

# THE ANTIRETROVIRAL PREGNANCY REGISTRY

## The Purpose of the Antiretroviral Pregnancy Registry (APR)

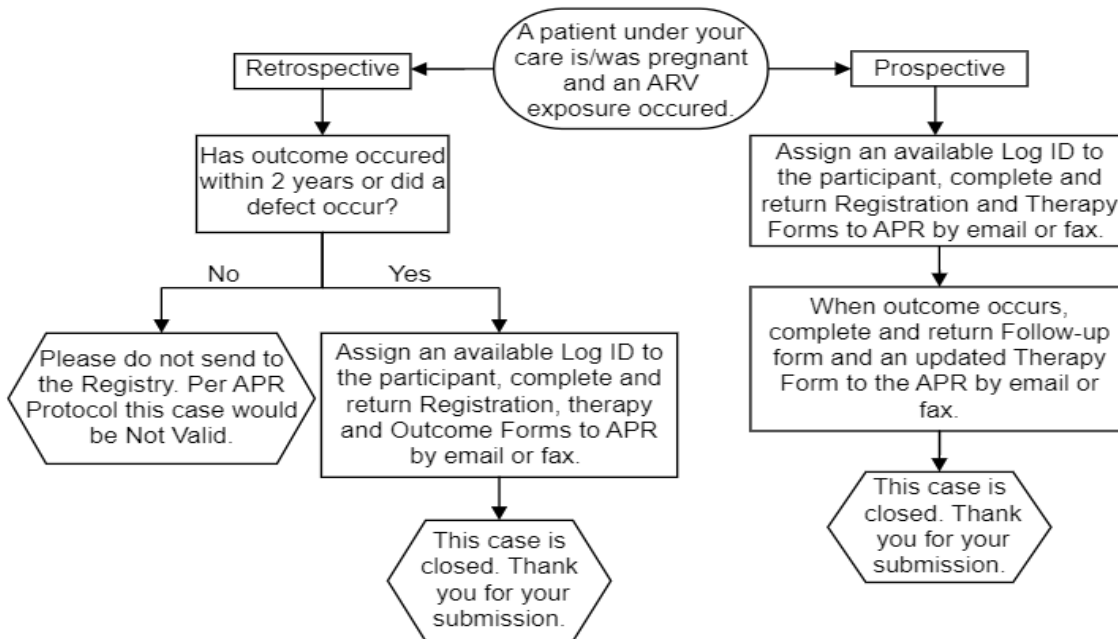
A voluntary prospective, exposure-registration, observational study designed to collect and evaluate data on the outcomes of pregnancies exposed to antiretroviral (ARV) products. The APR is intended to provide an early signal of any major teratogenic effect associated with a prenatal exposure to the products monitored through the Registry.

## Health Care Providers with Access to Medical Records can Report

Health Care Providers that can report to APR include: physician, pharmacist, physician assistant, nurse practitioner, registered nurse, medical assistant, etc., as long as you have access to ARV exposure and pregnancy outcome information.

To register, contact the APR by email or phone to provide basic contact information. You will then receive an HCP ID and Log IDs. The Log ID system keeps your patient's identity anonymous thus eliminating the need for informed consent.

## Enrolling your Patient in the Registry



## A Message From Dr. William R. Short, MD, MPH, FIDSA

### Associate Professor of Medicine and Obstetrics/Gynecology

“Often, pregnant persons are excluded from clinical trials; therefore, it is critical that we collect prospective data on new antiretroviral (ARV) agents as they are FDA approved. The Antiretroviral Pregnancy Registry (APR) plays a critical role in filling this gap but is dependent on providers, like you, to report exposures. Do your part and contribute to evaluating ARV safety by reporting all exposures to the APR.”



## Quick Links

- [Antiretroviral Pregnancy Registry \(APR\): ClinicalTrials.gov](#)
- [Antiretroviral Pregnancy Registry NIH \(hiv.gov\)](#)
- [2022 Safety Surveillance through Antiretroviral Pregnancy Registry Webinar](#)
- [31Jul2022 APR Interim Report](#)

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