

Operationalizing Use of Long-Acting Injectable ART

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Disclosures

No conflict of interest or relationships to disclose

Summary of Data from ATLAS and FLAIR Studies

- CAB/RPV IM q 4 weeks was non-inferior to oral options studied
- Injection site reactions not uncommon
- Satisfaction was high (97% and 99% preferred IM to PO)
- Low rates of virologic failure (3 in each study)
- When virologic failure occurred, NNRTI and INSTI resistance emerged (no cross resistance to DTG seen)

Case

- 35 year old woman is interested in LA-ART. She is tired of taking pills daily.

Your patient if interested in CAB/RPV

- Eligibility Criteria
 - Patient interest
 - Understands monthly visits for 2 injections each time
 - Understands that there is a 4-week oral lead-in
 - Has been suppressed for at least 3 months (although 6 months was used in the studies)

Your patient if interested in CAB/RPV

- Ineligibility criteria
 - Resistance (okay to have a K103N only)
 - HBV positive (if not planning to take oral HBV active agent)
 - Clade A1 or L74I polymorphism
 - h/o poor adherence (until LATITUDE trial data available)
 - Pregnancy or planning to become pregnant

Your patient meets criteria

- Place referral to pharmacy for CAB/RPV (Cabenuva)
- Pharmacy and provider team will clinically review
- PC (program coordinator) will obtain insurance approval (PA)
- Once approved, pharmacy will schedule in person visit to review
 - Prescribes the oral lead in from Viiv
 - Schedules visit for loading injections and subsequent first maintenance injections (pharmacy and nursing visits)

Referral to Pharmacy

- Reason for referral
- Therapy/Medication(s) to Review (Select all as default – can choose individual as needed):
 - Oral Lead-in:
 - Cabotegravir 30mg – Take 1 tablet daily #30 tablets
 - Rilpivirine 25mg – Take 1 tablet daily #30 tablets
 - Initiation Injection: 600-mg/900-mg kit:
 - 600-mg single-dose vial of cabotegravir IM x 1 dose +
 - 900-mg single-dose vial of rilpivirine IM x 1 dose
 - Continuation Injections: 400-mg/600-mg kit:
 - 400-mg single-dose vial of cabotegravir IM x 1 dose monthly +
 - 600-mg single-dose vial of rilpivirine IM x 1 dose monthly
- Most recent HIV RNA undetectable and within the last 3 months
- HIV genotype documented
- HBV status
- Pregnancy status

Your patient meets criteria

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Ensuring adherence with maintenance dosing

- RN champion to track patients
- What happens if patients missed doses
 - Bridge dosing for planned missed injections
 - If unplanned, has 7-day grace period to get f/u injection
 - If unplanned and more than 7 days after missed injections, then need oral bridge (which will need to be ordered)
 - Patients may need reloading

Considerations for Ineligibility

- Missed injections and requiring more than 2 additional loading doses in a 6-month period
- Pregnancy or desire to become pregnant

Status of LA-ART at Madison

- We have yet to start anyone
- Lots of provider interest, some patient interest

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