GILEAD | Medical Information

Descovy for PrEP® (FTC/TAF) Clinical Practice Guidelines

This document is in response to your request for information regarding Descovy for PrEP® (emtricitabine/tenofovir alafenamide [FTC/TAF] for HIV-1 pre-exposure prophylaxis) and clinical practice guidelines.

This document includes content from, or references to clinical practice guidelines and should not be interpreted as a treatment recommendation or an endorsement of the guidelines by Gilead Sciences.

The full indication, important safety information, and boxed warnings are available at: www.gilead.com/-/media/files/pdfs/medicines/hiv/descovy/descovy_pi.

Clinical Practice Guidelines

Guidelines from the International Antiviral Society–USA (IAS-USA) panel on antiretroviral drugs for the prevention and treatment of HIV in adults can be accessed at: jamanetwork.com/journals/jama/fullarticle/2827545.

The IAS-USA panel published a brief update of its 2024 recommendations for the prevention of HIV following new data on the use of FTC/TAF among cisgender women, which can be accessed at: jamanetwork.com/journals/jama/fullarticle/2835835.

The Centers for Disease Control and Prevention clinical practice guidelines of PrEP for the prevention of HIV in the US can be accessed at: stacks.cdc.gov/view/cdc/159891.

Guidelines from the World Health Organization for HIV treatment and PrEP can be accessed at: www.who.int/publications/i/item/9789240031593.

The European AIDS Clinical Society produces guidelines for the management of treatment and prevention of HIV in Europe, which can be accessed at: www.eacsociety.org/guidelines/eacs-guidelines.

Product Label

For the full indication, important safety information, and boxed warning(s), please refer to the Descovy US Prescribing Information available at: www.gilead.com/-/media/files/pdfs/medicines/hiv/descovy/descovy_pi.pdf.

Follow Up

For any additional questions, please contact Gilead Medical Information at:

☎1-866-MEDI-GSI (1-866-633-4474) or 🖱 www.askgileadmedical.com

Adverse Event Reporting

Please report all adverse events to:

Gilead Pharmacovigilance and Epidemiology ☎ 1-800-445-3235, option 3 or 🖱 www.gilead.com/utility/contact/report-an-adverse-event

FDA MedWatch Program by ☎ 1-800-FDA-1088 or ✉MedWatch, FDA, 5600 Fishers Ln, Rockville, MD 20852 or 🖱 www.accessdata.fda.gov/scripts/medwatch

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