Robert Vanderpool

RN
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#readytowork

Willing to relocate to: Florida - -
Authorized to work in the US for any employer

Work Experience

Research Nurse
System One - Lexington, KY
October 2020 to January 2021

Contract research Registered Nurse for clinical trial

Pine Meadows Acute Care
Pinnacle Health Care - Lexington, KY
August 2019 to October 2019

Work as a RN in a 110 bed nursing home facility. Responsible for 20 + Residents daily to ensure proper care is delivered.

RN- Unit Manager
Signature Healthcare at Heritage Hall - Lawrenceburg, KY
May 2017 to July 2019

Work as a unit manager in a 110 bed nursing home facility. Manage approximately 12 nurses and 25 Nursing assistants. Assist DON with all management of the facility.

Senior CRA
Chiltern US
February 2015 to May 2016

Responsibilities include management of study sites, conduct pre-investigational site visits, study initiation visits, routine monitoring visits and study close-out visits in accordance with corporate, sponsor, FDA and ICH guidelines. Provide site orientation and guidance to assure successful performance of study protocol; review source documents and case report forms in compliance with stated guidelines; assure completion, submission and maintenance of regulatory documents. Prepare and timely submit accurate monitoring reports and manage regional home office.

Therapeutic Area/ Indication/ Lab Experience
• GIST - Phase 1
• Solid Tumor- Phase 1
• Relapsed multiple Myeloma with end stage renal disease- Phase 1
• Head and Neck Cancer- Phase 1
• Thyroid Cancer- Phase 1

Key areas of Responsibility:
• To assist in protocol writing and review process.
• To prepare and conduct all site visits including, but not limited to, qualification visits, initiation visits, monitoring visits, motivational visits, audit support visits and close-out visits according to relevant SOP's.
• To undertake other project related tasks, as assigned by Project Manager and LCRA.
• To document all study activities including investigator contacts using relevant forms.

Senior CRA

United Biosource Corporation
July 2013 to December 2014

Responsibilities include management of study sites, conduct pre-investigational site visits, study initiation visits, routine monitoring visits and study close-out visits in accordance with corporate, sponsor, FDA and ICH guidelines. Provide site orientation and guidance to assure successful performance of study protocol; review source documents and case report forms in compliance with stated guidelines; assure completion, submission and maintenance of regulatory documents. Prepare and timely submit accurate monitoring reports and manage regional home office.

Therapeutic Area/ Indication /Lab Experience
• Pulmonary Hypertension-Phase IIIb-1.5 years

Key areas of Responsibility:
• Conduct various types of Monitoring visits including Pre-study, Initiation, Interim monitoring visits and Close-out visits according to Sponsor, FDA and ICH Guidelines.
• Provide timely monitoring visit reports as per guidelines.
• Review of regulatory documentation, lab data, subject data and drug accountability records on-site.

Senior CRA

ICON Clinical Research
April 2012 to February 2013

Responsibilities include management of study sites, conduct pre-investigational site visits, study initiation visits, routine monitoring visits and study close-out visits in accordance with corporate, sponsor, FDA and ICH guidelines. Provide site orientation and guidance to assure successful performance of study protocol; review source documents and case report forms in compliance with stated guidelines; assure completion, submission and maintenance of regulatory documents. Prepare and timely submit accurate monitoring reports and manage regional home office.

Therapeutic Area/ Indication /Lab Experience
• Alzheimer's-Phase III-1 year

Key areas of Responsibility:
• Conduct various types of Monitoring visits including Pre-study, Initiation, Interim monitoring visits and Close-out visits according to Sponsor, FDA and ICH Guidelines.
• Provide timely monitoring visit reports as per guidelines.
• Review of regulatory documentation, lab data, subject data and drug accountability records on-site.

Senior CRA

PRA International, Inc

July 2010 to April 2012

Responsibilities include management of study sites, conduct pre-investigational site visits, study initiation visits, routine monitoring visits and study close-out visits in accordance with corporate, sponsor, FDA and ICH guidelines. Provide site orientation and guidance to assure successful performance of study protocol; review source documents and case report forms in compliance with stated guidelines; assure completion, submission and maintenance of regulatory documents. Prepare and timely submit accurate monitoring reports and manage regional home office.

Therapeutic Area/ Indication /Lab Experience
• Inflammatory Breast Cancer-Phase II-1 year
• Bipolar- Phase III- 1 year

Key areas of Responsibility:
• Conduct various types of Monitoring visits including Pre-study, Initiation, Interim monitoring visits and Close-out visits according to Sponsor, FDA and ICH Guidelines.
• Provide timely monitoring visit reports as per guidelines.
• Review of regulatory documentation, lab data, subject data and drug accountability records on-site.

