15 December 2022
EMA/794451/2022
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): lamivudine / tenofovir disoproxil

Procedure No. PSUSA/00010751/202203

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Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lamivudin/Tenofovirdisoproxil Cipla 300/245 mg Filmtabletten	DE/H/5651/001	97614.00.00	CIPLA EUROPE NV	DE
Lamivudina/Tenofovir disoproxilo Cipla 300 mg/245 mg comprimidos recubiertos con película.	DE/H/5651/001	83802	CIPLA EUROPE NV	ES