Emtricitabine/tenofovir disoproxil fumarate 200 mg/300 mg tablets (Sun Pharmaceutical Industries Ltd†), HA551

## **Steps taken following Prequalification of [HA551 trade name]**<sup>1</sup>**:**

Changes	WHOPAR parts affected	Accepted on
Change in the name and/or corporate address of the supplier of the FPP	3, 4, 5	06 June 2015
Change in the name and/or address of a manufacturer of the FPP		
Change in the shelf-life of the FPP involving extension	4	11 November 2016
Change of the FPP manufacturing and testing site	3	10 April 2019
Change in the package size(s) of the FPP	3,4,5	10 April 2019
General update of the WHOPAR to reflect the current WHOPAR structure, product specifics, WHO guidelines and state of scientific knowledge	2, 3, 4	April 2020

<sup>&</sup>lt;sup>1</sup> Trade names are not prequalified by WHO. This is the national medicines regulatory authority's responsibility.

<sup>†</sup> Formerly Ranbaxy Laboratories Limited.