



**Swiss Public Summary of the
Risk Management Plan (RMP)**

for

Vosevi, film-coated tablets

(Sofosbuvir/Velpatasvir/Voxilaprevir Fixed-Dose Combination)
400mg/100mg/100mg

Version 4.0 (October 2021)
Based on EU RMP version 4.0 (October 2020)

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SUMMARY OF RISK MANAGEMENT PLAN FOR VOSEVI (SOFOSBUVIR/VELPATASVIR/VOXILAPREVIR)

The Risk Management Plan (RMP) is a comprehensive document submitted as part of the application dossier for market approval of a medicine. The RMP summary contains information on the medicine's safety profile and explains the measures that are taken in order to further investigate and follow the risks as well as to prevent or minimize them.

The RMP summary of Vosevi is a concise document and does not claim to be exhaustive.

As the RMP is an international document, the summary might differ from the „Arzneimittelinformation / Information sur le médicament“ approved and published in Switzerland, e.g. by mentioning risks occurring in populations or indications not included in the Swiss authorization.

Please note that the reference document which is valid and relevant for the effective and safe use of Vosevi in Switzerland is the „Arzneimittelinformation / Information sur le médicament“ (see www.swissmedic.ch) approved authorized by Swissmedic. Gilead Sciences Switzerland Sàrl is fully responsible for the accuracy and correctness of the content of the here published summary RMP of Vosevi.

I. The Medicine and What is it Used for

Vosevi is authorized for treatment of chronic hepatitis C (CHC) in adults (see SmPC for the full indication). It contains sofosbuvir (SOF), velpatasvir (VEL) and voxilaprevir (VOX) as active substances and it is given orally.

Further information about the evaluation of Vosevi's benefits can be found in Vosevi's EPAR, including in its plain-language summary, available on the EMA website, under the medicine's webpage: <https://www.ema.europa.eu/en/medicines/human/EPAR/vosevi>.

II. Risks Associated with the Medicine and Activities to Minimize or Further Characterize the Risks

Important risks of Vosevi, together with measures to minimize such risks and the proposed studies for learning more about Vosevi's risks, are outlined below.

Measures to minimize the risks identified for medicinal products can be:

- Specific Information, such as warnings, precautions, and advice on correct use, in the package leaflet and SmPC addressed to patients and healthcare professionals;
- Important advice on the medicine's packaging;
- The authorized pack size — the amount of medicine in a pack is chosen so to ensure that the medicine is used correctly;
- The medicine's legal status — the way a medicine is supplied to the public (e.g. with or without prescription) can help to minimize its risks.

Together, these measures constitute routine risk minimization measures.

In addition to these measures, information about adverse reactions is collected continuously and regularly analyzed including PSUR assessment so that immediate action can be taken as necessary. These measures constitute routine pharmacovigilance activities.

If important information that may affect the safe use of Vosevi is not yet available, it is listed under ‘missing information’ below.

II.A. List of important risks and missing information

Important risks of Vosevi are risks that need special risk management activities to further investigate or minimize the risk, so that the medicinal product can be safely administered. Important risks can be regarded as identified or potential. Identified risks are concerns for which there is sufficient proof of a link with the use of Vosevi. Potential risks are concerns for which an association with the use of this medicine is possible based on available data, but this association has not been established yet and needs further evaluation. Missing information refers to information on the safety of the medicinal product that is currently missing and needs to be collected (e.g. on the long-term use of the medicine).

Table Part VI.1. List of Important Risks and Missing Information

Important Identified Risks	Severe bradycardia and heart block when used with concomitant amiodarone
	HBV reactivation in HBV/HCV coinfecting patients
Important Potential Risks	Recurrence of HCC
	Emergence of HCC
Missing Information	Safety in pregnant women
	Safety in patients with previous HCC
	Safety in patients with moderate or severe hepatic impairment

II.B. Summary of Important Risks

Table Part VI 2. Summary of Important Risk(s) and Missing Information

Important Identified Risk	Severe bradycardia and heart block when used with concomitant amiodarone
Evidence for linking the risk to the medicine	Cases of severe bradycardia have been observed when SOF-containing regimens are used in combination with amiodarone.
Risk factors and risk groups	Patients also taking beta blockers, or those with underlying cardiac comorbidities and/or advanced liver disease may be at increased risk for symptomatic bradycardia with coadministration of amiodarone.
Risk Minimization Measure(s)	Routine risk minimization measures: SmPC section 4.4, 4.5, and 4.8

