

The Medical Letter[®]

on Drugs and Therapeutics

Volume 65

February 20, 2023

ISSUE No.
1670

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Volume 65 (Issue 1670)

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IN BRIEF

Expanded Access to Mifepristone

First approved in 2000 for termination of pregnancies of ≤ 49 days' gestation, the indication for the oral antiprogestin mifepristone (*Mifeprex*, and generics) was expanded in 2016 to include pregnancies of up to 10 weeks' gestation. A single 200-mg oral dose of mifepristone followed 24-48 hours later by a single 800-mcg buccal dose of the prostaglandin analog misoprostol terminates early intrauterine pregnancies in about 95% of women.

Before the COVID-19 pandemic, mifepristone could be dispensed to a patient only in-person by a certified prescriber. In late 2021, the FDA modified the Mifepristone Risk Evaluation and Mitigation Strategy (REMS) Program to allow mifepristone to be dispensed by mail by certified prescribers or pharmacies.¹ The FDA has now announced that certified retail pharmacies will be permitted to dispense the drug to patients who present a prescription from a certified prescriber.² To become certified, prescribers and pharmacies must complete agreement forms attesting to certain requirements. The patient is also required to complete an agreement cosigned by the prescriber. Retail pharmacies will not be permitted to dispense mifepristone in states that have passed laws forbidding medical termination of pregnancy. ■

1. In brief: Mifepristone by mail for pregnancy termination. *Med Lett Drugs Ther* 2022; 64:11.
2. FDA. Information about mifepristone for medical termination of pregnancy through ten weeks gestation. Available at: <https://bit.ly/3GSebTH>. Accessed January 9, 2023.

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