



# DIVISION SCOPE OF SERVICE

<b>Division: SAN ANTONIO</b>
<b>DHP Classification: RESEARCH COORDINATOR/ASSISTANT-(NON-LICENSED)</b>
<b>Name of Dependent Healthcare Professional (DHP):</b>

<p><b>Research Coordinator/Assistant-(Non-Licensed):</b>          The Research Coordinator/Assistant-(Non-Licensed) must have equivalent qualifications, competence and function in the same role as employed individuals performing the same or similar services at the facility, as defined by facility job description.</p>
<p><b>Definition of Care or Service:</b>          The Research Coordinator/Assistant-(Non-Licensed) practices under the direction and supervision of the physician (who is involved as the primary or sub-investigator in a clinical research trial, will perform delegated tasks as ordered by the investigator to provide certain aspects of care to patients participating in clinical research trials).          Scope of Service may include:</p> <ul style="list-style-type: none"> <li>• Demonstrates knowledge and skills to appropriately and effectively communicate and interact with patients, families, and visitors of all age groups maintaining cultural sensitivity and confidentiality.</li> <li>• Works in accordance with facility policies and procedures, state and federal regulations and laws.</li> <li>• Demonstrates Clinical and Service excellence behaviors to include code of HCA conduct core fundamentals in daily interactions with patients, families, co-workers and physicians.</li> </ul>
<p><b>Setting(s):</b></p> <ul style="list-style-type: none"> <li>• Healthcare facilities patient care areas</li> </ul>
<p><b>Supervision:</b></p> <ul style="list-style-type: none"> <li>• Direct/indirect supervision by Facility Medical Staff physician</li> <li>• Indirect supervision by department director, site manager or designee</li> <li>• The Research Assistant will secure a sponsoring/ supervising physician form for each physician they wish to provide.</li> </ul> <p><b>Evaluator:</b> Sponsoring physician/ supervising physician in conjunction with department director or designee</p>
<p><b>Qualifications:</b></p> <ul style="list-style-type: none"> <li>• Highest level of education verification required</li> <li>• American Heart Association health care provider BLS Certification</li> </ul>
<p><b>State Requirements:</b></p> <ul style="list-style-type: none"> <li>•</li> </ul>
<p><b>Experience:</b>          PREFERRED: One year experience as a Research Assistant</p>
<p><b>Competencies:</b>          The Research Coordinator/Assistant-(Non-Licensed) will demonstrate:</p> <ul style="list-style-type: none"> <li>• Collaboratively works with physicians and hospital staff to foster optimal patient outcomes</li> <li>• Serves as a resource on protocol issues</li> <li>• Perform, as needed, physician and staff in-services concerning the research study</li> <li>• Accurately obtains the physician requested patient medical records for rounds using at least two patient identifiers:             <ul style="list-style-type: none"> <li>○ Utilizes physician documentation, such as a rounding list, to confirm patient and their locations with the floor unit clerk or nurse</li> <li>○ Pulls and verifies correct patient charts</li> <li>○ Follows HIPAA policies</li> </ul> </li> <li>• Possesses an advanced working knowledge of fundamentals to include:             <ul style="list-style-type: none"> <li>○ Following universal precautions</li> <li>○ Following MHS hazardous materials management guidelines found on MHS Central (Intranet)</li> </ul> </li> </ul>



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- Follows MHS guidelines for hand hygiene
- Effectively communicates any necessary information to physicians and team members
- Appropriately handle, pick up, or deliver medications as directed by the research physician to the hospital pharmacy
- Conducts effective patient teaching as directed by the research physician.
- Safely handles any patient laboratory specimens and/or supplies.
- Reviews records of research patients
- Monitors research protocols
- Assures timely completion of all study activity as per research physician request
- Conducts patient intake to include:
  - Patient history collection
  - Phlebotomy
  - Obtaining an ECG
  - Obtaining vital signs
  - Phone calls related to the research patient
  - Answering questions related to the clinical trial
  - Obtaining informed consent for the clinical trial
- Infection Prevention
  - Practices consistent hand hygiene
  - Uses personal protective equipment (PPE)
  - Required immunizations per DHP Division requirements
  - Complies with Isolation precautions

**References:**

References: Cairns, Carol. (2007). Solving the AHP Conundrum: How to Comply with HR Standards Related to Non-privileged Practitioners (Appendix C). HCPPro, Inc.: Marblehead, MA.

**DHP Printed Name:** \_\_\_\_\_ **DHP Signature:** \_\_\_\_\_

**Company/Vendor:** \_\_\_\_\_ **Date:** \_\_\_\_\_