	<p>PL section 2.</p> <p>Additional risk minimization measures: None</p>
Important Identified Risk	HBV reactivation in HBV/HCV coinfecting patients
Evidence for linking the risk to the medicine	Cases of HBV reactivation have been reported in patients coinfecting with HBV/HCV during or after treatment with DAAs. HBV reactivation can potentially be life-threatening, as it could result in hepatitis, an increase in transaminase levels, an increase in bilirubin levels, hepatic failure and death.
Risk factors and risk groups	Due to the small number of cases of HBV reactivation with DAAs, risk factors have not been definitively established. However, some of the cases involving HBV reactivation with SOF-containing regimens involved patients who were immunocompromised (patients coinfecting with HIV or patients receiving immunosuppressants due to prior transplant). In addition, a case involving severe HBV reactivation had risk factors of NASH and Burkitt's lymphoma.
Risk Minimization Measure(s)	<p>Routine risk minimization measures: SmPC Section 4.4 PL Section 2</p> <p>Additional risk minimization measures: None</p>
Important Potential Risk	Recurrence of HCC
Evidence for linking the risk to the medicine	HCC has been reported in some patients who have previously had HCC while taking drugs used to treat hepatitis C virus (direct acting antivirals). It is unclear whether hepatitis C direct acting antivirals increase the risk of HCC returning in patients who previously had HCC and a study is being conducted to investigate this. The risk has not yet been confirmed.
Risk factors and risk groups	Risk factors associated with HCC recurrence include high alpha-fetoprotein (AFP) levels prior to HCC treatment, the size of the primary tumor, and the number of primary tumors. The risk of recurrence will also depend on the method used to treat the primary tumor.
Risk Minimization Measure(s)	<p>No risk minimization measures.</p> <p>The need for risk minimization measures will be reassessed following the availability of the results from a study for HCC recurrence.</p>
Additional Pharmacovigilance activities	<p>Additional pharmacovigilance activities:</p> <p>Study to assess the impact of DAA treatment on the incidence of HCC recurrence, relative to no DAA therapy</p> <p>See Section II.C of this summary for an overview of the post-authorization development plan.</p>

Important Potential Risk	Emergence of HCC
Evidence for linking the risk to the medicine	Some patients have developed HCC while taking drugs used to treat hepatitis C virus (direct acting antivirals). HCC is a known complication of hepatitis C virus especially in the presence of advanced liver disease. It is unclear whether hepatitis C direct acting antivirals increase the risk of developing HCC or not. The risk has not yet been confirmed.
Risk factors and risk groups	The presence of cirrhosis is a primary major risk factor for the development of HCC in CHC patients. Additional risk factors for the development of HCC in CHC patients includes older age, male sex, heavy alcohol use, diabetes, obesity, smoking, and HBV-coinfection {El-Serag 2007, Hiramatsu 2015, Shariff 2009, Yang 2010}. Clinical factors shown to influence the risk of HCC include advanced liver fibrosis, lower platelet count and albumin level; higher levels of alkaline phosphatase and α -fetoprotein; and the presence of esophageal varices. In CHC patients treated with DAAs, the presence of cirrhosis and treatment failure were associated with an increased risk of de novo HCC; treatment with DAAs with or without IFN was not a risk factor for de novo HCC.
Risk Minimization Measure(s)	No risk minimization measures The need for risk minimization measures will be reassessed following the availability of results from an investigation of the impact of DAA therapies on the incidence and type of de novo HCC.
Additional Pharmacovigilance activities	Additional pharmacovigilance activities: Study to evaluate the risk of de novo HCC in patients with compensated cirrhosis treated with DAAs for chronic hepatitis C. See Section II.C of this summary for an overview of the post-authorization development plan.
Missing information	Safety in pregnant women
Risk Minimization Measure(s)	Routine risk minimization measures SmPC section 4.6 PL section 2. Additional risk minimization measures: None
Missing information	Safety in patients with previous HCC
Risk Minimization Measure(s)	No risk minimization measures. The need for risk minimization measures will be reassessed following the availability of the results from a planned study for HCC recurrence.
Additional Pharmacovigilance activities	Additional pharmacovigilance activities: Study to assess the impact of DAA treatment on the incidence of HCC recurrence, relative to no DAA therapy See Section II.C of this summary for an overview of the post-authorization development plan.

Missing information	Safety in patients with moderate or severe hepatic impairment
Risk Minimization Measure(s)	Routine risk minimization measures: SmPC Sections 4.2, 4.4 and 5.2 PL Section 3 Additional risk minimization measures: None

II.C. Post-authorization Development Plan

II.C.1. Studies which are Conditions of the Marketing Authorization

Table Part VI 3. Studies as Condition of the Marketing Authorization

Short Study Name	Purpose of the Study
DAA-PASS: A post-authorisation safety study of early recurrence of hepatocellular carcinoma in HCV-infected patients after direct-acting antiviral therapy	To evaluate the impact of DAA treatment on the incidence of HCC recurrence, relative to no DAA therapy

II.C.2. Other Studies in Post-Authorization Development Plan

Table Part VI 4. Other Studies in Post-Authorization Development Plan

Short Study Name	Purpose of the Study
De Novo DAA PASS: A study to evaluate the risk of de novo hepatocellular carcinoma in patients with compensated cirrhosis treated with direct acting antivirals for chronic hepatitis C	To evaluate among compensated cirrhotic patients, whether DAA therapy for chronic HCV infection increases the risk of incident HCC compared to no treatment or treatment with IFN-based regimens