



# Veklury<sup>®</sup> (remdesivir)

## Osmolarity of the Final Solution

This document is in response to your request for information regarding the osmolarity of the final solution of Veklury<sup>®</sup> (remdesivir [RDV]).

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).**

## Available Data on the Osmolarity of the Final Solution of RDV<sup>1</sup>

The osmolarity of RDV diluted into 0.9% sodium chloride is presented in Table 1.

**Table 1. Osmolarity of the Final Solution of RDV<sup>1</sup>**

Formulation	RDV Dose	Final Diluted Volume <sup>a</sup>	Calculated Osmolarity <sup>b</sup>
RDV for injection 100 mg lyophilized powder	100 mg	100 mL	343 mOsm/L
		250 mL	324 mOsm/L
	200 mg	100 mL	377 mOsm/L
		250 mL	338 mOsm/L

<sup>a</sup>The final diluted volume is per the recommended preparation instructions for adults.

<sup>b</sup>Calculated from the measured osmolality.

## 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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