Nurse Manager

Thomson Hood Veterans Center
July 2009 to June 2010

Responsibilities include management of 60 resident wing in a 285 bed state long term care facility for veterans. Supervise forty nursing assistants and fifteen nurses to ensure that the veterans served are appropriately taken care of. Responsible for managing budget for the unit to ensure fiscal compliance.

Key areas of Responsibility:
• Manage 60 resident long term care unit.
• Supervise 40 nursing assistants and fifteen nurses
• Manage unit budget to ensure fiscal compliance.

Senior CRA

Kendle International, Inc
February 2008 to June 2009

Responsibilities include management of study sites, conduct pre-investigational site visits, study initiation visits, routine monitoring visits and study close-out visits in accordance with corporate, sponsor, FDA and ICH guidelines. Provide site orientation and guidance to assure successful performance of study protocol; review source documents and case report forms in compliance with stated guidelines; assure completion, submission and maintenance of regulatory documents. Prepare and timely submit accurate monitoring reports and manage regional home office.

Therapeutic Area/ Indication /Lab Experience
• Pediatric and adult ADHD-Phase III-1.25 years
• Colorectal Cancer- Phase II- 1 year

Key areas of Responsibility:
• Conduct various types of Monitoring visits including Pre-study, Initiation, Interim monitoring visits and Close-out visits according to Sponsor, FDA and ICH Guidelines.
• Provide timely monitoring visit reports as per guidelines.

• Review of regulatory documentation, lab data, subject data and drug accountability records on-site.

Independent CRA

REV Research LLC
April 2006 to January 2008

Independent CRA
REV Research LLC.

Responsibilities include management of study sites, conduct pre-investigational site visits, study initiation visits, routine monitoring visits and study close-out visits in accordance with corporate, sponsor, FDA and ICH guidelines. Provide site orientation and guidance to assure successful performance of study protocol; review source documents and case report forms in compliance with stated guidelines; assure completion, submission and maintenance of regulatory documents. Prepare and timely submit accurate monitoring reports and manage regional home office. Provide guidance and training for peers and inexperienced Clinical Research Associates. Prepare and timely submit accurate monitoring reports, manage regional home office and manage business affairs of LLC.

Therapeutic Area/ Indication /Lab Experience
• Numerous therapeutic areas such as GI, Rheumatology, Oncology, Cardiovascular and Infectious Disease- Phase I-III-2 years

Key areas of Responsibility:
• Conduct various types of Monitoring visits including Pre-study, Initiation, Interim monitoring visits and Close-out visits according to Sponsor, FDA and ICH Guidelines.
• Provide timely monitoring visit reports as per guidelines.
• Review of regulatory documentation, lab data, subject data and drug accountability records on-site.
• Act as Lead CRA for 2 trials.

Clinical Site Manager

Amgen
September 2004 to April 2006

Responsibilities include management of study sites, conduct pre-investigational site visits, study initiation visits, routine monitoring visits and study close-out visits in accordance with corporate, sponsor, FDA and ICH guidelines. Provide site orientation and guidance to assure successful performance of study protocol; review source documents and case report forms in compliance with stated guidelines; assure completion, submission and maintenance of regulatory documents. Prepare and timely submit accurate monitoring reports and manage regional home office.

Therapeutic Area/ Indication /Lab Experience
• Oncology(B-Cell lymphocytic Leukemia, Prostate and breast)- Phase I-1.5 years
• Chronic Renal Failure- Phase II and III-1 year
• Pediatric Psoriasis-Phase III-1 year

Key areas of Responsibility:
• Conduct various types of Monitoring visits including Pre-study, Initiation, Interim monitoring visits and Close-out visits according to Sponsor, FDA and ICH Guidelines.
• Provide timely monitoring visit reports as per guidelines.
• Review of regulatory documentation, lab data, subject data and drug accountability records on-site.

Independent CRA

REV Research LLC
April 2001 to September 2004

Responsibilities include management of study sites, conduct pre-investigational site visits, study initiation visits, routine monitoring visits and study close-out visits in accordance with corporate, sponsor, FDA and ICH guidelines. Provide site orientation and guidance to assure successful performance of study protocol; review source documents and case report forms in compliance with stated guidelines; assure completion, submission and maintenance of regulatory documents.
Provide guidance and training for peers and inexperienced Clinical Research Associates. Prepare and timely submit accurate monitoring reports, manage regional home office and manage business affairs of LLC.

