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Public statement

Vitekta

Withdrawal of the marketing authorisation in the European Union

On 3 November 2016, the European Commission withdrew the marketing authorisation for Vitekta (elvitegravir) in the European Union (EU). The withdrawal was at the request of the marketing authorisation holder, Gilead Sciences International Ltd, which notified the European Commission of its decision to permanently discontinue the marketing of the product for commercial reasons.

Vitekta was granted marketing authorisation in the EU on 13 November 2013 for treatment of human immunodeficiency virus-1 (HIV-1) infection in adults. The marketing authorisation was initially valid for a 5-year period.

The European Public Assessment Report (EPAR) for Vitekta will be updated to reflect that its marketing authorisation is no longer valid.

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