GILEAD | Medical Information

Veklury® (remdesivir) Use for COVID-19 Post-Exposure Prophylaxis

This document is in response to your request for information regarding Veklury® (remdesivir [RDV]) and its use for post-exposure prophylaxis of COVID-19.

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: www.gilead.com/-/media/files/pdfs/medicines/covid-19/veklury/veklury_pi.

Product Labeling[1]

Indications and Usage

RDV is indicated for the treatment COVID-19 in adults and pediatric patients (birth to <18 years of age weighing ≥1.5 kg) who are:

- Hospitalized, or
- Not hospitalized and have mild-to-moderate COVID-19, and are at high risk for progression to severe COVID-19, including hospitalization or death.

Clinical Data

Currently there are no clinical data on the use of RDV for post-exposure prophylaxis of COVID-19.

Additionally a literature search was conducted in Ovid MEDLINE, BIOSIS Previews and Embase databases for studies published between 1946 and December 19, 2024 using search terms that included Veklury remdesivir, GS-441524, GS-704277, post-exposure prophylaxis and related search terms. No published clinical trials in humans were identified.

References

1. Enclosed. Gilead Sciences Inc. VEKLURY® (remdesivir) for injection, for intravenous use. VEKLURY® (remdesivir) injection, for intravenous use. U.S. Prescribing Information. Foster City, CA.

Abbreviations

COVID-19=coronavirus disease 2019

RDV=remdesivir

Product Label

For the full indication, important safety information, and boxed warning(s), please refer to the Veklury US Prescribing Information available at: www.gilead.com/-/media/files/pdfs/medicines/covid-19/veklury/veklury_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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