This document provides an overview of the data included on UF StudyConnect and explains how studies are identified for inclusion. The UF CTSI maintains UF StudyConnect in collaboration with the UF IRBs, UF Health and research teams across the university to provide a comprehensive, searchable listing of actively enrolling human-subject research studies led by UF investigators. In addition to being displayed on UF StudyConnect, study listings appear on UFHealth.org.

UF StudyConnect: [http://studyconnect.ctsi.ufl.edu](http://studyconnect.ctsi.ufl.edu) • UFHealth.org Research Studies & Clinical Trials: [http://ufhealth.org/research-studies-clinical-trials](http://ufhealth.org/research-studies-clinical-trials)

### How are studies identified for inclusion on UF StudyConnect?

1. **IRB data collection by the CTSI Study Registry team.**
   - The team uses three criteria from IRB records to identify studies for possible inclusion: study received full-board or expedited review; end date for enrollment has not passed or was not specified; study is listed as active.

2. **Submission of IRB-approved studies by research teams.**

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**What is this?**
- IRB-approved study title
  - Source: IRB database, intro questionnaire or informed consent/protocol

**What is this?**
- Study description for lay audience
  - Source: Study Registry team (drafted based on IRB informed consent/protocol)

**What is this?**
- Keywords for lay audience
  - Source: Study Registry team

**What is this?**
- Name of principal investigator (PI)
  - Source: IRB intro questionnaire or informed consent/protocol

**What is this?**
- PI department
  - Source: IRB database, intro questionnaire or informed consent/protocol

**What is this?**
- PI title, phone and email
  - Source: UF Directory