

IMPORTANT RECALL NOTICE FOR BREXAFEMME® 150 MG TABLETS

Dear Member,

Your health is important to us. On September 28, 2023, the U.S. Food and Drug Administration (FDA) announced a drug recall notice. It is for two lots of **BREXAFEMME®**. The FDA issued the recall because the product may have a safety concern.

Products that may be affected by this recall:

Drug Name	NDC	Lot#	Expiration Date
Brexafemme 150 mg tablet	NDC 75788-115-04	LF21000008	November 30, 2023
Brexafemme 150 mg tablet	NDC 75788-115-04	LF22000051	November 30, 2025

Additional information is available at: https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/scynexis-issues-voluntary-nationwide-recall-brexafemmer-ibrexafungerp-tablets-due-potential-cross?utm_medium=email&utm_source=govdelivery

Please call your pharmacy to find out if your drug could be part of this recall. The pharmacy may replace the drug. We are writing this letter to give you information. This should not replace your doctor's advice. Only your doctor can decide what drugs are right for you.

Please disregard this letter if you are not currently taking **BREXAFEMME®** or if this safety warning does not apply to you. Our records may not be complete, or your doctor may have discontinued this medication.

Sincerely,

Doctors HealthCare Plans, Inc.

ATENCIÓN: Si habla español, tiene a su disposición servicios gratuitos de asistencia lingüística. Llame al 786-460-3427 o 833-342-7463(TTY: 711).

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