The Board of Pharmacy has received notice of the following product recall. The Board strongly encourages pharmacies to immediately review their quality assurance and recall policies and procedures to determine if any corrective action is required.

| Description | Lot # | Exp Date | NDC | UPC |
|---|---|---|---|---|
| ACCURETIC TAB 10/12.5MG 90 | EA6686 | 04/30/2022 | 00071022223 | 30071022223 |
| ACCURETIC TAB 10/12.5MG 90 | FG5379 | 08/31/2024 | 00071311223 | 30071311223 |
| ACCURETIC TAB 20/12.5MG 90 | EA6665 | 04/30/2022 | 00071022023 | 30071022023 |
| ACCURETIC TAB 20/12.5MG 90 | CN0640 | 04/30/2022 | 00071022023 | 30071022023 |
| ACCURETIC TAB 20/12.5MG 90 | FG5381 | 08/31/2024 | 00071521223 | 30071521223 |
| ACCURETIC TAB 20/25MG 90 | ET6974 | 02/28/2023 | 00071022323 | 30071022323 |
| QUINAPRIL/HCTZ 20/12.5 GRE 90 | ED3905 | 03/31/2023 | 59762022001 | 35976202201 |
| QUINAPRIL/HCTZ 20/12.5 GRE 90 | ED3904 | 03/31/2023 | 59762022001 | 35976202201 |
| QUINAPRIL/HCTZ 20/12.5 GRE 90 | DN6931 | 03/31/2023 | 59762022001 | 35976202201 |
| QUINAPRIL/HCTZ 20/25MG GRE 90 | DP3414 | 02/28/2023 | 59762022301 | 35976202231 |
| QUINAPRIL/HCTZ 20/25MG GRE 90 | FE3714 | 02/28/2023 | 59762522509 | 35976252259 |

Pfizer is recalling the above item(s)/lot(s) due to the presence of n-nitroso-quinapril above the Acceptable Daily Intake (ADI) level. This recall is to the consumer level. Affected product started shipping November 2019.