Indications	Patients with HIV-1 infection who have had an undetectable (<50 copies/mL) viral load, on antiretroviral therapy Required duration of prior virologic suppression not defined, but was 1-6 months in registrational trials
Contraindications	Hepatitis B co-infection [requires a regimen containing tenofovir] Known resistance to either cabotegravir or rilpivirine Pregnancy or planning pregnancy [not studied in this population]
Drug Interactions	Contraindicated: - anticonvulsants [e.g. carbamazepine, oxcarbazepine, phenytoin, phenobarbital] - anti-mycobacterials [e.g. rifampin, rifapentine, rifabutin] - glucocorticoids [e.g. dexamethasone] - macrolides (except for azithromycin) - St. John's Wort Caution when co-administered with: - methadone - methotrexate - QT prolonging medications
Dosing and Administration	 Oral lead-in (optional): 30mg cabotegravir + 25mg rilpivirine once daily x 1 month, followed by: Intra-gluteal, intra-muscular (IM) injections: 600mg cabotegravir IM + 900mg rilpivirine IM x 1, followed by: 400mg cabotegravir IM + 600mg rilpivirine IM every 1 month OR 600mg cabotegravir IM + 900mg rilpivirine IM at 1 month, and then 600mg cabotegravir IM + 900mg rilpivirine IM every 2 months Patients may receive cabotegravir/rilpivirine up to 7 days before- or after- the scheduled date
Known Risk Factors for Virologic Failure despite on-time injections	BMI >30 kg/m2 HIV subtype A6/A1 Pro-viral rilpivirine resistance-associated mutations

Appendix 1: Long-acting, injectable cabotegravir/rilpivirine for treatment of HIV-1 infection in adults(1,2)

Appendix 1, as supplied by the authors. Appendix to: Halani S, Tan D, Andany N. Long-acting injectable antiretroviral therapy for HIV-1 infection in adults. *CMAJ* 2024. doi: 10.1503/cmaj.231498. Copyright © 2024 The Author(s) or their employer(s). To receive this resource in an accessible format, please contact us at cmajgroup@cmaj.ca.

Adverse Events	Injection site reactions (ISRs) - median duration of ISRs is 3 days. - mostly mild and reduce in frequency over time
	Less common adverse events (<2%): - injection-associated reactions (pyrexia, headache, musculoskeletal pain, nausea, dizziness, rash) - gastrointestinal disorders - hepatotoxicity - psychiatric disorders - hypersensitivity

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