

# Epclusa® (sofosbuvir/velpatasvir) Coadministration with Doxorubicin

This document is in response to your request for information regarding Epclusa® (sofosbuvir/velpatasvir [SOF/VEL]) and coadministration with doxorubicin.

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/liver-disease/epclusa\\_pi](http://www.gilead.com/-/media/files/pdfs/medicines/liver-disease/epclusa_pi).**

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## PK DDI Evaluation

Drug interaction studies have not been conducted between the single-tablet regimen SOF/VEL and doxorubicin. Based on the PK profile of each active ingredient within SOF/VEL and doxorubicin, a PK interaction would be predicted.<sup>1,2</sup> VEL is an inhibitor of drug transporter P-gp. P-gp inhibitors may increase the serum concentration of doxorubicin. For more information about doxorubicin, please refer to its product labeling.<sup>2</sup>

### SOF/VEL PK<sup>1</sup>

DDI Mechanism		SOF	VEL
Drug Transporters	P-gp/BCRP	Substrate	Substrate/Inhibitor
	OATP1B1	N/A	Inhibitor
	OATP1B3	N/A	Inhibitor
	OATP2B1	N/A	Inhibitor
Drug Metabolizing Enzymes	CYP1A2	N/A	N/A
	CYP2B6	N/A	Substrate
	CYP2C8	N/A	Substrate
	CYP2C19/19	N/A	N/A
	CYP2D6	N/A	N/A
	CYP3A4	N/A	Substrate

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## Relevant SOF/VEL Label Information<sup>1</sup>

There is no information in the SOF/VEL product labeling about the coadministration of SOF/VEL and doxorubicin.

Clearance of HCV infection with direct acting antivirals may lead to changes in hepatic function, which may impact safe and effective use of concomitant medications. Frequent monitoring of relevant laboratory parameters (INR or blood glucose) and dose adjustments of certain concomitant medications may be necessary. For more information, please refer to

Section 7.3 of the SOF/VEL US Prescribing Information (Established and Potentially Significant Drug Interactions).

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## Available Data

There are no Gilead studies evaluating the coadministration of SOF/VEL and doxorubicin.

Additionally, a literature search was conducted in Ovid MEDLINE and Embase databases for studies published between 1946 and January 14, 2026 using search terms that included Epclusa, velpatasvir, sofosbuvir, doxorubicin, and related search terms. No relevant citations were found.

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## References

1. Enclosed. Gilead Sciences Inc, EPCLUSA® (sofosbuvir and velpatasvir) tablets, for oral use. US Prescribing Information. Foster City, CA.
2. Pfizer Labs, Division of Pfizer Inc, DOXORUBICIN HYDROCHLORIDE injection, for intravenous use. DOXORUBICIN HYDROCHLORIDE for injection, for intravenous use. US Prescribing Information. New York, NY.

## Abbreviations

BCRP=breast cancer resistance protein

DDI=drug-drug interaction

OATP=organic anion transporting polypeptide  
P-gp=P-glycoprotein  
PK=pharmacokinetic(s)

SOF=sofosbuvir  
VEL=velpatasvir

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## Product Label

For the full indication, important safety information, and Boxed Warning(s), please refer to the Epclusa US Prescribing Information available at:

[http://www.gilead.com/-/media/files/pdfs/medicines/liver-disease/epclusa/epclusa\\_pi](http://www.gilead.com/-/media/files/pdfs/medicines/liver-disease/epclusa/epclusa_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 Pharmacovigilance and Epidemiology ☎ 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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