium tablets 02315564503 02315564509 02315564510 02315564509 02315564603 02315564609 02315564610 | CGMP Deviations | If you have questions Avet Pharmaceuticals, Inc., 1-800-967-5952 Option 1 | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |
| Lisinopril Tablets 06818098201 | Presence of Foreign Tablet/ Capsule | If you have questions about this recall, Lupin Pharmaceuticals Inc., 1-800-399-2561 | $\begin{array}{l\|} \hline \text { April } \\ 2020 \end{array}$ | Class II |

[^28]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| HYDROmorphone in 0.9\% Sodium Chloride HCl 06155316644 | lack of sterility assurance. | If you have questions about this recall, PharMEDium Services, 1-800-523-7749 Option 1 | $\begin{array}{l\|} \hline \text { April } \\ 2020 \end{array}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Glycopyrrolate Tablets 49884006501 | lack of sterility assurance. | If you have questions about this recall, Par Pharmaceutical Inc., 1-845-573-5500 | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |
| fentaNYL Citrate Injection 06155330633 06155330328 | lack of sterility assurance. | If you have questions about this recall, PharMEDium Services, LLC, 1-800-523-7749 Option 1 | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |
| fentaNYL Citrate 06155311350 06155367244 | lack of sterility assurance. | If you have questions about this recall, PharMEDium Services, LLC, 1-800-523-7749 Option 1 | $\begin{array}{l\|} \hline \text { April } \\ 2020 \end{array}$ | Class II |
| ePHEDrine Sulfate <br> 07103000312 <br> 07103000309 <br> 07103000302 <br> 07103000109 | lack of sterility assurance. | If you have questions about this recall, PharMEDium Services, LLC, 1-800-523-7749 Option 1 | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |
| C-*Vancomycin Opthal 14 mg drops $67457034001$ | Lack of Assurance of Sterility | If you have questions about this recall, Medical Center Pharmacy, Inc., 1-800-523-7749 | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |

## * Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| C-*Gentamicin/Bacitracin Bladder Irrigation in N.S., 250 mL 63323001002 | Lack of Assurance of Sterility | If you have questions about this recall, Medical Center Pharmacy, Inc., 1-800-523-7749 | $\begin{array}{l\|} \hline \text { April } \\ 2020 \end{array}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| C-*Albumin Eye Drop 10\% S 44206025105 | Lack of Assurance of Sterility | If you have questions about this recall, Medical Center Pharmacy, Inc., 1-800-523-7749 | $\begin{array}{l\|} \hline \text { April } \\ 2020 \end{array}$ | Class II |
| *Morphine $2 \mathrm{mg} / \mathrm{mL}$ Cassette 00409113403 | Lack of Assurance of Sterility | If you have questions about this recall, Medical Center Pharmacy, Inc., 1-800-523-7749 | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |
| *Mitomycin 0.04\% Ophth DR eye drops 067457051805 | Lack of Assurance of Sterility | If you have questions about this recall, Medical Center Pharmacy, Inc., 1-800-523-7749 | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |
| Nystatin Oral Suspension 0021081016 | SubPotent Drug: Low out-ofspecification results for assay testing. | If you have questions about this recall, PAI Holdings, LLC. dba Pharmaceutical Associates Inc, 1-864-277-7282. | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |
| Theophylline (Anhydrous) Extended-Release 02903300101 | CGMP Deviations: poor manufacturing practices resulted in Labeling | If you have questions about this recall, Jubilant Draximage Inc, 1-800-361-2356. | $\begin{array}{l\|} \hline \text { April } \\ 2020 \end{array}$ | Class II |

[^29]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| DRAXIMAGE DTPA (KIT FOR THE PREPARATION OF TECHNETIUM TC 99M PENTETATE INJECTION) 06517428805 06717528830 | Failed Stability Specifications | If you have questions about this recall, Nostrum Laboratories Inc, 1-816-841-4636. | $\begin{array}{l\|} \hline \text { April } \\ 2020 \end{array}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Advil Liqui-Gel Mini 160+20+20 CT 00573171559 | Labeling: Lacks Warning or Rx Legend- Certain lots does not include required safety warning information in the Drug Facts Panel. | If you have questions about this recall, Glaxosmithkline Consumer Healthcare Holdings, +44 208047 5000. | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |
| Ibuprofen 50 mg per 1.25 mL Oral Suspension Advil Infant Concentrated Drops White Grape 00573019175 00573019150 | Labeling: Lacks Warning or Rx Legend- Certain lots does not include required safety warning information in the Drug Facts Panel. | If you have questions about this recall, Glaxosmithkline Consumer Healthcare Holdings, +44 208047 5000. | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |
| Ibuprofen 200 mg liquid filled capsules Advil Liqui-Gel Minis $\begin{aligned} & 00573176989 \\ & 00573176995 \end{aligned}$ | Labeling: Lacks Warning or Rx Legend- Certain lots does not include required safety warning information in the Drug Facts Panel. | If you have questions about this recall, Glaxosmithkline Consumer Healthcare Holdings, +44 208047 5000. | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |
| Advil Allergy \& Congestion Relief 00573019610 | Labeling: Lacks Warning or Rx Legend- Certain lots does not include required safety warning information in the Drug Facts Panel. | If you have questions about this recall, Glaxosmithkline Consumer Healthcare Holdings, +44 208047 | $\begin{array}{l\|} \hline \text { April } \\ 2020 \end{array}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

|  |  | 5000. |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Phenylephrine 10 mg tablets. 00573019620 | Labeling: Lacks Warning or Rx Legend- Certain lots does not include required safety warning information in the Drug Facts Panel. | If you have questions about this recall, Glaxosmithkline Consumer Healthcare Holdings, +44 208047 5000. | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |
| Ibuprofen 200 mg Chlorpheniramine Maleate 4 mg 00573019610 | Labeling: Lacks Warning or Rx Legend- Certain lots does not include required safety warning information in the Drug Facts Panel. | If you have questions about this recall, Glaxosmithkline Consumer Healthcare Holdings, +44 208047 5000. | $\begin{aligned} & \text { April } \\ & 2020 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { Doxycycline } \\ & 06330461501 \\ & 06330461650 \end{aligned}$ | CGMP Deviations: During manufacturing Solifenacin Succinate Tablets might convert to Solifenacin Tartrate Tablets. | If you have questions about this recall, Sun Pharmaceutical Industries, Inc., 1-800-818-4555. | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Solifenacin Succinate 05199189333 05199189390 | CGMP Deviations: During manufacturing Solifenacin Succinate Tablets might convert to Solifenacin Tartrate Tablets. | If you have questions about this recall, Breckenridge Pharmaceutical, Inc, 1-860-828-8140. | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Atorvastatin Calcium 06330482905 | Presence of foreign substance: Foreign matter has been identified as latex glove in one lot of Atorvastatin Calcium Tablets USP 40 mg . | If you have questions about this recall, Sun Pharmaceutical Industries, Inc., 1-800-818-4555. | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Pantoprazole Sodium DelayedRelease | CGMP Deviations: Presence of dark brown discoloration on edges of tablets | If you have questions about this recall, Jubilant Cadista Pharmaceuticals, Inc., 1-410-860-2836. | $\begin{aligned} & \text { March } \\ & 0000 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \hline \text { Sotalol HCL } \\ & 00378512301 \end{aligned}$ | Presence of particulate matter. presence of metal particles. | If you have questions about this recall, Mylan Pharmaceuticals Inc., 1-551-233-2700. | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Daytrana (methylphenidate transdermal system) <br> 06896855523 <br> 06896855533 <br> 06896855543 <br> 06896855553 | Defective Delivery System: Out of specification for mechanical peel and shear. | If you have questions about this recall, Noven Therapeutics, LLC, 1 -800-796-9526. | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Ranitidine 00113785282 05977954082 03780850782 02113011682 00363085282 | CGMP Deviation; Possible contamination with impurity N nitrosodimethylamine. | If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Ranitidine } \\ & 05977995001 \end{aligned}$ | CGMP Deviation; Possible contamination with impurity N nitrosodimethylamine. | If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Ranitidine <br> 00113785201 <br> 06839185256 <br> 05977954001 <br> 05591001101 | CGMP Deviation; Possible contamination with impurity N nitrosodimethylamine. | If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| 06398185256 03014260056 04125085201 01182208524 00363085201 |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \hline \text { Ranitidine } \\ & 04903580075 \\ & 01167395075 \end{aligned}$ | CGMP Deviation; Possible contamination with impurity N nitrosodimethylamine. | If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Ranitidine } \\ & 04903560875 \\ & 01167302375 \end{aligned}$ | CGMP Deviation; Possible contamination with impurity N nitrosodimethylamine. | If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260 | $\begin{array}{\|l\|} \hline \text { March } \\ 2020 \end{array}$ | Class II |
| Ranitidine 00113795009 05977995009 04903580009 04612253309 03014289109 03680095009 01167395009 00363095009 | CGMP Deviation; Possible contamination with impurity N nitrosodimethylamine. | If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Ranitidine 04152039209 05977954009 05531985209 04612253209 03780850709 04934810954 | CGMP Deviation; Possible contamination with impurity N nitrosodimethylamine. | If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260 | $\begin{array}{\|l\|} \hline \text { March } \\ 2020 \end{array}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class $\mathbf{2}$ Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

