Considerations for Implementation of Cabenuva **Use in Clinical Practice**



Cabenuva is co-packaged into a kit containing cabotegravir extended-release (CAB LA) injectable suspension with rilpivirine extended-release (RPV LA) injectable suspension¹





CAB LA 600 mg/3 mL RPV LA 900 mg/3 mL





CAB LA 400 mg/2 mL RPV LA 600 mg/2 mL

Figure 1. Dosing Kits

Cabenuva, a 2-drug co-packaged product of cabotegravir, an HIV-1 INSTI, and rilpivirine, an HIV-1 NNRTI, is indicated as a complete regimen for the treatment of HIV-1 infection in adults to replace the current antiretroviral regimen in those who are virologically suppressed (HĪV-1 RNA < 50 copies/mL) on a stable antiretroviral regimen with no history of treatment failure and with no known or suspected resistance to either cabotegravir or rilpivirine.

Monthly Dosing Schedule of Cabenuva¹ Every 2 Month Dosing Schedule of Cabenuva¹ Month 2 and 3 Month 1 Month 2 Month ≥ 3 Month 1 OLI* Initiation Continuation OLI* Initiation phase Continuation (minimum 28 days) phase phase (minimum 28 days) **CAB** tablet **CAB LA** CAB LA **CAB** tablet **CAB LA CAB LA** CAB LA Daily oral dose Single IM Q1M IM Daily oral dose Single IM Q2M IM Q2M IM (30 mg) injection injections (30 mg) injection injections injections Previous Previous (600 mg, 3 mL) (600 mg, 3 mL) (600 mg, 3 mL) (600 mg, 3mL) (400 mg, 2 mL) ART ART RPV LA RPV tablet **RPV LA RPV LA RPV** tablet **RPV LA RPV LA** 8 8 Daily oral dose Single IM Q1M IM Daily oral dose Single IM Q2M IM Q2M IM (25 mg) taken injection injections (25 mg) taken injection injections injections (600 mg, 2mL) (900 mg, 3mL) with a meal (900 mg, 3mL) with a meal (900 mg, 3mL) (900 mg, 3mL) Initiate injections on Both IM injections Initiate injections on *OLI (oral lead-in) *OLI (oral lead-in) given once a month, Second injection is the final day of the the final day of the All subsequent is optional and is optional and OLI (if used) or the at separate gluteal OLI (if used) or the given one month injections are given may be used to may be used to after the first patient may go injections sites patient may go every two months assess tolerability assess tolerability directly to injections during same visit directly to injections

Overview of the Implementation Journey of *Cabenuva* in Clinical Practice¹⁻²

Preparation



- 1. Practice preparation
- 2. Patient selection

Acquisition



- 3. Benefits Investigation
- & Product Acquisition
- 4. Inventory Management

Initiation



- 5. Oral Lead-in
- 6. Injection visits

Continuation



- 7. Patient Adherence
- 8. Alternate Site for Administration

Click here

Month ≥ 5

phase

To access the Cabenuva Guidance **Document for** Implementation

Considerations for Clinic Preparation³

CUSTOMIZE was an implementation-effectiveness study to inform clinical practice around implementation of Cabenuva. Some of this information may be useful for varying clinic types. Additional information on CUSTOMIZE:

For Office Staff Perspectives, please click



For Patient Perspectives, please click





Injection education programs are available through ViiV Healthcare by contacting a regional sales representative..





For higher BMI individuals (e.g. > 30 kg/m²), consider a 2-inch needle (not included in kit). For more details, please click the hand icon.



Important safety information is found in the attached Prescribing Information.

For more information



MI Letter









CLICK FOR VIIV US Medical Portal

Some information contained in this response may not be included in the approved Prescribing information. This response is not intended to offer recommendations for administering this product in a manner inconsistent with its approved labeling. Please note that reports of adverse events in the published literature often lack causality assessments and may contain incomplete information; therefore, conclusions about causality generally cannot be drawn. In order for ViiV Healthcare to monitor the safety of our products, we encourage healthcare professionals to report adverse events or suspected overdoses to the company at 877-844-8872. Please consult the attached Prescribing Information. This response was developed according to the principles of evidence-based medicine and, therefore, references may not be all-inclusive.

Abbreviations: ART = antiretroviral therapy; CAB = cabotegravir tablets; CAB LA = long-acting cabotegravir; cu ft = cubic-foot; HCP = health care professional; HIV-1 = human immunodeficiency virus type 1; IM = intramuscular; INSTI = integrase strand transfer inhibitor; NNRTI = non-nucleoside reverse transcriptase inhibitor; RPV = rilpivirine tablets; RPV LA = long-acting rilpivirine; RNA = ribonucleic acid; OLI = oral lead-in; Q1M = once-monthly; Q2M = every 2 month.

References: 1. ViiV Healthcare Local Label; 2. Data on File. 2019N425811; 3. Data on File. CUSTOMIZE (Study 209493). Available at http://www.viiv-studyregister.com