## 1. Purpose:

1.1. To describe the process for requesting and scheduling Clinical Research Center (CRC) support.

# 2. Scope:

2.1. This procedure applies to clinical research protocols for which CRC support is requested/provided.

### 3. Procedures:

- 3.1. CRC support is initiated after study team has received all regulatory and administrative approval.
- 3.2. CRC provides support to investigators for nursing as well space need.
- 3.3. Requests for CRC Support and Scheduling:
  - 3.3.1. A request for CRC support is sent via email at <u>crc-support@bcm.edu</u> at least two weeks before the planned visit date.
  - 3.3.2. Request includes the following information:
    - 3.3.2.1. Study Number/Name
    - 3.3.2.2. Subject Demographic (Patient name, Date of Birth/MRN, Race and ethnicity, Address, Phone number)
    - 3.3.2.3. Visit Name
    - 3.3.2.4. Length of visit
    - 3.3.2.5. Description of nursing procedure
  - 3.3.3. Request includes the following documents as applicable:
    - o <u>Subject Registration form</u>
    - Physician order
    - Signed consent form
    - Signed eligibility checklist (not applicable for screening visit)
    - Any other documents as applicable for the visit.



- 3.4. Based on availability of nursing personnel and space, priority may be given to studies with high acuity subject population.
- 3.5. CRC nurse manager sends confirmation email to study team along with assign CRC nurse for that visit as applicable.
- 3.6. All CRC visits are registered in the Electronic Medical Record (EMR) and shared CRC outlook calendar.
- 3.7. Study team may get read only access to CRC outlook calendar to view visit and study assignment.
- 3.8. Any changes in scheduling or cancellations is communicated to CRC nurse manager as soon as possible.
- 3.9. Study team requesting CRC space are responsible for cleaning CRC room after the visit.

## 4. MATERIALS AND REFERENCES:

4.1. Subject Registration form

## 5. VERSION/REVISION HISTORY:

Approval Date	Version	Revision Summary
10/10/22	1	