



**Topic:** PowerTrials - Clinical Trial Identification in PowerChart

**Facility:** IU Health Facilities

**Audience:** Clinical Staff

**What:** Patients enrolled in research trials are identified on the Demographics Bar in PowerChart. Additional research study information can be accessed on the Clinical Research page.

**Why:** Facilitates patient care by providing information to clinicians about the progress of the patient through the clinical trial

## Clinical Trial Information

The Clinical Trial indicator contains information about the patient’s current participation status in clinical research trials. It is located in the center of the demographics banner bar as shown below. The details are being expanded to include the current research treatment status for interventional trials. The 5 possible statuses are defined below:

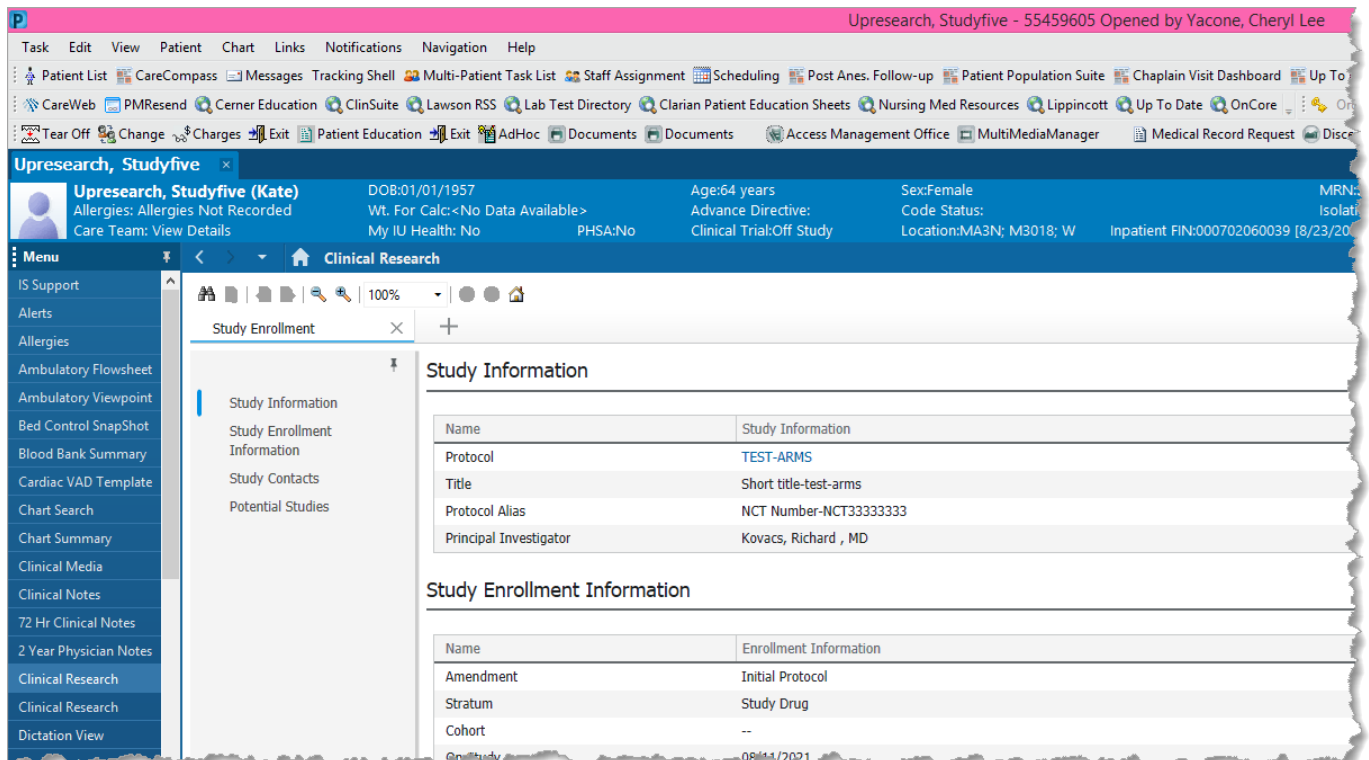
- **Clinical Trial: <No Data Available** Never shown in a clinical trial in PowerTrials
- **Clinical Trial: On Study** Currently enrolled (on study), on treatment, or off treatment in a clinical trial
- **Clinical Trial: Off Study** Currently off study for all clinical trials in PowerTrials

Additional information including study dates, protocol number, study contact information and details about arm enrollment if applicable may be found in the Clinical Research page available on the TOC for most positions.

