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AccessMatters Applauds FDA Decision to Update Protocol for Medication Abortion

PHILADELPHIA, PA (April 1, 2016): Earlier this week, the U.S. Food and Drug Administration announced new guidelines for prescribing Mifepristone, a drug used for medication abortion. Melissa Weiler Gerber, President and CEO of AccessMatters released the following statement:

“AccessMatters supports the U.S. Food and Drug Administration’s new guidelines which will expand access to medication abortion. As a public health advocate and Title X safety-net provider, AccessMatters is committed to ensuring women are able to exercise a full range of reproductive health options, including prenatal care and delivery; foster care; adoption; and abortion services. AccessMatters works to ensure providers are trained to deliver unbiased pregnancy options counseling in safe, confidential settings. The FDA’s decision to reduce the number of visits to a healthcare provider needed to access the medication, to extend the period of time into a pregnancy that a woman can take the medication, and to reduce the dosage needed, all align with standard medical practice. This decision is a victory for science over politics and for all women nationwide.”

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[AccessMatters](http://www.AccessMatters.org) is the catalyst for providing access to sexual and reproductive health care for teens and adults in need reaching over 200,000 individuals annually. For more information about AccessMatters programs, visit www.AccessMatters.org, [@AccessMatters4U](https://twitter.com/AccessMatters4U) on Twitter, and [@AccessMatters](https://www.facebook.com/AccessMatters) on Facebook.