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Public statement

Atripla

Withdrawal of the marketing authorisation in the European Union

On 15 November 2021, the European Commission withdrew the marketing authorisation for Atripla (efavirenz / emtricitabine / tenofovir disoproxil) in the European Union (EU). The withdrawal was at the request of the marketing authorisation holder, Gilead Sciences Ireland UC, which notified the European Commission of its decision to permanently discontinue the marketing of the product for commercial reasons.

Atripla was granted marketing authorisation in the EU on 13 December 2007 for treatment of HIV-1 infection. The marketing authorisation was initially valid for a 5-year period. It was then granted unlimited validity in 2012.

The European Public Assessment Report (EPAR) for Atripla will be updated to indicate that the marketing authorisation is no longer valid.

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