Therapeutic Area/ Indication /Lab Experience
• Numerous Therapeutic areas such as GI, Cardiovascular, Oncology, Rheumatology and Women's health- Phase I-IV-

Key areas of Responsibility:
• Conduct various types of Monitoring visits including Pre-study, Initiation, Interim monitoring visits and Close-out visits according to Sponsor, FDA and ICH Guidelines.
• Provide timely monitoring visit reports as per guidelines.
• Review of regulatory documentation, lab data, subject data and drug accountability records on-site.

Regional Clinical Monitor

Ingenix, Inc.
March 2000 to April 2001

Responsibilities include management of study sites, conduct pre-investigational site visits, study initiation visits, routine monitoring visits and study close-out visits in accordance with corporate, sponsor, FDA and ICH guidelines. Provide site orientation and guidance to assure successful performance of study protocol; review source documents and case report forms in compliance with stated guidelines; assure completion, submission and maintenance of regulatory documents. Prepare and timely submit accurate monitoring reports and manage regional home office.

Therapeutic Area/ Indication /Lab Experience
• Infectious Disease- Phase II and III- 1 year

Key areas of Responsibility:
• Conduct various types of Monitoring visits including Pre-study, Initiation, Interim monitoring visits and Close-out visits according to Sponsor, FDA and ICH Guidelines.
• Provide timely monitoring visit reports as per guidelines.
• Review of regulatory documentation, lab data, subject data and drug accountability records on-site.

Senior CRA

Phoenix International Health Sciences
December 1997 to March 2000

Responsibilities include management of study sites, conduct pre-investigational site visits, study initiation visits, routine monitoring visits and study close-out visits in accordance with corporate, sponsor, FDA and ICH guidelines. Provide site orientation and guidance to assure successful performance of study protocol; review source documents and case report forms in compliance with stated guidelines; assure

completion, submission and maintenance of regulatory documents. Prepare and timely submit accurate monitoring reports and manage regional home office.

Therapeutic Area/ Indication /Lab Experience
• GI- Phase II and III- 2 years

Key areas of Responsibility:
• Conduct various types of Monitoring visits including Pre-study, Initiation, Interim monitoring visits and Close-out visits according to Sponsor, FDA and ICH Guidelines.
• Provide timely monitoring visit reports as per guidelines.
• Review of regulatory documentation, lab data, subject data and drug accountability records on-site.

CRA I
Clintrials Research, Inc
January 1997 to December 1997

Responsibilities include management of study sites, conduct pre-investigational site visits, study initiation visits, routine monitoring visits and study close-out visits in accordance with corporate, sponsor, FDA and ICH guidelines. Provide site orientation and guidance to assure successful performance of study protocol; review source documents and case report forms in compliance with stated guidelines; assure completion, submission and maintenance of regulatory documents. Prepare and timely submit accurate monitoring reports and manage regional home office.

Therapeutic Area/ Indication /Lab Experience
• Infectious Disease- Phase III- 1 year

Key areas of Responsibility:
• Conduct various types of Monitoring visits including Pre-study, Initiation, Interim monitoring visits and Close-out visits according to Sponsor, FDA and ICH Guidelines.
• Provide timely monitoring visit reports as per guidelines.
• Review of regulatory documentation, lab data, subject data and drug accountability records on-site.

Education

Bachelor's in Healthcare Services
Trinity College
September 2000 to September 2001

Associate in Nursing
Lexington Community College
May 1987 to May 1993

Nursing Licenses

RN

Skills

- Nursing Home
- RN
- Acute Care
- Medical Surgical
- Oncology
- Nurse Management
- Alzheimer's Care
- Clinical Research
- Nursing
- Medication Administration
- Hospice Care
- ICU Experience
- Dementia Care
- Clinical Trials
- Laboratory Experience
- Epic
- GCP
- Travel nursing

Certifications and Licenses

RN License
September 1993 to October 2020

BLS Certification

Additional Information

2000-2001 Bachelor of Science Trinity College and University
1987-1993 Associate of Science Lexington Community College

ADDITIONAL SKILLS
- Microsoft Office (Word, Excel, Access, PowerPoint)

- Operating Systems: MS Windows 97, 2000, XP Vista and 7.
- EDC Systems: DataTrak, eClinical, Inform (Phase Forward), UBC Capture and Medidata Rave.
- CTMS: eClinical
- US English, Fluent
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LOCATION

Location Nicholasville, KY