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| 04125089109 01182208523 03701285209 05059485209 04934810954 01167385209 00363085209 |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Ranitidine 05591085271 04612222471 00113085271 06925604171 03014260071 00904671651 04125085271 05606209971 01182208522 02113011671 03680085271 | CGMP Deviation; Possible contamination with impurity N nitrosodimethylamine. | If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Ranitidine } \\ & 01167395058 \end{aligned}$ | CGMP Deviation; Possible contamination with impurity N nitrosodimethylamine. | If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Ranitidine 04125095002 01182209500 03701295062 02113056862 | CGMP Deviation; Possible contamination with impurity N nitrosodimethylamine. | If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Blue Cross <br> Blue Shield <br> Blue Care Network of Michigan |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & 03680095062 \\ & 00363095002 \end{aligned}$ |  |  |  |  |
| Ranitidine 04699485262 04152039202 05977954002 05591085202 04116385262 04903560802 05530185202 05531985202 04612222462 00113085262 06925604162 04934810904 03014260002 00904671624 04125085202 05606209902 01182208525 01020285262 04119085262 03701285262 02113011602 05059485202 06201102821 03680085202 | CGMP Deviation; Possible contamination with impurity N nitrosodimethylamine. | If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
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Class $\mathbf{3}$ Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| $\begin{aligned} & 01167302302 \\ & 00363085262 \end{aligned}$ |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \text { Ranitidine } \\ & 00113085251 \\ & 00363085251 \end{aligned}$ | CGMP Deviation; Possible contamination with impurity N nitrosodimethylamine. | If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260 | $\begin{array}{\|l\|} \hline \text { March } \\ 2020 \end{array}$ | Class II |
| $\begin{aligned} & \hline \text { Ranitidine } \\ & 06984229306 \end{aligned}$ | CGMP Deviation; Possible contamination with impurity N nitrosodimethylamine. | If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260 | $\begin{array}{\|l\|} \hline \text { March } \\ 2020 \end{array}$ | Class II |
| $\begin{aligned} & \hline \text { Ranitidine } \\ & 03780887647 \\ & 04903587647 \end{aligned}$ | CGMP Deviation; Possible contamination with impurity N nitrosodimethylamine. | If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260 | $\begin{array}{\|l\|} \hline \text { March } \\ 2020 \end{array}$ | Class II |
| $\begin{aligned} & \text { Ranitidine } \\ & 03780887647 \\ & 04903587647 \end{aligned}$ | CGMP Deviation; Possible contamination with impurity N nitrosodimethylamine. | If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260 | $\begin{array}{\|l\|} \hline \text { March } \\ 2020 \end{array}$ | Class II |
| Ranitidine 00113787627 07000003752 01167387627 00363187627 | CGMP Deviation; Possible contamination with impurity Nnitrosodimethylamine. | If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260 | $\begin{array}{\|l\|} \hline \text { March } \\ 2020 \end{array}$ | Class II |

[^30]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Ranitidine 06201102832 09046715524 00125025272 04934813612 | CGMP Deviation; Possible contamination with impurity N nitrosodimethylamine. | If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260 | $\begin{aligned} & \hline \text { March } \\ & 2020 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Ranitidine 06984229365 04116393165 05531987665 00113087665 03780887665 06925687665 06201102831 06201102831 00904671546 02113011865 04934813644 01167387665 00363187665 | CGMP Deviation; Possible contamination with impurity N nitrosodimethylamine. | If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Lisinopril/HCTZ 68180051902 70518038203 | Presence of Foreign Tablets/Capsules | If you have questions about this recall, RemedyRepack Inc., 1-866-845-3791 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Calcium Chloride Injection 07128302253 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |

[^31]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Cholecalciferol (Vitamin D3) <br> Injection, VITAMIN D3 [P] 1,000 <br> IU/ML INJECTABLE <br> 07128302103 | Lack of Assurance of Sterility | If you have questions <br> about this recall, Fusion IV <br> Pharmaceuticals, Inc, 1- <br> $877-685-8222$ | March <br> 2020 | Class II |
| :--- | :--- | :--- | :--- | :--- |
| Zinc Chloride Injection, ZINC <br> CHLORIDE 10MG/ML <br> INJECTABLE <br> 07128302243 | Lack of Assurance of Sterility | If you have questions <br> about this recall, Fusion IV <br> Pharmaceuticals, Inc, 1- <br> $877-685-8222$ | March <br> 2020 | Class II |
| Cholecalciferol (Vitamin D3) <br> Injection, VITAMIN D3 [P] <br> 100,000 IU/ML INJECTABLE <br> 07128302513 | Lack of Assurance of Sterility | If you have questions <br> about this recall, Fusion IV <br> Pharmaceuticals, Inc, 1- <br> $877-685-8222 ~$ | March <br> 2020 | Class II |
| Elelyso (taliglucerase alfa) for <br> injection <br> 069010601 | Lack of Assurance of Sterility | If you have questions <br> about this recall, Pfizer <br> Labs, 1-877-225-9750 | March <br> 2020 | Class II |
| Mesalamine Delayed-Release <br> Tablets591224522 | Failed Dissolution Specifications: <br> Low out of specification dissolution <br> result observed during stability <br> testing. | If you have questions <br> about this recall, Teva <br> Pharmaceuticals USA, <br> Inc, 1-877-685-8222 | March <br> 2020 | Class II |
| Triamcinolone Diacetate <br> Injectable Suspension, | Lack of Assurance of Sterility | If you have questions <br> about this recall, Fusion IV <br> TRIAMCINOLONE DIACETATE <br> Pharmaceuticals, Inc, 1- <br> (PF) [2ML] CMC 40 MG/ML INJ <br> SUSP, For IM, IA7 | March <br> 0128306282 | 2020 |

[^32]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Triamcinolone Diacetate Injectable Suspension, TRIAMCINOLONE DIACETATE [10ML] CMC $80 \mathrm{MG} / \mathrm{ML}$ INJ SUSP, For IM, IA 07128306341 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \hline \text { March } \\ & 2020 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Triamcinolone Diacetate Injectable Suspension, TRIAMCINOLONE Diacetate [10ML] CMC 10MG/ML INJ SUSP, For IM, IA 7128306251 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Triamcinolone Acetonide (PF) Injectable Suspension, <br> TRIAMCINOLONE ACETONIDE (PF) [2ML] 40MG/ML INJ SUSP, For IM, IA7128306352 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Triamcinolone Acetonide (PF) Injectable Suspension, <br> TRIAMCINOLONE ACETONIDE <br> (PF) [10ML] 50MG/ML INJ <br> SUSP, For IM, IA7128306331 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Triamcinolone <br> Acetonide/Bupivacaine Hydrochloride Injectable Suspension, TRIAMCINOLONE ACET/BUPIVACAINE HCL | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |

## * Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
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| [10ML] 40MG/5MG/ML INJ SUSP, For IM, IA 07128306321 |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Testosterone Cypionate Injection, TESTOSTERONE CYP IN GRAPESEED OIL [1ML] 200 MG/ML INJECTABLE 07128305302 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Triamcinolone Acetonide (PF) Injectable Suspension, TRIAMCINOLONE ACETONIDE (PF) [10ML] 50MG/ML INJ SUSP, For IM, IA 07128306331 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Triamcinolone Acetonide (PF) Injectable Suspension, <br> TRIAMCINOLONE ACETONIDE (PF) [2ML] 40MG/ML INJ SUSP, For IM, IA 07128306352 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Triamcinolone Diacetate Injectable Suspension, TRIAMCINOLONE Diacetate [10ML] CMC 10MG/ML INJ SUSP, For IM, IA | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class $\mathbf{2}$ Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| 071283062511 |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Testosterone Cypionate/Progesterone Injection, TESTOSTERONE CYP/PROGESTERONE [2ML] 200MG/2.5MG/ML INJECTABLE7128305312 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Testosterone Cypionate Injection,TESTOSTERONE CYP IN GRAPESEED OIL [10ML] 200 MG/ML INJECTABLE 07128305301 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Selenium Injection, SELENIUM 200MCG/ML INJECTABLE 07128302273 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Super MIC Injection, SUPER MIC* INJECTABLE <br> 07128302343 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Nicotinamide Adenine Dinucleotide (PF) Injection, NICOTINAMIDE ADENINE DINUCLEOTIDE (PF) 20 MG/ML INJECTABLE | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class $\mathbf{2}$ Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| 07128303361 |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Iohexol (PF) <br> Injection,OMNIPAQUE <br> INJECTION [5ML] 300MG I/ML <br> INJECTABLE <br> 07128314135 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Pyridoxine Hydrochloride Injection, PYRIDOXINE HCL 100 MG/ML INJECTABLE $07128302163$ | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Nicotinamide Adenine Dinucleotide (PF) Injection, NICOTINAMIDE ADENINE DINUCLEOTIDE (PF) $50 \mathrm{MG} / \mathrm{ML}$ INJECTABLE 07128303061 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| MIC-PLUS Injection, Vitamin Complex, MIC-COMBO* 25MG/50MG/50MG/1MG/20MG/ 5MG/ML INJECTABLE 07128302313 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |

[^33]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Mitomycin-C (PF) Irrigation Solution, MITOMYCIN-C (PF) 0.5 MG/ML PF SYRINGE 07128308128 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | March 2020 | Class II |
| :---: | :---: | :---: | :---: | :---: |
| MIC-PLEX Injection, Vitamin Complex, MIC-COMBO* 25MG/50MG/50MG/1MG/20MG/ 5MG/ML <br> INJECTABLE7128302333 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | March 2020 | Class II |
| MIC-B12 Injection, MIC-B12 25MG/50MG/50MG/1MG/ML INJECTABLE7128302303 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | March 2020 | Class II |
| Methylprednisolone Acetate Injectable Suspension, METHYLPREDNISOLONE ACETATE [10ML] CMC 50 MG/ML INJ SUSP, For IM, IA 07128306361 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | March 2020 | Class II |
| Methylprednisolone Acetate Injectable Suspension, METHYLPREDNISOLONE ACETATE [10ML] CMC 100 MG/ML INJ SUSP 07128306191 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |

[^34]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Methylprednisolone Acetate/Bupivacaine Hydrochloride Injectable Suspension 07128306241 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Methylprednisolone Acetate Injectable Suspension, METHYLPREDNISOLONE ACETATE (PF) CMC [2ML] 80 MG/ML INJ SUSP, For IM, IA 07128306392 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Methylprednisolone Acetate/Bupivacaine Hydrochloride Injectable Suspension, METHYLPRED ACETATE/BUPIV [10ML] CMC 40MG/5MG/ML INJ SUSP, For IM, IA 07128306231 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Methylcobalamin Injection, METHYLCOBALAMIN [CD] 10MG/100MG/ML INJECTABLE, For IV, IM 07128303383 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |

