Contact

www.linkedin.com/in/dat-nguyen-7617a04 (LinkedIn)

Top Skills

Clinical Development
Project Management
Clinical Trials

Dat Nguyen

Director, Medical Communications at Corcept Therapeutics
Menlo Park, California

Summary

-Extensive operational knowledge of drug development process and clinical study management from initial study design (including registries) to publication of study results
-Effective in matrix environments working with Marketing, Sales, Medical, Legal, Regulatory and Research and Development
-Successful development of strategic, scientific, and commercially aligned communication and publication plans to publish and disseminate clinical/scientific data
-Proficient in current practices in medical and regulatory affairs, drug development, FDA and regulatory guidelines (i.e., PhRMA guidelines, ACCME, ICJME etc.)
-Strong leadership and project management skills, ability to work collaboratively in cross functional teams in addition to operating independently
-Trained in Six Sigma and Change Management methodologies to facilitate organizational efficiencies and process improvements
-Diverse therapeutic knowledge: Transplantation, Immunology, Urology, CNS, Metabolism (Obesity), Cardiovascular, Orthopedics (Hand Surgery)

Specialties: Medical Affairs, Drug Information, Publications, Clinical Development, Project Management, Registries

Experience

Corcept Therapeutics
6 years 8 months

Director, Medical Communications
January 2017 - Present (3 years 5 months)
Menlo Park, CA

Associate Director, Medical Communications

PRA International

Scientific Affairs Director, Late Phase Services

June 2011 - September 2013 (2 years 4 months)

Supports global and regional post-approval studies by planning and conducting safety-surveillance studies, large simple trials, registries, restricted access programs, risk management programs, and diagnostic and biomarker research.

Auxilium Pharmaceuticals

Assoc. Director,Medical Affairs

April 2008 - June 2011 (3 years 3 months)

-Established the foundation of the Medical Affairs department by implementing policies and procedures to ensure proper compliance with current regulations while servicing the needs of a growing company

-Instituted and established the Drug Information Center, Educational and Research Grants Portal, and enhanced the Patient Assistance Program

oCreated product information support documents, including position papers and standard response letters

oManaged investigator-initiated research proposals through review/ approval process, while identifying and supporting research opportunities for future collaboration

oSupervised the selection/support of independent accredited medical education activities to ensure that the scope is consistent with the company educational objectives, and compliant with all applicable guidelines and regulations

-Medical representative & signatory for promotional review of marketing materials to ensure evidence-based approach, medical accuracy and fair-balance

-Involved in publication strategy and execution, including writing, reviewing and approving abstracts, posters, and manuscripts for scientific accuracy and adherence to regulations

-Effectively collaborated with Key Opinion Leaders, external advisors, and professional societies to address medical and commercial needs

-Served as a scientific resource for the commercial team by attending scientific meetings, conducting competitive product analysis, and identifying market opportunities

-Executed and managed a large observational post-marketing registry

March 2006 - April 2008 (2 years 2 months)

Functioning as a project manager for multiple Phase III global clinical trials across therapeutic areas (Immunology-Transplant; Metabolism-Obesity; Cardiovascular - DVT)

Novartis
Sr. Clinical Research Scientist
July 2002 - March 2006 (3 years 9 months)

Education

Rutgers University
· (2002 - 2004)

Albany College of Pharmacy and Health Sciences
PharmD · (1996 - 2002)