



# Veklury<sup>®</sup> (remdesivir) Coring During Preparation

This document is in response to your request for information regarding Veklury<sup>®</sup> (remdesivir [RDV]) and coring during preparation.

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](http://www.gilead.com/-/media/files/pdfs/medicines/covid-19/veklury/veklury_pi).**

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## Available Data on Coring During RDV Preparation<sup>1</sup>

Follow your local hospital or pharmacy procedures if coring occurs during preparation.

A filter can be used to remove the cored rubber stopper piece. There are no data on the type of filter that should be used to remove the cored rubber stopper piece.

After reconstitution of RDV, inspect the vial to ensure the container closure is free from defects and the solution is free of particulate matter.

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## Reference

1. Gilead Sciences Inc. Data on File.

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## 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](http://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](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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