[^35]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

Blue Cross Blue Shield
Blue Care Network
of Michigan

| Methylcobalamin Injection, METHYLCOBALAMIN 1 MG/ML INJECTABLE 07128303313 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Lysine Hydrochloride Injection, <br> LYSINE HCL 100MG/ML <br> INJECTABLE <br> 07128303423 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Levocarnitine Injection, LEVOCARNITINE $500 \mathrm{MG} / \mathrm{ML}$ INJECTABLE 07128392603 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Lidocaine HCL (PF) Injection, LIDOCAINE HCL 4\% (PF) 40 MG/ML INJECTABLE 07128310115 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Polyoxyl Lauryl Ether (Polidocanol) Injection, <br> LAURETH-9 (POLIDOCANOL) <br> 5\% INJECTABLE <br> 0712830603 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Hydroxyprogesterone Caproate Injection, <br> HYDROXYPROGESTERONE CAPROATE [4ML] $250 \mathrm{MG} / \mathrm{ML}$ INJECTABLE | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| 07128305325 |  |  |  |  |
| :--- | :--- | :--- | :--- | :--- |
| Arginine Hydrochloride Injection, <br> L-ARGININE HCL 100MG/ML <br> INJECTABLE <br> 07128303413 | Lack of Assurance of Sterility | If you have questions <br> about this recall, Fusion IV <br> Pharmaceuticals, Inc, 1- <br> $877-685-8222$ | March <br> 2020 | Class II |
| Human Chorionic Gonadotropin <br> (hCG) Injection, HCG [10ML] <br> 1000 IU/ML INJECTABLE <br> 07128305331 |  | Lack of Assurance of Sterility | If you have questions <br> about this recall, Fusion IV <br> Pharmaceuticals, Inc, 1- <br> $877-685-8222$ | March <br> 2020 |
| Glycine Injection, GLYCINE USP <br> 50MG/ML INJECTABLE <br> 07128303443 | Lack of Assurance of Sterility | If you have questions <br> about this recall, Fusion IV <br> Pharmaceuticals, Inc, 1- <br> $877-685-8222$ | March <br> 2020 | Class II |
| Glycerin (PF) Injection, <br> GLYCERIN 99\% INJECTABLE <br> 07128304641 | Lack of Assurance of Sterility | If you have questions <br> about this recall, Fusion IV <br> Pharmaceuticals, Inc, 1- <br> $877-685-8222$ | March <br> 2020 | Class II |
| Glutathione <br> Injection,GLUTATHIONE <br> 200MG/ML INJECTABLE <br> 07128312313 |  | If you have questions <br> about this recall, Fusion IV <br> Pharmaceuticals, Inc, 1- <br> $877-685-8222 ~$ | March <br> 2020 | Class II |

[^36]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Hydroxocobalamin Injection, <br> HYDROXOCOBALAMIN <br> 5MG/ML INJECTABLE | Lack of Assurance of Sterility | If you have questions <br> about this recall, Fusion IV <br> Pharmaceuticals, Inc, 1- <br> 07128303393 |  | March <br> 2020 |
| :--- | :--- | :--- | :--- | :--- |
| Glutathione <br> Injection,GLUTATHIONE <br> 200MG/ML INJECTABLE <br> 07128312313 | Lack of Assurance of Sterility | If you have questions <br> about this recall, Fusion IV <br> Pharmaceuticals, Inc, 1- <br> $877-685-8222$ | March <br> 2020 | Class II |
| Dexpanthenol Injection, <br> DEXPANTHENOL 250 MG/ML <br> INJECTABLE <br> 07128302143 |  | If you have questions <br> about this recall, Fusion IV <br> Pharmaceuticals, Inc, 1- <br> $877-685-8222 ~$ | March <br> 2020 | Class II |
| Dexamethasone (LA) Injectable <br> Suspension, DEXAMETHASONE | Lack of Assurance of Sterility | If you have questions <br> about this recall, Fusion IV <br> LA [10ML] 16MG INJ SUSP |  | Pharmaceuticals, Inc, 1- <br> 07128306301 |

* Drug Recall Class

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Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

|  |  | 877-685-8222 |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Cyanocobalamin/Folinic Acid Injection, CYANOCOBALAMIN : FOLINIC ACID 2000 MCG/ML: 500MCG/ML INJECTABLE 07128303333 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Injection,COENZYME Q-10 20MG/ML OIL INJ SOLN 07128304111 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| Biotin (Vitamin H) Injectable Suspension, BIOTIN 10 MG/ML INJ SUSP, For IM 07128302203 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |
| BETAMETHASONE <br> ACETATE/BETAMETHASONE <br> (PF) Injectable Suspension, <br> Betamethasone <br> Acetate/Betamethasone (PF) <br> CMC [5ML] 7MG/ML INJ SUSP, <br> For IM, IA <br> 07128306215 | Lack of Assurance of Sterility | If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222 | $\begin{aligned} & \text { March } \\ & 2020 \end{aligned}$ | Class II |

[^37]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| BETAMETHASONE | Lack of Assurance of Sterility | If you have questions <br> about this recall, Fusion IV <br> Pharmaceuticals, Inc, 1- <br> (PF) Injectable Suspension, <br> Betamethasone <br> Acetate/Betamethasone (PF) <br> CMC [10ML] 7MG/ML INJ SUSP, <br> For IM, IA <br> 07128306211 |  | $877-685-8222$ |
| :--- | :--- | :--- | :--- | :--- |

## * Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

|  |  |  |  |  |
| :--- | :--- | :--- | :--- | :--- |
| Alprostadi//Papaverine <br> Hydrochloride/Phentolamine <br> Mesylate Injection <br> 07128305462 | Lack of Assurance of Sterility | If you have questions <br> about this recall, Fusion IV <br> Pharmaceuticals, Inc, 1- <br> $877-685-8222$ | March <br> 2020 | Class II |
| Alprostadil/Papaverine <br> Hydrochloride/Phentolamine <br> Mesylate Injection <br> 07128305452 | Lack of Assurance of Sterility | If you have questions <br> about this recall, Fusion IV <br> Pharmaceuticals, Inc, 1- <br> $877-685-8222$ | March <br> 2020 | Class II |
| Alprostadil (prostaglandin E1) 80 <br> MCG/ML Injectable <br> 07128305432 | Lack of Assurance of Sterility | If you have questions <br> about this recall, Fusion IV <br> Pharmaceuticals, Inc, 1- <br> $877-685-8222$ | March <br> 2020 | Class II |
| Alprostadil (prostaglandin E1)150 <br> MCG/ML Injectable <br> 027128305382 | Patent ductus arteriosus | If you have questions <br> about this recall, Fusion IV <br> Pharmaceuticals, Inc, 1- <br> $877-685-8222$ | March <br> 2020 | Class II |
| Alprostadil/Papaverine <br> Hydrochloride/Phentolamine <br> Mesylate/Atropine Sulfate <br> Injection <br> 07128305502 |  | If you have questions <br> about this recall, Fusion IV <br> Pharmaceuticals, Inc, 1- <br> $877-685-8222 ~$ | March <br> 2020 | Class II |

## * Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Alprostadil/Papaverine <br> Hydrochloride/Phentolamine <br> Mesylate/Atropine Sulfate <br> Injection <br> 07128305492 | Lack of Assurance of Sterility | If you have questions <br> about this recall, Fusion IV <br> Pharmaceuticals, Inc, 1- <br> $877-685-8222$ | March <br> 2020 | Class II |
| :--- | :--- | :--- | :--- | :--- |
| Alprostadil/Papaverine <br> Hydrochloride/Phentolamine <br> Mesylate/Atropine Sulfate <br> Injection 07128305472 | Lack of Assurance of Sterility | If you have questions <br> about this recall, Fusion IV <br> Pharmaceuticals, Inc, 1- <br> $877-685-8222 ~$ | March <br> 2020 | Class II |
| Phenytoin Oral Suspension <br> 05167240691 | Resuspension Problems | If you have questions <br> about this recall, Taro <br> Pharmaceuticals U.S.A, 1- <br> $866-923-4914$. | March <br> 2020 | Class II |
| Glycopyrrolate Tabs <br> 00615817039 | If you have questions <br> about this recall, NCS <br> Healthcare of Kentucky <br> Inc, 1-270-651-6188 | February <br> 2020 | Class II |  |
| Desmopressin Acetate Tablets <br> 06808460421 | GMP Deviations | If you have questions <br> about this recall, American <br> Health Packaging, 1-800- <br> $967-5952$. | February <br> 2020 | Class II |
| Desmopressin Acetate Tablets <br> 06808460621 | GMP Deviations | If you have questions <br> about this recall, American <br> Health Packaging, 1-800- <br> $967-5952$. | February <br> 2020 | Class II |

[^38]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| $\begin{aligned} & \text { Lamotrigine } \\ & 05167241311 \end{aligned}$ | Cross contamination | If you have questions about this recall, Taro Pharmaceuticals U.S.A, 1 -866-923-4914. | $\begin{aligned} & \text { February } \\ & 2020 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Methylphenidate hydrochloride Extended-Release 06203772501 | Product bottle may be absent of desiccant. | If you have questions about this recall, Teva Pharmaceuticals, 1-888-838-2872. | $\begin{aligned} & \text { February } \\ & 2020 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Fentanyl Citrate Inj } \\ & 0409909412 \end{aligned}$ | Defective container | If you have questions about this recall, Pfizer Inc, 1-800-879-3477. | $\begin{aligned} & \text { February } \\ & 2020 \end{aligned}$ | Class II |
| Ethacrynate Sodium for Injection 06838224601 | cGMP Deviations | If you have questions about this recall, Zydus Pharmaceuticals, 1-877-993-8779. | $\begin{aligned} & \text { February } \\ & 2020 \end{aligned}$ | Class II |
| Caduet (amlodipine besylate/atorvastatin calcium) 00069218030 | Defective container | If you have questions about this recall, Pfizer Inc, 1-800-879-3477. | $\begin{aligned} & \text { February } \\ & 2020 \end{aligned}$ | Class II |
| Hydrocortisone and Acetic Acid Otic <br> 05038390110 | Subpotent Drug | If you have questions about this recall, TECH PHARMACAL CO., INC, 1-800-932-5676. | $\begin{aligned} & \text { February } \\ & 2020 \end{aligned}$ | Class II |
| Ranitidine Hydrochloride 04306384414, <br> 04306384430 , <br> 04306384490, <br> 04306384401 | cGMP Deviations | If you have questions about this recall, PD-Rx Pharmaceuticals, Inc., 1-405-942-3040. | $\begin{aligned} & \text { February } \\ & 2020 \end{aligned}$ | Class II |

* Drug Recall Class

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Class $\mathbf{2}$ Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Olmesartan Medoxomil Tablets <br> 06787744690 | cGMP Deviations | If you have questions <br> about this recall, Ascend <br> Laboratories LLC, 1-201- <br> 476-1977. | February <br> 2020 | Class II |
| :--- | :--- | :--- | :--- | :--- |
| Atorvastatin Calcium Tablets <br> 07037702711 | Presence of Foreign Tablets | If you have questions <br> about this recall, Graviti <br> Pharmaceuticals Private <br> Limited, 040 6815 5555. | February <br> 2020 | Class II |
| Walgreens Acne Cleansing Bar, <br> Benzoyl Peroxide 10\% <br> 00363013711 | Presence of Foreign Substance | If you have questions <br> about this recall, Shandex <br> Personal Care <br> Manufacturing Inc., 1-613- <br> 267-1881 | February |  |

