GILEAD | Medical Information

Yeztugo® (lenacapavir) Use After Bariatric Surgery

This document is in response to your request for information regarding the use of Yeztugo® (lenacapavir [LEN]) after bariatric surgery.

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 warning are available at: www.gilead.com/-/media/files/pdfs/medicines/hiv/yeztugo/yeztugo_pi.

Available Data on LEN Use After Bariatric Surgery

Currently, there are no data available regarding LEN use after bariatric surgery.

There were no reports of participants with a history of bariatric surgery in the PURPOSE 1 and PURPOSE 2 publications.[1,2]

A literature search was conducted in Ovid MEDLINE and Embases databases for studies published between 1946 and March 5, 2026 using search terms that included Yeztugo, lenacapavir, PrEP, pre-exposure prophylaxis, gastric bypass, gastric sleeve surgery, bariatric surgery, sleeve gastrectomy, roux-en-y, absorption and related search terms. No relevant citations were found.

References

1. Bekker LG, Das M, Abdool Karim Q, et al. Twice-Yearly Lenacapavir or Daily F/TAF for HIV Prevention in Cisgender Women. *N Engl J Med.* 2024;391(13):1179-1192.
2. Kelley CF, Acevedo-Quinones M, Agwu AL, et al. Twice-Yearly Lenacapavir for HIV Prevention in Men and Gender-Diverse Persons. *N Engl J Med.* 2025;392(13):1261-1276.

Abbreviations

LEN=lenacapavir

Product Label

For the full indication, important safety information, and boxed warning, please refer to the Yeztugo US Prescribing Information available at: www.gilead.com/-/media/files/pdfs/medicines/hiv/yeztugo/yeztugo_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

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

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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