JANE SMITH, RN

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HEALTH CARE PROFESSIONAL

AREAS OF EMPHASIS - CLINICAL RESEARCH COORDINATION & NURSE MANAGEMENT

kegi terd. Nurse with more than 12 years of experience looking to transition into a full time Clinical Research Coordination role. Seeking to leverage 7 years of experience in Clinical Research and 2 ½ years in managen en and teaching. Well-developed skills collaborating with research organizations and managing studies from illitiation brough close-out. Dedicated leader with outstanding communication skills and a proven track record of success a takiting patients for studies, enrolling patients, collecting medical data, performing patient education, and entering information into electronic databases.

Areas of Expertise

Clinical I esearch Coordination | In-Service Coordination | Nurse Management Staff Development & Training | FDA Regulations & Compliance | Quality Improvements

EXPERIENCE

ABC Company, Dallas, TX

January 2002 - Present

A corporate health care facility specializing in the treatment of a variety of immune conditions.

Special Projects Coordinator/Registered Nurs (Nay 2011 – Present)

- Spearheaded the launch of a new line of business to ouy, hill, and dispense a specialty infusion drug for use by patients at home.
- Secured contracts and agreements with Pharmacy Benefi Managers for major insurance companies.
- Located medical and pharmaceutical suppliers as well as sof ware distributors.
- Purchased medical supplies for infusion pumps and other ancill xy equipment.
- Created and wrote reports to track progress.
- Develop decision trees, flow charts, and procedures from the ground up.
- Managed all purchasing of the specialty drug and coordinating necessary equipment.
- Trained and educated patients and clinical staff regarding sub-cutaneous if fusion societies.

Key Accomplishments

- Instrumental in fulfilling pharmacy requirements to obtain a Physician Dispensity Pharmacy License.
- Contacted Pharmacy Benefit Managers for major insurance companies for contracts an agreements including Prime Therapeutics, Medco, and Medicare Durable Medical Equipment (DME).
- Located and met with Pharmacy Software Distributors including McKesson, QS/1, and Healix

Senior Nurse Manager (January 2009 – May 2011)

- Managed and oversaw the daily activity and scheduling of up to 16 Registered Nurses and Medical Assistant for 10 different clinic locations.
- Hired, trained, and mentored staff including training on products, drugs, procedures, services and equipment.
- Routinely met with pharmaceutical sales representatives, medical science liaisons, and other industry vendors.
- Responsible for the procurement of all medical supplies and drugs.
- Performed patient education and supported clinicians on the floor.
- Developed, updated, and revised clinical documents such as Medical History Questionnaire, Recurrent Infection Questionnaires, and Outpatient Lab and Diagnostic Order forms; these patient forms were uploaded to the business website and are still used today.

December 2000

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Key Accomplishments

- Developed and created a Nursing Procedure book for the clinic that listed different processes for each procedure performed at the clinic.
- A sted and implemented a new employee orientation program that included procedures for Fire, Tornado, Inclement Weather, Safety, and other nursing guidelines.
- acted educational seminars and in-services for medical staff.
- Developed and implemented the use of Excel spreadsheets for the scheduling of all medical staff at 10 clinics with a positimately 15 staff members.
- Created guidelines for scheduling testing and procedures for the front desk staff with different spreadsheets to track inflation dues, insurance coverage and other relevant data.
- Developed and implemented a blood borne pathogens exposure control plan.

Senior Clinical Research Coordinator (January 2002 – January 2009)

- Coordinated and conjucted multiple clinical trials for asthma, allergy and immunology.
- Recruited patients for todie, enrolled patients, performed patient education, collected medical data, and entered information into a software program for Electronic Data Capture (EDC).
- Conducted and managed Phase III are Phase IV studies from initiation thru close-out working directly with Contract Research Associates carsite
- Attended investigator meetings and net with contract research organizations, pharmaceutical sponsors, and institutional review boards.
- Gathered and submitted adverse events and stripes adverse events data.
- Conducted studies in accordance to study guide he; completed case report forms.
- Developed industry contacts for leads on new tribs
- Tracked site payments and studied revenues.
- Developed and wrote quarterly reports.

Key Accomplishments

- Selected to serve on a Pharmaceutical Advisory Board for Res are coordinators out of 712 sites.
- Selected to speak at an Investigator Meeting on Patient Retention Stretegies with an audience of approximately 300 that included physician, pharmaceutical leaders, and other heatileare professionals.
- Represented the clinic during a Food and Drug Administration 3-day audit; clinic passed audit with 100% success and no fines.
- Instrumental in increasing clinical research revenues by 50% from 2002 to 200

ABC Company, Arlington, TX

Staff Registered Nurse

- Assisted surgeons with in-office procedures.
- Scheduled procedures at hospitals and worked with other healthcare departments and professional request medical records/history prior to surgery.
- Performed a variety of nursing procedures such as removing staples and assisting with biopses.

EDUCATION & PROFESSIONAL LICENSE

University of Nebraska Medical Center, Omaha, NE

Bachelor of Science Degree in Nursing