

The Board of Pharmacy has received notice of the following product recall:

| Description | Lot # | Exp Date | NDC | UPC |
|--------------------------|-------------|----------|------------------|------------|
| HM FEXOFEN TAB 180MG 15 | 067180010B | 3/31/21 | 62011031501 | 5256914055 |
| HM FEXOFEN TAB 180MG 30 | 067180016A | 5/31/21 | 62011031502 | 5256914056 |
| SM FEXOFEN TAB 180MG 15 | 067180024F1 | 7/31/21 | 70677000801 | 1093983144 |
| SM FEXOFEN TAB 180MG 30 | 067180016B | 5/31/21 | 70677000802 | 1093983244 |
| FEXOFENADINE HCl TABLETS | 06718027A1 | 7/31/21 | 00363009755 | |
| FEXOFENADINE HCl TABLETS | 067180024A1 | 7/31/21 | 46122038723 | |
| FEXOFENADINE HCl TABLETS | 067180010A | 3/31/21 | 49035099562 | |
| | 067180024D1 | 7/31/21 | | |
| | 067180023C1 | 7/31/21 | | |
| FEXOFENADINE HCl TABLETS | 067180024B1 | 7/31/21 | 53943002109 | |
| FEXOFENADINE HCl TABLETS | 067180008A | 3/31/21 | 58602071121 | |
| FEXOFENADINE HCl TABLETS | 067180025B1 | 7/31/21 | 60000040930 | |
| FEXOFENADINE HCl TABLETS | 067180025C1 | 7/31/21 | 60000040945 | |
| FEXOFENADINE HCl TABLETS | 067180025A1 | 7/31/21 | 60000040948 | |
| FEXOFENADINE HCl TABLETS | 067180025D1 | 7/31/21 | 60000040953 | |
| FEXOFENADINE HCl TABLETS | 067180009A | 3/31/21 | 68196097691 | |
| | 067180013A | 4/30/21 | | |
| | 067180014A | 4/30/21 | | |
| | 067180015A | 4/30/21 | | |
| | 067180018A | 5/31/21 | | |
| | 067180020A | 6/30/21 | | |
| | 067180021A1 | 7/31/21 | | |
| | 067180022A1 | 7/31/21 | | |
| | 06718028A1 | 9/30/21 | | |
| | 06718028B1 | 9/30/21 | | |
| 06719001A3 | 1/31/22 | | | |
| FEXOFENADINE HCl TABLETS | 067180011A | 4/30/21 | BULK BRITE STOCK | |
| FEXOFENADINE HCl TABLETS | 067180012A | 4/30/21 | BULK BRITE STOCK | |
| FEXOFENADINE HCl TABLETS | 06718027B1 | 9/30/21 | BULK SHIPMENT | |

Aurobindo is recalling the above items/lots due to a stability failure. This recall is to the retail pharmacy level. Affected product started shipping June 20, 2018.

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