* Drug Recall Class

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Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

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Blue Cross
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| 06878863820 |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Nizatidine Capsules 00378515091 , <br> 00378530093 | Trace amounts of an impurity | If you have questions about this recall, Mylan Pharmaceuticals Inc, 1-304-599-2595. | $\begin{aligned} & \hline \text { January } \\ & 2020 \end{aligned}$ | Class II |
| Ranitidine Tablets <br> 52959050207 <br> 52959050214 <br> 52959050220 <br> 52959050230 <br> 52959050260 | Trace amounts of an impurity | If you have questions about this recall, H.J. Harkins Company, Inc, 1-800-841-5554. | $\begin{aligned} & \hline \text { January } \\ & 2020 \end{aligned}$ | Class II |
| Ranitidine Tablets 07093401704, 07093401720, 07093401724, 07093401730, 07093401790, 07093428715, 07093428790 | Trace amounts of an impurity | If you have questions about this recall, Denton Pharma, Inc., 1-800-7220772. | $\begin{aligned} & \hline \text { January } \\ & 2020 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { Netspot } \\ & 06948800140 \end{aligned}$ | Defective container | If you have questions about this recall, Advanced Accelerator Applications USA, Inc., 1-862-263-0820. | $\begin{aligned} & \hline \text { January } \\ & 2020 \end{aligned}$ | Class II |

* Drug Recall Class

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| Estriol USP Micronized 62991215906, 62991215903, 62991215905, 62991215902 | Subpotent | If you have questions about this recall, Letco Medical LLC, 1-800-2395288. | $\begin{array}{\|l} \hline \text { January } \\ 2020 \end{array}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Dutasteride 5026828213 | cGMP Deviations | If you have questions about this recall, AvKARE, Inc, 1-931-292-6222 | $\begin{aligned} & \text { January } \\ & 2020 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Testosterone Cypionate } \\ & 06275601740,06275601540, \\ & 06275601640 \end{aligned}$ | cGMP Deviations | If you have questions about this recall, Sun Pharmaceutical Industries, Inc., 1-609-495-2800. | $\begin{aligned} & \text { January } \\ & 2020 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Sumatriptan Succinate } \\ & 06275652169,06275652188 \end{aligned}$ | Failed Impurities/Degradation Specifications; out-of-specification results obtained for related substance. | If you have questions about this recall, Sun Pharmaceutical Industries, Inc., 1-609-495-2800. | $\begin{aligned} & \text { January } \\ & 2020 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Ranitidine } \\ & 06255969060,06255969005 \end{aligned}$ | Impurities | If you have questions about this recall, Appco Pharma LLC, 1-732-2537735. | $\begin{aligned} & \text { January } \\ & 2020 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Ranitidine } \\ & 03780819601,03780819604 \end{aligned}$ | Impurities | If you have questions about this recall, AAA Pharmaceutical, Inc, 1-609-288-6060. | $\begin{aligned} & \text { January } \\ & 2020 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { Blisovi Fe 1.5/30 } \\ & 06818086611 \end{aligned}$ | Failed tablet | If you have questions about this recall, Lupin Pharmaceuticals Inc, 1-410-576-2000. | $\begin{aligned} & \text { January } \\ & 2020 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Estriol 07642004936,07642004937, 07642004938 | Presence of impurities | If you have questions about this recall, Asclemed USA Inc, 1-310-320-0100. | $\begin{aligned} & \text { January } \\ & 2020 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Ranitidine 06846224860,68486224860, 06846224801, 68486224860, 06846224930, 68486224860 06846224901, 68486224860 06846224920, 68486224860, 06846224805,68486224860 | Presence of impurities | If you have questions about this recall, Glenmark Pharmaceuticals Inc, 1-888-721-7115 | $\begin{aligned} & \text { January } \\ & 2020 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Ranitidine } \\ & 06220777332 \end{aligned}$ | Presence of impurities | If you have questions about this recall, Granules India Limited, 1-877-7703183. | $\begin{aligned} & \text { January } \\ & 2020 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { Myorisan® } \\ & 06174830213 \end{aligned}$ | Unit dose mispackaging | If you have questions about this recall, Akorn, Inc, 1-800-932-5676. | $\begin{aligned} & \text { January } \\ & 2020 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Ranitidine } \\ & 06068726069 \end{aligned}$ | Presence of impurities | If you have questions about this recall, American Health Packaging, 1-800-707-4621. | $\begin{aligned} & \text { January } \\ & 2020 \end{aligned}$ | Class II |
| Amantadine Hydrochloride 5974669901 | Presence of foreign substance | If you have questions about this recall, Jubilant Cadista Pharmaceuticals, Inc. 1-410-860-8500 | $\begin{aligned} & \text { December } \\ & 2019 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| $\begin{aligned} & \hline \text { 25\% Dextrose } ® \\ & 00409177510 \end{aligned}$ | Labeling error | If you have questions about this recall, Pfizer Inc1-877-225-9750 | $\begin{aligned} & \text { December } \\ & 2019 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Vancomycin Hydrochloride 05515020420 | Product discoloration | If you have questions about this recall, AuroMedics Pharma LLC, 888-238-7880. | $\begin{aligned} & \text { December } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Lidocaine Hcl } \\ & 70004072309 \end{aligned}$ | Presence of foreign substance | If you have questions about this recall, SCA Pharmaceuticals, LLC, 1-877-550-5059. | $\begin{aligned} & \text { December } \\ & 2019 \end{aligned}$ | Class II |
| Zantac®  <br> 06671597362,  <br> 06671597363,60671597368,  <br> 00597012201 00597012208, <br> 00597012213 00597012237, <br> 00597012240 00597012254, <br> 00597012261 00597012281, <br> 00597012296  | Presence of impurities | If you have questions about this recall, SanofiAventis U.S. LLC, 1-800-633-1610 | $\begin{aligned} & \text { December } \\ & 2019 \end{aligned}$ | Class II |
| Regular Strength Zantac $®$ 05026922225,06775115101, 06775115201,06775115202 , 06815125840 | Presence of impurities | If you have questions about this recall, SanofiAventis U.S. LLC, 1-800-633-1610 | $\begin{aligned} & \text { December } \\ & 2019 \end{aligned}$ | Class II |

## * Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

|  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Maximum Strength Zantac ${ }^{\circledR}$ 00597012101, 00597012106, 00597012108, 00597012109, 00597012111, 00597012124, 00597012138, 00597012150, 00597012164, 00597012166, 00597012168, 00059701218, 00597012180, 00597012182, 00597012185, 00597012190, 0059701294 | Presence of impurities | If you have questions about this recall, SanofiAventis U.S. LLC, 1-800-633-1610 | $\begin{aligned} & \text { December } \\ & 2019 \end{aligned}$ | Class II |
| Regular Strength Zantac ${ }^{\circledR}$ 04116703000, 04116703001, 04116703003, 04116703005, 04116703006, 04116703007, 04116703008,00526922025 | Presence of impurities | If you have questions about this recall, SanofiAventis U.S. LLC, 1-800-633-1610 | $\begin{aligned} & \text { December } \\ & 2019 \end{aligned}$ | Class II |
| Cool Mint Tablets Maximum Strength Zantac ${ }^{\circledR}$ 00597012006,00597012008, 00597012009,00597012024, 00597012038,00597012050, 00597012076, 00597012078 00597012080,00597012082, 00597012087, 04116703201, 04116703202,04116703203 04116703204,04116703205 04116703206, 04116703207 | Presence of impurities | If you have questions about this recall, SanofiAventis U.S. LLC, 1- 800-633-1610 | $\begin{aligned} & \text { December } \\ & 2019 \end{aligned}$ | Class II |

[^39]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Ibuprofen 068788726801 | Presence of foreign substance | If you have questions about this recall, Preferred Pharmaceuticals Inc 1-714-777-3729 | $\begin{aligned} & \text { December } \\ & 2019 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Lidocaine hcl 05515016505 | Presence of foreign substance | If you have questions about this recall, AuroMedics Pharma, 1-888-238-7880 | $\begin{aligned} & \text { December } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Ranitidine } \\ & 04229172460,04229172530, \end{aligned}$ | Presence of impurities | If you have questions about this recall, AvKARE, Inc, 1-931-292-6222 | $\begin{aligned} & \text { December } \\ & 2019 \end{aligned}$ | Class II |
| Ranitidine <br> 06516225306, 06516225310, <br> 06516225318,06516225350, <br> 06516225311,06516225430, <br> 06516225410,06516225425, <br> 06516266490,05374625310, <br> 05374625402 <br> Ral | Presence of impurities | If you have questions about this recall, Amneal Pharmaceuticals, Inc., 1-908-947-3120 | $\begin{aligned} & \text { December } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Ranitidine } \\ & 068788707803,068788707806 \end{aligned}$ | Presence of impurities | If you have questions about this recall, Preferred Pharmaceuticals Inc 1-714-777-3729 | $\begin{aligned} & \text { December } \\ & 2019 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class $\mathbf{2}$ Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| $\begin{aligned} & \hline \text { Ranitidine } \\ & 05591009279 \end{aligned}$ | Presence of impurities | If you have questions about this recall, Dolgencorp LLC, Inc, 1-615-855-4000 | $\begin{aligned} & \text { December } \\ & 2019 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Ranitidine 05965114460, 059651145-30, 06586243174, 05965114405, | Presence of impurities | If you have questions about this recall, Aurobindo Pharma, 1-866-850-2876 | $\begin{aligned} & \text { December } \\ & 2019 \end{aligned}$ | Class II |
| Walgreens Sodium Chloride Ophthalmic Solution 00363019313 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November $2019$ | Class II |
| Walgreens Sodium Chloride Ophthalmic Ointment 00363750050 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November 2019 | Class II |
| Walgreen's Lubricant Eye Ointment, Mineral Oil 00363019150 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November $2019$ | Class II |
| Equate Support Harmony Lubricant Eye Drops 04903514510 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1- | November 2019 | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

