



# Biktarvy® (BIC/FTC/TAF)

## Effects of Adherence on PK

This document is in response to your request for information regarding Biktarvy® (bictegravir/emtricitabine/tenofovir alafenamide [BIC/FTC/TAF]) and the effects of adherence on pharmacokinetics (PK).

Some data may be outside of the US FDA-approved Prescribing Information. In providing this data, Gilead Sciences, Inc. is not making any representation as to its clinical relevance or to the use of any Gilead product(s). For information about the approved conditions of use of any Gilead drug product, please consult the FDA approved prescribing information.

**The full indication, important safety information, and boxed warnings are available at:**  
**[http://www.gilead.com/~media/files/pdfs/medicines/hiv/biktarvy/biktarvy\\_pi](http://www.gilead.com/~media/files/pdfs/medicines/hiv/biktarvy/biktarvy_pi)**

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## Available Data

Currently, there is no data available regarding effects of adherence on PK and BIC/FTC/TAF.

A literature search was conducted in Ovid MEDLINE and Embase databases for studies published between 1946 and June 16, 2025 using search terms that included bictegravir, tenofovir alafenamide, emtricitabine, adherence, pharmacokinetics, and other related search terms. No citations relevant to your inquiry were identified.

## Product Label

For the full indication, important safety information, and boxed warning(s), please refer to the Biktarvy US Prescribing Information available at:

[http://www.gilead.com/~media/files/pdfs/medicines/hiv/biktarvy/biktarvy\\_pi](http://www.gilead.com/~media/files/pdfs/medicines/hiv/biktarvy/biktarvy_pi)

## Follow Up

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

☎ 1-866-MEDI-GSI (1-866-633-4474) or 🌐 [www.askgileadmedical.com](http://www.askgileadmedical.com)

## Adverse Event Reporting

Please report all adverse events to:

Gilead Global Patient Safety ☎ 1-800-445-3235, option 3 or

🌐 [www.gilead.com/utility/contact/report-an-adverse-event](http://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](http://www.accessdata.fda.gov/scripts/medwatch)

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