GILEAD | Medical Information

Yeztugo® (lenacapavir) Intramuscular Administration

This document is in response to your request for information regarding Yeztugo® (lenacapavir [LEN]) and intramuscular (IM) administration.

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.

Product Labeling[1]

LEN injection is only for subcutaneous administration into the abdomen by a healthcare provider. The thigh can be used as an alternative injection site if preferred. Do NOT administer intradermally due to risk of serious injection site reactions.

For complete information regarding the approved dosage and administration of LEN, please refer to Section 2 Dosage and Administration of the Yeztugo U.S. Prescribing Information.

Available Data on LEN IM Administration

Currently, there are no clinical data available regarding LEN and IM administration.

Additionally, a literature search was conducted in Ovid MEDLINE and Embase databases for studies published between 1946 and March 5, 2026 using search terms that included Yeztugo, lenacapavir, PrEP, pre-exposure prophylaxis, intramuscular and related search terms. No relevant citations were found.

References

1. Enclosed, Gilead Sciences Inc. YEZTUGO® (lenacapavir) tablets, for oral use. YEZTUGO® (lenacapavir) injection, for subcutaneous use. U.S. Prescribing Information. Foster City, CA.

Abbreviations

IM=intramuscular
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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