|  |  | 631-722-5988 Ext 16 |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Equate Support Advanced 04903588549 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November $2019$ | Class II |
| Equate Support Advanced Lubricating Eye Drops 04903588254 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November 2019 | Class II |
| Equate Support Advanced Lubricant Gel Drops 0903588252 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Equate Sterile Lubricant Stye Ointment $04903587550$ | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Equate Restore Tears Lubricant Eye Drops Twin Pack 04903518949 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Equate Restore PM Nighttime Lubricant Eye Ointment 04903519150 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Equate Night \& Day Restore Tears Lubricant Eye Pack 04903588359 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Equate Eye Allergy Relief Drops 4903587413 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Equate Eye Allergy Relief Drops 04903588713 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November 2019 | Class II |
| Equate Comfort Gel Lubricant Eye Gel 04903519749 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November $2019$ | Class II |
| Puralube Petrolatum Ophthalmic Ointment ${ }^{\text {B }}$ $00574402535$ | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1- | November $2019$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

|  |  | 631-722-5988 Ext 16 |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Polycin (bacitracin zinc and polymyxin B sulfate) Ophthalmic Ointment ${ }^{8}$ 00574402135 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November 2019 | Class II |
| Tetcaine (Tetracaine Hydrochloride) Ophthalmic Solution (8) 05479950215 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November $2019$ | Class II |
| CVS Health Preservative Free Lubricant Eye Drops® $50428302958$ | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November $2019$ | Class II |
| TRP Stye Relief® 01731201413 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| TRP Pink Eye Relief® 01731201315 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| TRP Blur Relief 01731200211 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Perrigo Sulfacetamide Sodium Ophthalmic Ointment® 00574419035 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Perrigo Sterile Neo-Polycin HC (neomycin and polymixin B sulfates, bacitracin zinc and hydrocortisone acetate) Ophthalmic ${ }^{\text {B }}$ 00574414435 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Perrigo Neo-Polycin neomycin and polymixin B sulfates and bacitracin zinc Ophthalmic Ointment ${ }^{8}$ 0574425035 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Perrigo Neomycin and Polymixin B Sulfates and Dexamethasone Ophthalmic® 0574416035 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |

[^40]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Perrigo Bacitracin Ophthalmic Ointment ${ }^{\text {B }}$ 0574402235 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Ofloxacin Ophthalmic Solution 05939014005 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \hline \text { November } \\ & 2019 \end{aligned}$ | Class II |
| OCuSOFT Tetravisc Tetracaine $\mathrm{HCl} 0.5 \%$ Sterile Anesthetic ${ }^{\circledR}$ 05479950505 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November $2019$ | Class II |
| OCuSOFT Tetravisc Tetracaine $\mathrm{HCl} 0.5 \%$ Sterile Anesthetic $05479950501 ®$ | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November $2019$ | Class II |
| OCuSOFT Tetravisc Forte Tetracaine HCl 0.5 \% Sterile Anesthetic ${ }^{\circledR}$ 05479950405 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November $2019$ | Class II |
| OCuSOFT Tetravisc Forte (Tetracaine HCl) 0.5\% Sterile Anesthetic ${ }^{\circledR}$ | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1 - | November $2019$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| 05479950401 |  | 631-722-5988 Ext 16 |  |  |
| :---: | :---: | :---: | :---: | :---: |
| OCuSOFT Tears Again Sterile Lubricant Eye Drops® 05479990430 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November 2019 | Class II |
| Ocusoft Tears Again Lubricant Eye Drops ${ }^{\circledR}$ 05479990415 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November 2019 | Class II |
| OCuSOFT Homatropine Hydrobromide Ophthalmic Solution (8) $05479943105$ | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November 2019 | Class II |
| Ocusoft Goniosoft Hypromellose 2.5\% Opthalmic Demulcent Solution (8) 05479950315 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November 2019 | Class II |
| OCuSOFT Flucaine <br> Proparacaine Hydrochloride and Fluorescein Sodium Ophthalmic Solution (8) $05479950721$ | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November $2019$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| OCuSOFT Eye Wash Sterile Isotonic Buffered Solution ${ }^{\circledR}$ 05479956501 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| OCuSOFT Eye Wash Sterile Isotonic 05479956559 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Natural Ophthalmics Women's Tear Stimulation Dry Eye Drops® 06877010315 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Natural Ophthalmics Tear Stimulation Forte Dry Eye Drops® $06877010415$ | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Natural Ophthalmics Ortho-K Thin Eye Drops ${ }^{\circledR}$ 06877014415 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Natural Ophthalmics Ortho-K Thick Comfort Gel 06877014315 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1- | November $2019$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

|  |  | 631-722-5988 Ext 16 |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Natural Ophthalmics Cataract Eye Drops® 06877013015 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November $2019$ | Class II |
| Natural Ophthalmics Allergy Desensitization Eye Drops® 06877012015 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \hline \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Nano Tears XP Clear Emollient Lubricant Gel Drops® $05939014351$ | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Nano Tears TF Clear Emollient Lubricant Gel Drops® $05939014249$ | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Nano Tears TF Clear Emollient Lubricant Gel Drops® $05939014156$ | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Lubricant Eye Drops Moisturizing <br> Twin Pack® <br> 00363018549 | Lack of Assurance of Sterility | If you have questions <br> about this recall, Altaire <br> Pharmaceuticals, Inc, 1- <br> $631-722-5988$ Ext 16 | November <br> 2019 | Class II |
| :--- | :--- | :--- | :--- | :--- |
| iSolutions NanoTears HA <br> Preservative Free Multi - Dose <br> Lubricant Gel Drops® <br> 05939020810 |  | Lack of Assurance of Sterility | If you have questions <br> about this recall, Altaire <br> Pharmaceuticals, Inc, 1- | November <br> 631-722-5988 Ext 16 |
| iSolutions Nano Tears XP Clear <br> Emollient Lubricant Gel Drops® |  | Class II |  |  |
| 05939014310 |  |  |  |  |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| 05939014756 |  | 631-722-5988 Ext 16 |  |  |
| :---: | :---: | :---: | :---: | :---: |
| iSolutions Nano Tears MXP Forte Clear Emollient Lubricant Gel Drops® 05939014410 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| iSolutions Nano Tears MO Clear Emollient Lubricant Drops ${ }^{\circledR}$ 05939014510 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \hline \text { November } \\ & 2019 \end{aligned}$ | Class II |
| iSolutions ActivEyes Nighttime Lubricant Eye Ointment® 05939019050 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \hline \text { November } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { FreshKote Lubricant Eye Drops® } \\ & 01582110115 \end{aligned}$ | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Altaire Sterile Eye Wash® } \\ & 05939017513,05939017535 \text {, } \\ & 05939017518 \end{aligned}$ | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Altaire Homatropaire Homatropine Hyrdobromide Opthalmic Solution ${ }^{\text {B }}$ 05939019205 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November $2019$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Altaire Goniotaire Hypromellose 2.5\% Opthlamic Demulcent Solution ${ }^{\text {® }}$ 05939018213 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Altaire Fluorescein Sodium with Proparacaine Hydrochloride Ophthalmic Solution 05939020505® | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Altaire Diclofenac Sodium Opthalmic Solution ® 5939014905 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November $2019$ | Class II |
| Altaire Diclofenac Sodium Opthalmic Solution ® 05939014902 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November $2019$ | Class II |
| Altaire Ciprofloxacin Ophthalmic Solution ${ }^{\circledR}$ $05939021710$ | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, | November $2019$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

|  |  | 631-722-5988 Ext 16 |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Altaire Ciprofloxacin Ophthalmic Solution ${ }^{\circledR}$ $05939021702$ | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November $2019$ | Class II |
| Altaire Ciprofloxacin HCl Ophthalmic Solution ${ }^{\circledR}$ 05939021705 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November $2019$ | Class II |
| Altacaine (Tetracaine Hydrochloride) Ophthalmic Solution ( ${ }^{8}$ 05939018113 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November 2019 | Class II |
| ActivEyes Sterile Altalube Ointment ${ }^{\text {® }}$ $05939019850$ | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November $2019$ | Class II |
| ActivEyes Preservative Free Multi-Dose Lubricant Gel Drops® 05939014852 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November $2019$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| ActivEyes Preservative Free Multi-Dose Lubricant Drops® 05939014652 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| ActivEyes Sterile Altalube Ointment ${ }^{(8)}$ $05939019850$ | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November 2019 | Class II |
| ActivEyes Preservative Free Multi-Dose Lubricant Gel Drops® ${ }^{\circledR}$ 05939014852 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November 2019 | Class II |
| ActivEyes Preservative Free Multi-Dose Lubricant Drops ${ }^{\circledR}$ 05939014652 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November 2019 | Class II |
| $\begin{aligned} & \text { ActivEyes Lubricant Eye } \\ & \text { Ointment® } \\ & 05939018950 \end{aligned}$ | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November $2019$ | Class II |
| ActivEyes Altachlore Solution 05939018313® | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1- | November $2019$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

|  |  | 631-722-5988 Ext 16 |  |  |
| :---: | :---: | :---: | :---: | :---: |
| ActivEyes Altachlore Sodium Chloride Hypertonicity Opthalmic Ointment ${ }^{8}$ 05939018450 | Lack of Assurance of Sterility | If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16 | November $2019$ | Class II |
| $\begin{aligned} & \hline \text { Ranitidine } \\ & 05140709705 \end{aligned}$ | impurity detected | If you have questions about this recall, Golden State Medical Supply Inc. 1-805-477-9866 | November $2019$ | Class II |
| Novitium Pharma Ranitidine®® <br> 07095400120 <br> 07095400140 <br> 07095400210 <br> 07095400240 | impurity detected | If you have questions about this recall, Novitium Pharma LLC. 1-855-2041431 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Rite Aid Pharmacy Maximum Strength Ranitidine ${ }^{\circledR}$ $\begin{aligned} & 01182260522 \\ & 01182260518 \end{aligned}$ | Possible contamination | If you have questions about this recall, Apotex Inc. 1-800-706-5575 | November $2019$ | Class II |
| Equate Maximum Strength <br> Ranitidine® <br> 04903511706 | Possible contamination | If you have questions about this recall, Apotex Inc. 1-800-706-5575 | November $2019$ | Class II |

* Drug Recall Class

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Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Rite Aid Pharmacy Maximum Strength Ranitidine® 01182261074 | Possible contamination | If you have questions about this recall, Apotex Inc. 1-800-706-5575 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Equate Maximum Strength Ranitidine® $04903510007$ $04903510000$ | Possible contamination | If you have questions about this recall, Apotex Inc. 1-800-706-5575 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Walgreens Regular Strength Wal-Zan 75 Ranitidine® 00363102903 | Possible contamination | If you have questions about this recall, Apotex Inc. 1-800-706-5575 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Walgreens Maximum Strength Wal-Zan 150 Ranitidine ${ }^{\circledR}$ $00363103007$ $00363103006$ | Possible contamination | If you have questions about this recall, Apotex Inc. 1-800-706-5575 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Alprazolam tablets 00378400305 | Presence of foreign substance | If you have questions about this recall, Mylan Pharmaceuticals. 1-304-599-2595 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Ranitidine tablets 00363001061,00363001062 ,00363001001, 04903540461, 04903540413,04903540465, 03014250534, 03014250550, 00150062076, 06984287130, 06984287180,06984287137, | Impurity detected in product. | If you have questions about this recall, Dr. Reddy's Laboratories, Inc. 1-309-375-9900 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |

