

## Statistician Position Description & Responsibilities

Vanda is seeking a mid-level Clinical Statistician who will primarily be responsible for statistical operations, analysis of clinical studies, data presentation, report writing, and providing statistical representation to the FDA, as well as statistical consulting support in preclinical discovery work. Roles and responsibilities include the following: Liaise with CRO trial statisticians and data managers. Develop Statistical Analysis Plans with contractors. Prepare the statistical section of clinical protocols. Assist with the design of Clinical Report Forms and contribute to Clinical Study Reports. Position Requirements Successful candidates will have the following: A M.S. or Ph.D. in Statistics (if M.S., then 3-4 years of additional relevant pharmaceutical experience will be needed). Will be familiar with clinical trial methodology and preparation of data for FDA/CDER submission. Experience liaising with the FDA/CDER. SAS programming skills. Excellent practical, organizational, interpersonal, oral and written communication skills are a must. Formal training in pharmacology or chemistry a plus, but not required. Vanda Pharmaceuticals offers a competitive compensation package with excellent benefits and a 401(k) matching program. Vanda is an equal opportunity employer, committed to the hiring, advancement and fair treatment of individuals without regard to race, color, religion, sex, sexual preference, age, national origin, ethnicity, disability, or any other protected status designated by law.

Lyndsay Ferro  
Account Manager  
The Cambridge Group Ltd  
a part of On Assignment Clinical Research  
(800) 525-3396 ext. 285  
(203) 226-4243  
(203) 226-3856 FAX  
lferro@cambridgegroup.com  
[www.cambridgegroup.com](http://www.cambridgegroup.com) / [www.oaclinicalresearch.com](http://www.oaclinicalresearch.com)

Best wishes,

Sutan Wu  
Ph.D Candidate  
Statistics Department  
Florida State University

(850)5667129  
sutanwu@stat.fsu.edu