21 May 2026
EMADOC-1700519818-3140849
Committee for Medicinal Products for Human Use (CHMP)

Summary of opinion[1] (post authorisation)

Maviret

glecaprevir / pibrentasvir

On 21 May 2026, the Committee for Medicinal Products for Human Use (CHMP) adopted a positive opinion recommending a change to the terms of the marketing authorisation for the medicinal product Maviret. The marketing authorisation holder for this medicinal product is Abbvie Deutschland GmbH & Co. KG.

The CHMP adopted an extension to the existing indication as follows:[2]

> Maviret is indicated for the treatment of **acute and** chronic hepatitis C virus (HCV) infection in adults and children aged 3 years and older (see sections 4.2, 4.4. and 5.1).

Detailed recommendations for the use of this product will be described in the updated summary of product characteristics (SmPC), which will be published on the EMA website in all official European Union languages after a decision on this change to the marketing authorisation has been granted by the European Commission.

[1] Summary of positive opinion are published without prejudice to the Commission decision, which will normally be issued 67 days from adoption of the opinion.

[2] New text in bold.

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