* Drug Recall Class

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NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| 03014213130, 06386848230, 06386848260, 04359880862 04359880865, 07171320302 07171320305, 05789671524, 01167384940, 05511113160, 05511140434,00363013130 00363013180, 06386848024 06386848050,05789671705 |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Ranitidine capsules 05511112960, 05511112905, 05511113030, 05511113001 | Impurity detected in product. | If you have questions about this recall, Dr. Reddy's Laboratories, Inc. 1-309-375-9900 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Atorvastatin calcium 06050525808 | Presence of foreign substance | If you have questions about this recall, Apotex, Inc. 1-800-706-5575 | November $2019$ | Class II |
| Gatifloxacin Ophthalmic Solution ${ }^{(8)}$ 06131467225 | Labeling error | If you have questions about this recall, Sandoz, Inc. 1-609-627-8500 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Neomycin and Polymyxin B Sulfates and Dexamethasone Ophthalmic Suspension 06131463006 | Incorrect or missing package insert | If you have questions about this recall, Sandoz, Inc. 1-609-627-8500 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |

## * Drug Recall Class

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Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.
of Michigan

| Bimatoprost Ophthalmic Solution 00781620675 | Incorrect or missing package insert | If you have questions about this recall, Sandoz, Inc. 1-609-627-8500 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Testosterone Cypionate Injection 05253662501 | Labeling error | If you have questions about this recall, Arbor Pharmaceuticals. 1-866-284-8792 | November $2019$ | Class II |
| Ranitidine tablets 05591009279 | Empty bottles | If you have questions <br> recall, <br> about this  <br> AuroMedeics Pharma <br> LLC.  | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Ranitidine hydrochloride capsules <br> 04229173550,04229173650 | Impurities | If you have questions about this recall, AvKare, Inc. 1-931-292-6222 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Estradiol vaginal inserts 06846271188 | Impurities | If you havequestions <br> about this <br> Glecall,Pharmaceuticals. <br> $721-888$ -$l$ | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Prasugrel tablets ${ }^{\circledR}$ 00378518593 | Failed dissolution specification | If you have questions about this recall, Mylan Pharmaceuticals. 1-304- | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |

* Drug Recall Class

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NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

|  |  | 599-2595 |  |  |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \text { Ibuprofen suspension } \\ & 05167213858,05167221308, \\ & 51672138509,05167221301 \text {, } \\ & 6709132104 \end{aligned}$ | Presence of foreign substance | If you have questions about this recall, Taro Pharmaceuticals. 1-866-923-4914 | $\begin{aligned} & \text { November } \\ & 2019 \end{aligned}$ | Class II |
| Leucovorin Calcium 05074246450 | Presence of particulate matter | If you have questions about this recall, Ingenus Pharmaceutical. 1-877-748-1970 | $\begin{aligned} & \hline \text { October } \\ & 2019 \end{aligned}$ | Class II |
| Pioglitazone Hydrochloride 03334205407 | Superpotent | If you have questions about this recall, Macleods Pharmaceuticals Ltd, 91-22-6676-2800 | $\begin{aligned} & \hline \text { October } \\ & 2019 \end{aligned}$ | Class II |
| Dextroamphetamine Sacharate, Amphetamine Aspartate, Dextroamphetamine Sulfate and Amphetamine Sulfate 01310707301 | Superpotent Drug: Amphetamine Mixed Salts 20mg have been found to be out of specification for weight and thickness. | If you have questions about this recall, Aurobindo Pharma USA Inc. 1-732-839-9400 | $\begin{aligned} & \hline \text { October } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Fentanyl Citrate } ® \\ & 49452003206 \end{aligned}$ | Potential contamination | If you have questions about this recall, Spectrum Laboratory Products. 1-800-772-8786 | $\begin{aligned} & \hline \text { October } \\ & 2019 \end{aligned}$ | Class II |

* Drug Recall Class

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| Ascorbic Acid 07159150050 | Labeling issue | If you have questions about this recall, Atlas Pharmaceutical. 1-410-860-8500 | $\begin{aligned} & \text { October } \\ & 2019 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Pantoprazole Sodium Delayed Release 05974628490 | Discoloration on edges of tablets | If you have questions about this recall, Jubilant Cadista Pharmaceuticals, Inc. 1-410-860-8500 | $\begin{aligned} & \hline \text { October } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { Vivitrol® } \\ & 65757030001 \end{aligned}$ | Mislabeling | If you have questions about this recall, Alkermes, Inc. 1-800-8484876 | $\begin{aligned} & \text { September } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Relpax® } \\ & 00049234005,00049234045 \end{aligned}$ | Microbial contamination | If you have questions about this recall, Pfizer Inc1-877-225-9750 | $\begin{aligned} & \text { September } \\ & 2019 \end{aligned}$ | Class II |
| Hydrocortisone cream with aloe, Hydrocortisone 00363062003, 00363062004 | Potential microbial contamination | If you have questions about this recall, US Pharmaceuticals Inc. 1-888-337-7464 | $\begin{aligned} & \text { August } \\ & 2019 \end{aligned}$ | Class II |
| Lisinopril and hydrochlorothiazide tablets <br> 68180051801, 68180051802 | Brownish/blackish stains on the tablets | If you have questions about this recall, Lupin Limited 1-866-587-4617 | $\begin{aligned} & \text { August } \\ & 2019 \end{aligned}$ | Class II |

[^41]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Blue Cross Blue Care Network of Michigan |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Nitrofurantoin monohydrate/macrocrystals capsules 47781030301 | Failed dissolution specifications | If you have questions about this recall, Alvogen, Inc. 1-866-770-3024 | $\begin{aligned} & \hline \text { August } \\ & 2019 \end{aligned}$ | Class II |
| Macrobid Urinary Tract Antibacterial $52427028501$ | Failed dissolution specifications | If you have questions about this recall, Alvogen, Inc. 1-866-770-3024 | $\begin{aligned} & \text { August } \\ & 2019 \end{aligned}$ | Class II |
| Aspirin and extended-release dipyridamole capsules 60687030532 | Failed impurities/degradation specifications | If you have questions about this recall, American Health Packaging 1-614-492-8177 | $\begin{aligned} & \text { August } \\ & 2019 \end{aligned}$ | Class II |
| Doxycycline hyclate 62584069321 | Failed dissolution specifications | If you have questions about this recall, American Health Packaging | $\begin{aligned} & \text { August } \\ & 2019 \end{aligned}$ | Class II |
| Neomycin $3.5 \mathrm{mg} / \mathrm{g} /$ polymyxin b10000 usp units/g / dexamethasone $1 \mathrm{mg} / \mathrm{g}$ ophthalmic ointment 52959040701 | Insufficient quality | If you have questions about this recall, H.J. Harkins Company 1-805-929-1333 | July 2019 | Class II |
| $\begin{aligned} & \text { Diphenhydramine } \\ & 49035033002,49035033096 \text {, } \\ & 49035033045 \end{aligned}$ | Contamination | If you have questions about this recall, LNK International 1-631-5433787 | July 2019 | Class II |

[^42]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Blue Cross Blue Shield Blue Care Network of Michigan |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Norethindrone and ethinyl estradiol $68180090313$ | Failed Impurities | If you have questions about this recall, Lupin Pharmaceuticals 1-800-399-2561 | July 2019 | Class II |
| Milrinone lactate 00409277602,00409277623 | Bags have the potential to leak | If you have questions about this recall, Pfizer Inc. 1-800-879-3477 | $\begin{aligned} & \text { July } \\ & 2019 \end{aligned}$ | Class II |
| Fluorouracil injection 63323011710, 63323011719, 63323011728, 63323011761, 63323011761, 63323011769, 63323011718, 63323011720, 63323011751, 63323011759, 63323011768 | Glass Particles | If you have questions about this recall, Fresenius Kabi 1-888-3861300 | $\begin{aligned} & \hline \text { June } \\ & 2019 \end{aligned}$ | Class I |
| $\begin{aligned} & \text { Heparin Sodium® } \\ & 00264958720 \end{aligned}$ | Not potent | If you have questions about this recall, Braun Medical Inc 1-800-5239676 | $\begin{aligned} & \text { June } \\ & 2019 \end{aligned}$ | Class II |
| Estradiol vaginal inserts 68462071171, 68462071188 | Defective delivery | If you have questions about this recall, Glenmark Pharmaceuticals Inc 1-201-684-8000 | $\begin{aligned} & \text { June } \\ & 2019 \end{aligned}$ | Class II |

* Drug Recall Class

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Class $\mathbf{3}$ Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Blue Cross Blue Care Network of Michigan |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Zyflo Cr $®$ ® 10122090212,10122090220 | Failed dissolution | If you have questions about this recall, Chiesi USA, 1-919-678-6611 | $\begin{aligned} & \text { June } \\ & 2019 \end{aligned}$ | Class II |
| Pramipexole dihydrochloride 68084079325, 68084097425 | Possible cross contamination | If you have questions about this recall, American Health Packaging, 1-800-707-4621 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { Zileuton® } \\ & 66993048532 \end{aligned}$ | Failed dissolution | If you have questions about this recall, Chiesi USA, 1-919-678-6611 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { Testosterone } \\ & 69699170230,69699170210 \end{aligned}$ | Lack of sterility | If you have questions about this recall, Pharm D. Solutions 1-713-790-1693 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| Hydrocodone bitartrate and homatropine methylbromide oral solution $59741026216,13668057710$ | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { Risperidone } \\ & 23155031751 \end{aligned}$ | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |

[^43]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| $\begin{aligned} & \hline \text { Cetirizine hydrochloride } \\ & 23155029251,23155029252, \\ & 13668059607,13668059611 \end{aligned}$ | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Phenylephrine hydrochloride 00536138912, 00536138935, 00904768822 | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{array}{\|l\|} \hline \text { Bisacodyl } \\ 00536135501,00536135512 \end{array}$ | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| Hydrocortisone acetate suppositories 59741030101, 59741030124, $59741030112$ | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| Memantine hydrochloride 39328055112 <br> 13668057309 | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| Lactulose solution 13668058008, 13668058012, 13668057408, 13668057412, | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-912- | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |

## * Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| 13668057410 |  | 9561 |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Pseudoephedrine Hydrochloride 00536185085, 00536185097 | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| Diphenhydramine hydrochloride 00536077085, 00536077097 | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| Guaifenesin \& dextromethorphan hydrobromide 00536097085,00536097097 | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \hline \text { May } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Guaifenesin } \\ & 00536082585,00536082597 \end{aligned}$ | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Acetaminophen } \\ & 00536012285,00536012297 \end{aligned}$ | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| $\begin{aligned} & \hline \text { Phenobarbital } \\ & 16571033016 \end{aligned}$ | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Guaifenesin, Codeine Phosphate \& Pseudoephedrine Hydrochloride 16571030116 | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Guaifenesin AC } \\ & 16571030216 \end{aligned}$ | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Acetic Acid } \\ & 64980042415 \end{aligned}$ | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Hyoscyamine sulfate } \\ & 39328004715,39328004816 \end{aligned}$ | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| Cyproheptadine hydrochloride 39328004416 | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-912- | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
Class $\mathbf{2}$ Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

|  |  | 9561 |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Bisacodyl 00904505812,00904505860 | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| Pedia Relief 00904505020 | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| Robafen dm cough sugar free clear $00904630620$ | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| Oxymetazoline Hydrochloride 00904571130,00904571135 | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| Pseudoephedrine hydrochloride 00536185085,00536185097 | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |

* Drug Recall Class

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Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
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| $\begin{aligned} & \hline \text { Banofen } \\ & 904517416 \end{aligned}$ | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| :---: | :---: | :---: | :---: | :---: |
| Robafen DM 00904005300,00904005316 00904005309 | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { Robafen AC } \\ & 00904647916 \end{aligned}$ | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { Risperidone } \\ & 13668058906 \end{aligned}$ | Contamination with burkholderia | If you have questions about this recall, Torrent Pharma Inc 1-800-9129561 | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { Pecgen dmx } \\ & 52083063016 \end{aligned}$ | Incorrect labeling instructions | If you have questions about this recall, Novis PR 1-800-727-6711 | $\begin{aligned} & \hline \text { May } \\ & 2019 \end{aligned}$ | Class I |
| Cefdinir for oral suspension 68180072310, 68180072320 | Metal piece identified in the product bottle | If you have questions about this recall, Lupin Pharmaceuticals Inc 1- | $\begin{aligned} & \text { May } \\ & 2019 \end{aligned}$ | Class II |

* Drug Recall Class

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Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.
Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

|  |  | $410-576-2000$ |  |  |
| :--- | :--- | :--- | :--- | :--- |

* Drug Recall Class

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Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

|  |  | 875-0123 option 5 |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Losartan Potassium tab 68788004809 | Presence of an impurity | If you have questions about this recall, Preferred Pharmaceuticals Inc 1-714-777-3729 | $\begin{aligned} & \text { April } \\ & 2019 \end{aligned}$ | Class II |
| Carvedilol tablets 65841061605 | Label Mix-up | If you have questions about this recall, Zydus Pharmaceuticals 1-877-993-8779 | $\begin{aligned} & \text { April } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Ketorolac tromethamine } \\ & 25021070101,25021070102 \end{aligned}$ | Lack of sterility | If you have questions about this recall, Sagent Pharmaceuticals 1-866-625-1618 | $\begin{aligned} & \text { April } \\ & 2019 \end{aligned}$ | Class II |
| Acyclovir tablets 68382079101, 68382079106, 68382079116, 68382079105, 68382079110,68382079130 | Label mix-up | If you have questions about this recall, Zydus Pharmaceuticals 1-877-993-8779 | $\begin{aligned} & \text { April } \\ & 2019 \end{aligned}$ | Class II |
| Fentanyl transdermal system, 12 $\mathrm{mcg} / \mathrm{h}$ $47781042347$ | Label mix-up | If you have questions about this recall, Alvogen 1-866-770-3024 | $\begin{aligned} & \text { April } \\ & 2019 \end{aligned}$ | Class I |
| Losartan Potassium/hctz 70518156000 | Presence of an impurity | If you have questions about this recall, Remedy | $\begin{aligned} & \text { April } \\ & 2019 \\ & \hline \end{aligned}$ | Class II |

* Drug Recall Class

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Class 3 Recall: Using the drug is not likely to cause adverse health consequences.
NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

|  |  | Repack 1-866-845-3791 |  |  |
| :--- | :--- | :--- | :--- | :--- |
| Losartan Potassium and <br> hydrochlorothiazide tablets <br> 13668011610, 13668011674, <br> 13668011630, 13668011690, <br> 13668011710, 13668011774, <br> 13668011730, 13668011790, <br> 13668011810, 13668011874, <br> 13668011830, 13668011890 |  | If you have questions <br> about this recall, Torrent <br> Pharmaceuticals 1-800- <br> $912-9561$ | April <br> 2019 | Class II |
| Losartan Potassium <br> 68645049454 |  | Presence of an impurity |  |  |

* Drug Recall Class

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| 43386-0050-19 |  | about this recall, Lupin Pharmaceuticals 1-800-399-2561 | 2019 |  |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \text { Combigan } \\ & 00023921103,00023921110, \\ & 00023921105,00023921115 \end{aligned}$ | Trace amounts of an impurity | If you have questions about this recall, Ecolab 1-800-433-8871 | $\begin{aligned} & \text { March } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { Pravastatin Sodium } \\ & \text { 68462-0196-05, 68462-0196-90 } \end{aligned}$ | Foreign tablet in bottle | If you havequestions <br> about this recall,GlenmarkPharmaceuticals, Inc. 201-$684-8000$ | $\begin{aligned} & \text { March } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Losartan } \\ & \text { 70518-0588-01 } \end{aligned}$ | Trace amounts of impurity | If you have questions about this recall, Remedy repack Inc 1-886-8453791 | $\begin{aligned} & \text { March } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \text { Phenobarbital } \\ & \text { 70166-0536-02 } \end{aligned}$ | The label contains the incorrect expiration date | If you have questions about this recall, Lohxa LLC 1-800-641-5564 | $\begin{aligned} & \text { March } \\ & 2019 \end{aligned}$ | Class II |
| Losartan Potassium 31722-0702-30 | Impurity found | If you have questions about this recall, Pharma Pac 1-805-929-1333 | $\begin{aligned} & \text { March } \\ & 2019 \end{aligned}$ | Class II |
| Volumex | Lack of assurance | If you have questions | March | Class II |

* Drug Recall Class

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| 50914-7720-08 |  | about this recall, Iso-Tex Diagnostics $1-800-477-$ 4539 | 2019 |  |
| :---: | :---: | :---: | :---: | :---: |
| Hydrocortisone and Acetic Acid 50383-0301-10 | Sub Potent Drug | If you have questions about this recall, call Akon Inc. 1-800-477-4539 | $\begin{array}{\|l\|} \hline \text { March } \\ 2019 \end{array}$ | Class II |
| $\begin{aligned} & \text { Losartan Potassium } \\ & 68645-0578-54 \end{aligned}$ | Trace amounts of impurity | If you have questions about this recall, Call Legacy 1-877-538-8443 | $\begin{aligned} & \text { February } \\ & 2019 \end{aligned}$ | Class II |
| Losartan <br> Potassium/Hydrochlorothiazide <br> 13668-0118-10, 13668-0118-74, <br> 13668-0118-30,13668-0118-90, <br> 13668-0117-10, 13668-0117-74, <br> 13668-0117-30, 13668-0117-90, <br> 13668-0116-10, 13668-0116-74, <br> 13668-0116-30, 13668-0116-90 | Presence of Impurity | If you have questions about this recall, Call Torrent Pharmaceuticals 1-800-912-9561 | $\begin{aligned} & \text { February } \\ & 2019 \end{aligned}$ | Class II |
| Losartan Potassium <br> 13668-0113-10, 13668-0113-90, <br> 13668-0113-74, 13668-0409-10, <br> 13668-0409-74, 13668-0409-30, <br> 13668-0409-90, 13668-0115-10, <br> 13668-0115-74, 13668-0115-30, <br> 13668-0115-90 | Presence of an impurity | If you have questions about this recall, Call Torrent Pharmaceuticals 1-800-912-9561 | $\begin{aligned} & \text { February } \\ & 2019 \end{aligned}$ | Class II |
| Valsartan | Presence of an impurity | If you have questions | February | Class II |

* Drug Recall Class

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| $\begin{aligned} & \text { 52343-0122-30, 52343-0123-90, } \\ & 52343-0124-90,52343-0125-90 \end{aligned}$ |  | about this recall, Call Preferred Pharmaceuticals 1-855-544-9419 | 2019 |  |
| :---: | :---: | :---: | :---: | :---: |
| Losartan Potassium 68788-6882-03, 68788-6882-09 | Presence of an impurity | If you have questions about this recall, Call Preferred Pharmaceuticals 1-855-544-9419 | $\begin{aligned} & \text { February } \\ & 2019 \end{aligned}$ | Class II |
| $\begin{aligned} & \hline \text { Valsartan } \\ & \text { 60687-0139-01 } \end{aligned}$ | Trace amounts of an impurity | If you have questions about this recall, call Aurobindo 1-800-9129572 | $\begin{aligned} & \text { February } \\ & 2019 \end{aligned}$ | Class II |
| Losartan Potassium <br> 31722-0702-05, 31722-0702-30, <br> 31722-0702-90, 31722-0702-10, <br> 31722-0702-60, 50268-0517-15, <br> 50268-0513-15, 50268-0514-15, <br> 68645-0577-54 | Trace amounts of an impurity | If you have questions about this recall, call Aurobindo 1-800-9129572 | $\begin{aligned} & \text { February } \\ & 2019 \end{aligned}$ | Class II |
| Alprazolam 51079-0788-20 | Failed impurities/degradation | If you have questions about this recall, call Mylan 1-888-406-9305 | $\begin{array}{\|l} \hline \text { January } \\ 2019 \end{array}$ | Class II |
| Ceftriaxone for Injection 68180-0611-01 | Presence of Particulate Matter: Product complaints received of grey flecks, identified as shredded rubber particulate matter from the stopper | If you have questions about this recall, Lupin Pharmaceuticals Inc. 111 S Calvert St FI 21ST | $\begin{array}{\|l} \hline \text { January } \\ 2019 \end{array}$ | Class I |

