RAPID ART START PROTOCOL

Evidence-Informed for the Linking and Retention in HIV Care Chapter Evidence-Informed for the Structural Interventions Chapter



POPULATION

> Patients with HIV at a Veterans' health clinic

KEY INTERVENTION EFFECTS

- Decreased time to engage in HIV care
- Decreased time to ART initiation
- Decreased time to viral suppression

BRIEF DESCRIPTION

Rapid Antiretroviral Therapy (ART) Start Protocol streamlines HIV treatment for patients who have a new HIV diagnosis. The protocol workflow is as follows:

- The Infectious Disease Clinic receives notification, confirms HIV diagnosis, and contacts the patient within 72 hours (preferably on the same day of diagnosis).
- During the first visit, a multidisciplinary team (i.e., nurse, scheduler, medical provider, pharmacist, psychologist, and social worker) provides care.
 - The provider performs initial assessment, opportunistic infection screening, HIV education, sexually transmitted infection screening, and counseling (e.g., prophylaxis for partners).
 - Initial lab tests are conducted, and ART is prescribed.
 - Social worker assists patient with partner notification and assesses potential barriers to care with linkage to further resources.
 - Psychologist addresses mental health and substance use concerns.
- At day 14, pharmacist and social worker connect with client via telephone and provide side-effect management, adherence education, and counseling.
- Follow-up visit occurs within 4-6 weeks with provider to address adherence, medication interactions, and comorbidities; lab tests are conducted and results are discussed with patient. Further follow-up lab tests and visits scheduled around 6-8 weeks later.

DURATION: at least 4 sessions (3 in person and 1 telephone) over the course of 10-14 weeks
 SETTING: Veteran's Health Administration Infectious Disease clinic (Atlanta, GA)
 STUDY YEARS: 2012 – 2020
 STUDY DESIGN: Retrospective cohort design
 DELIVERERS: Multidisciplinary team of clinical care staff (nurses, schedulers, medical providers, pharmacists, psychologists, and social workers)
 DELIVERY METHODS: Appointment scheduling, Case management, In-person visits, Phone calls

STUDY SAMPLE

The baseline study sample of 116 patients was characterized by the following:

- 85% Black or African American persons
- 15% White persons
- 1% persons identifying as Hispanic, Latino or Latina, regardless of race
- 76% male persons, 5% female persons
- Median age = 44 years

STRUCTURAL COMPONENTS

- Access HIV care
 - Expedited access to HIV care and ART prescription
- Institutional Policy/Procedure Institutional Procedure
 - $_{\odot}$ Changed clinical procedures to increase access to care and HIV treatment

KEY INTERVENTION EFFECTS (see Primary Study for all outcomes)

- The median (Interquartile interval, IQI) time from referral to first attended clinic appointment was reduced from 20 days (10-43) pre-intervention to 1 day (0-3) post-intervention (p < 0.001).
- The median (IQI) time from first attended visit to ART initiation (measured as the ART dispense date) decreased from 27.5 days (3-50) pre-intervention to 0 days (0-0) post-intervention (p = 0.01).
- The median (IQI) time to viral suppression from diagnosis decreased from 180.5 days (102.5-338.5) preintervention to 62 days (40-105) post-intervention (p < 0.001).
- Patients who received Rapid ART Start were more likely to achieve viral suppression at any given time during the study period compared to pre-intervention participants (Hazard Ratio = 2.65, 95% Confidence Interval: 1.69 - 4.16, p < 0.001).

CONSIDERATIONS

- Mortality: More deaths were seen pre-intervention (n = 6), compared to post intervention (n = 2) groups.
- Fidelity measures of both pre-intervention and post-intervention groups were very high with 100% of patients having a first appointment visit with a subsequent follow-up visit, and 95% of patients were retained in care.
- Resources that may be needed to implement rapid ART programs include: 1) dedicated point of contact for
 efficient and reliable notification of new diagnosis of HIV; 2) peers or navigators to assist through the clinic
 and rapid ART process; 3) training of staff to assist with pharmaceutical assistance program applications; and
 4) a multidisciplinary team that includes a social worker, eligibility/insurance specialist, and a dedicated
 medical provider.

ADVERSE EVENTS

The author did not report adverse events.

FUNDING

• Emory c-FAR–Emory Center for AIDS Research (P30 AI050409)

PRIMARY STUDY

O'Shea, J. G., Gallini, J. W., Cui, X., Moanna, A., & Marconi, V. C. (2022). <u>Rapid Antiretroviral Therapy</u> <u>Program: Development and evaluation at a Veterans Affairs Medical Center in the southern United States</u>. *AIDS Patient Care and STDs*, *36*(6), 219–225. doi.org/10.1089/apc.2022.0039

PLEASE CONTACT STUDY AUTHOR FOR TRAINING AND INTERVENTION MATERIALS.

Contact information

Jesse G. O'Shea, MD, MSc Division of Infectious Diseases Emory University School of Medicine 49 Jesse Hill Jr. Drive Atlanta, GA 30303 Email: jesseosheamd@gmail.com