**Step**

**Action**

**1** Click the **Clinical Research** band on the chart’s Table of Contents (TOC).



The **Clinical Research** page displays and the **Clinical Trial/Study Enrollment History for Patient** is displayed; a list of protocols that the patient is or has been enrolled in.

Continued on next page



Step

Action

1  
continued

Study coordinator contact information is displayed in the **Study Contacts** section.

The screenshot displays the PowerChart interface for a patient named Upresearch, Studyfive (Kate). The patient's demographic information is shown at the top: DOB: 01/01/1957, Age: 64 years, Sex: Female, and MRN: [redacted]. The interface includes a navigation menu on the left with options like 'IS Support', 'Alerts', and 'Clinical Research'. The main content area is titled 'Study Enrollment' and contains a 'Study Contacts' section. This section lists the contact name 'Yacone, Cheryl Lee' with the role 'PowerTrials Study Contact'. Below this, there is a 'Potential Studies' section with a message: 'Patient interest has not been documented. Patient is automatically included in prescreening.' There are also radio buttons for 'Interested' and 'Not Interested', and a 'View Activity' link.

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## Access Clinical Notes

### Research Recruitment Clinical Note

Per IU Health *Clinical Research Recruitment Policy*, when recruiting IU Health patients as potential human research subjects a clinical note will be filed within the **Research/Clinical Trials Records** folder of the patient’s electronic medical record (EMR) and will include the following information-

- Full Study Title
- Sponsor
- Sponsor Protocol Number
- NCT number (if applicable)\*
- Investigational Review Board (IRB) Number
- Principal Investigator’s Name

\*Additional information about the study can be obtained about protocols with NCT numbers by accessing the *Clinical Trials.gov* website at <https://clinicaltrials.gov>

This note can be found under the Progress Notes folder in a sub-folder called Research/Clinical Trials Records.

Upresearch, Studyfive - 55459605 Opened by Yacone, Cheryl Lee

Task Edit View Patient Chart Links Notifications Index Documents Help

Upresearch, Studyfive (Kate) DOB:01/01/1957 Age:64 years Sex:Female  
 Allergies: Allergies Not Recorded Wt. For Calc:<No Data Available> Advance Directive:  
 Care Team: View Details My IU Health: No PHSA:No Clinical Trial:Off Study Location:MA3N; M3018; W Inpatient FIN:00

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Document Type: Research/Clinical Trial Records  
 Document Subject: Test Study Enrollment Note PI Dr. Kovacs  
 Performed By: Buckley, Katherine E, RN on September 07, 2021 11:59 EDT  
 Verified By: Buckley, Katherine E, RN on September 07, 2021 11:59 EDT  
 Encounter Info: 000702060039, IUH Methodist Hosp, Inpatient, 08/23/2021 -

**\* Final Report \***

Test Study  
 Roche (ABC-123)  
 NCT Number: NCT111111111  
 IRB: 111111111111111111  
 PI: Dr. Kovacs

A thorough screening of the patient's chart was completed and patient meets inclusion and has no exclusions per proto Patient and spouse given information about the study treatment and requirements of the protocol as well as information to the HIPAA Authorization. All aspects of the study including study procedures, benefits, risks, confidentiality issues and alternatives to study participation were explained to patient. Patient is aware that participation is voluntary and he/she withdraw from study at any time. The patient was provided adequate time and opportunity to review the consent and