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NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Blue Cross Blue Care Network of Michigan |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
|  | observed in reconstituted vials | Baltimore, MD 21202- 6174 |  |  |
| Vecuronium Bromide for Injection 10 mg 47335-0931-44 | Presence of Particulate Matter: Foreign matter identified as glass detected in Vecuronium Bromide for Injection. | If you have questions about this recall, Sun Pharmaceutical Industries, Inc. <br> 270 Prospect Plains Rd Cranbury, NJ 08512-3605 | $\begin{aligned} & \text { January } \\ & 2019 \end{aligned}$ | Class I |
| $\begin{aligned} & \text { OZURDEX } \\ & \text { 0023-3348-07 } \end{aligned}$ | Deviations: A silicone particulate was noted in Ozurdex | If you have questions about this recall, Allergan, PLC. <br> 5 Giralda Farms Madison, NJ 07940-1027 (714) 246-4500 | $\begin{aligned} & \text { January } \\ & 2019 \end{aligned}$ | Class II |
| Estradiol Vaginal Inserts 68462-711-71 68462-711-88 | Defective Delivery System: Customer complaints of malfunctioning plunger of the applicator | If you have questions <br> about this recall, <br> Glenmark  <br> Pharmaceuticals Inc., <br> USA  <br> 750 Corporate Dr <br> Mahwah, NJ 07430-2009  | $\begin{aligned} & \text { January } \\ & 2019 \end{aligned}$ | Class II |
| Dianeal Low Calcium 00941-0424-52 | Lack of Assurance of Sterility: Confirmed customer complaints for | If you have questions about this recall, Baxter | $\begin{aligned} & \hline \text { January } \\ & 2019 \\ & \hline \end{aligned}$ | Class II |

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| Blue Cross Blue Care Network of Michigan |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Lot \#: Y281477 <br> Expiration: 02/2020 | leaks on the tubing | Healthcare Corporation 1 Baxter Pkwy Deerfield, IL 60015-4625 |  |  |
| Cefdinir for Oral Suspension USP <br> $125 \mathrm{mg} / 5 \mathrm{~mL}$ 68180-0722-10 68180-0722-20 | Deviations: Product complaints received indicating reconstituted suspension was observed to be thick | If you have questions about this recall, Lupin Pharmaceuticals Inc. 111 S Calvert St FI 21ST Baltimore, MD 212026174 866-587-4617 | $\begin{array}{\|l} \hline \text { January } \\ 2019 \end{array}$ | Class II |
| Cidofovir Injection 375mg 23255-0216-31 | Lack of Assurance of Sterility: complaints received about dried powder on the outside of bottle | If you have questions about this recall, Heritage Pharmaceuticals, Inc. 1 Tower Center Blvd Ste 1700 <br> East Brunswick, NJ 08816-1145 <br> 732) 429-1000 | $\begin{array}{\|l} \hline \text { January } \\ 2019 \end{array}$ | Class II |
| $\begin{aligned} & \text { Cephalexin for Oral Suspension } \\ & \text { USP } \\ & 68180-0124-01 \\ & 68180-0124-02 \end{aligned}$ | Deviation; manufacturing batch record could not be located | If you have questions about this recall, Lupin Pharmaceuticals Inc. 111 S Calvert St FI 21ST Baltimore, MD 21202 | $\begin{array}{\|l} \hline \text { January } \\ 2019 \end{array}$ | Class II |

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NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

| Blue Sross Blue Care Network of Michigan |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \text { Valsartan tablets USP } 320 \mathrm{mg} \\ & 65862-0573-05 \\ & 65862-0573-90 \end{aligned}$ | Deviations: FDA lab confirmed <br> presence an impurity, presence an impurity, Nnitrosodimethylamine (NDEA) contained in the API used to manufacture the product above the interim acceptable daily intake level of 0.083 parts per million. | If you have questions about this recall, Aurobindo Pharma USA Inc. <br> 279 Princeton Hightstown <br> Rd <br> East Windsor, NJ 08520- <br> 1401 $732-839-9400$ | $\begin{aligned} & \text { January } \\ & 2019 \end{aligned}$ | Class II |
| Valsartan and Hydrochlorothiazide tablets $\begin{aligned} & 65862-0549-10 \\ & 65862-0549-90 \\ & 65862-0549-99 \end{aligned}$ | Deviations: FDA lab confirmed presence an impurity, Nnitrosodimethylamine (NDEA) contained in the API used to manufacture the product above the interim acceptable daily intake level of 0.083 parts per million | If you have questions about this recall, Aurobindo Pharma USA Inc. <br> 279 Princeton Hightstown <br> Rd <br> East Windsor, NJ 08520 <br> 732-839-9400 | $\begin{aligned} & \text { January } \\ & 2019 \end{aligned}$ | Class II |
| Amlodipine and Valsartan Tablets USP $10 \mathrm{mg} / 320 \mathrm{mg}$ $\begin{aligned} & 65862-0740-03 \\ & 65862-0740-30 \\ & 65862-0740-90 \end{aligned}$ | Deviations: FDA lab confirmed presence an impurity, Nnitrosodimethylamine (NDEA) contained in the API used to manufacture the product above the interim acceptable daily intake level | If you have questions about this recall, Aurobindo Pharma USA Inc. <br> 279 Princeton Hightstown <br> Rd <br> East Windsor, NJ 08520- | $\begin{aligned} & \text { January } \\ & 2019 \end{aligned}$ | Class II |

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Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

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| $\text { (i) (\$) } \begin{aligned} & \text { Blue Cross } \\ & \text { Bue Shield } \\ & \text { Biue Care } \\ & \text { of Michigan Network } \end{aligned}$ |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
|  | of 0.083 parts per million. | $\begin{aligned} & 1401 \\ & \underline{7328399400} \\ & \hline \end{aligned}$ |  |  |
| EEMT (esterified estrogens and methyltestosterone 15310-0010-01 | The combination of esterified estrogens and methyltestosterone is used to treat symptoms of menopause such as hot flashes, and vaginal dryness, burning, and irritation. | If you have questions about this recall, Syntho Pharmaceuticals, Inc. 230 Sherwood Ave Farmingdale, NY 117351718 631-755-9898 | $\begin{aligned} & \text { January } \\ & 2019 \end{aligned}$ | Class II |
| Olmesartan Medoxomil and Hydrochlorothiazide Tablets, 40 $\mathrm{mg} / 25 \mathrm{mg}$ 00093761756 00093761798 | Failed dissolution specifications | If you have questions about this recall, Teva Pharmaceuticals USA 1090 Horsham Rd North Wales, PA 194541505 <br> 888-TEVA-USA (888-8382872) | $\begin{aligned} & \text { January } \\ & 2019 \end{aligned}$ | Class II |
| Fluocinolone Acetonide Topical Solution, USP, 0.01 \%, 60 mL bottle <br> Lot \#: S700214, Exp Apr-19; | Failed Impurities/Degradation Specifications: Expansion of October 2018 recall due to elevated out of specification results for total impurities that have been chemically | If you have questions about this recall, Contact: <br> LUPIN SOMERSET <br> 400 Campus Dr <br> Somerset, NJ 08873-1145 | $\begin{array}{\|l\|} \hline \text { January } \\ 2019 \end{array}$ | Class II |

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| S700447, Exp Jun-19; S700787, Exp Oct-19; S701057, Exp Nov19; S800107, Exp Feb-20; <br> S800266, Exp Mar-20; S800524, Exp May-20; S800791 Kul Exp Jul-20 | identified as oxidative degradation products of the fluocinolone active pharmaceutical ingredient |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Ezetimibe and Simvastatin Tablets $10 \mathrm{mg} / 80 \mathrm{mg}, 1000$-count bottles <br> Lot \#: 43E021 and 43E023, Exp. 01/2020 | Presence of Foreign Substance: Product complaint of black speckles observed on tablets. | If you have questions about this recall, Contact: <br> Dr. Reddy's Laboratories, Inc. <br> 107 College Rd E <br> Princeton, NJ 08540-6623 | $\begin{aligned} & \text { January } \\ & 2019 \end{aligned}$ | Class II |
| Losartan potassium and hydrochlorothiazide $1000 \mathrm{mg} / 25 \mathrm{mg}$ tablets <br> ONLY Lot: JB8912 | This product is being recalled due to th presence of an impurity, N nitrosodiethylamine. NDEA occurs naturally in manufacturing processes and could cause cancer based on laboratory test results. No adverse effects related to this recall have been reported to date. | If you have questions about this recall, Contact Sandoz Inc. at 1-800-5258747 Monday-Friday 8:30 AM - 5:00 PM (EST) or email usdrugsafety.operations@ novartis.com. | November 2018 | Class I |
| Montelukast Sodium 10Mg | This medicine is made by Camber Pharmaceuticals, Inc. This medicine is being recalled | If you have questions about this recall, call Camber Pharmaceuticals | $\begin{aligned} & \text { September } \\ & 2018 \end{aligned}$ | Class I |

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.
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Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

|  | because sealed bottles labeled as Montelukast sodium tablets, 10 mg , 30-count bottle from Camber were found to instead contain 90 tablets of losartan potassium tablets, 50 mg . This could result in a life-threatening condition. | Med Line at 1-866-4951995 |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Diphenoxylate Hydrochloride and Atropine Sulfate tablets, 2.5 $\mathrm{mg} / 0.025 \mathrm{mg}$ (Lomotil). | This medication is made by Greenstone LLC, a wholly owned subsidiary of Pfizer, Inc. This medication is being recalled due to super potency. The use of the impacted super potent product when used as labeled has a low probability of being associated with adverse events of limited severity such as lethargy, skin flush, and drowsiness. Serious adverse events such as coma and respiratory depression are improbable. | If you have questions about this recall, please contact Pfizer Customer Support at 800-879-3477. | $\begin{aligned} & \text { January } \\ & 2018 \end{aligned}$ | Class I |
| EpiPen and EpiPen Jr AutoInjector | This medication is made by Mylan. This medication is being recalled due to failure to activate the device or increased force needed to activate | If you have questions about this recall, please contact your physician, pharmacy or Mylan | May 2017 | Class I |

[^44]NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

|  | due to a potential defect in a supplier <br> component. The potential defect <br> could make the device difficult to <br> activate in an emergency. This could <br> result in a life-threatening condition. | Customer Relations at <br> $800-796-9526$ or <br> customer.service@mylan. <br> com. |  |  |
| :--- | :--- | :--- | :--- | :--- |
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|  |  |  |  |  |
|  |  |  |  |  |

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