Call Toll-free 1-800 856-7842 or 828-254-0881 with Extension listed

Name of Study: HCV-TARGET- Hepatitis C Therapeutic Registry and Research Network (UF-UNCCH)

LINK: https://clinicaltrials.gov/ct2/show/NCT01474811?term=Target+HCV&recr=Open&rank=1

PI: William R. Harlan, III, MD

Study Coordinator: Kylee Diaz x **3426** or Direct-828-350-3669

Enrollment Begins: 2012

Enrollment Ends: Approximately Dec 2016

Summary: The primary purpose of the **HCV-TARGET** study is to establish a nationwide registry of patients undergoing treatment with antiviral therapies for chronic **hepatitis C** (**HCV**) at both academic and community practices.

Inclusion:

- All adult patients (age 18 or older) being treated with antiviral HCV treatment regimens treated with triple therapy, including newer DAA's

Exclusion:

- Inability to provide written informed consent.
- Currently participating in another clinical trial of hepatitis C therapeutics. Studies comparing HCV RNA assays are not considered exclusionary.