

## Drug Recall

**DATE OF RECALL:** September 28, 2023

**DRUG NAME:** BREXAFEMME® tablets

**RECALLING FIRM:** SCYNEXIS, Inc.

**REASON FOR RECALL:** This recall was issued due to potential cross contamination with a non-antibacterial  $\beta$ -lactam drug substance in the ibrexafungerp citrate used to manufacture the BREXAFEMME® tablets.

**FDA LINK:** <https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/scynexis-issues-voluntary-nationwide-recall-brexafemmer-ibrexafungerp-tablets-due-potential-cross>

**OTHER DETAILS:** N/A

**RECALLED PRODUCT:**

| Product Description | NDC Number    | Lot Number               | Expiration Date    |
|---------------------|---------------|--------------------------|--------------------|
| BREXAFEMME® tablets | 75788-0115-04 | LF21000008<br>LF22000051 | 11/2023<br>11/2025 |