Drug Safety Research Analyst

AdverseEvents makes previously unavailable post-approval drug side effect information accessible and actionable with tools and analytics that allow decision makers throughout the healthcare industry to quickly understand and analyze the impact of drug safety on a comparative and predictive basis.

With proprietary technologies and unique product offerings, AdverseEvents is positioned to disrupt the entire post-approval drug industry with the broad roll out of AdverseEvents Explorer and accompanying service offerings.

We are currently seeking a **Drug Safety Research Analyst** to work on site in our Santa Rosa, CA office. The primary role of the Drug safety Research Analyst is to work with our analyst and tech teams to conduct online research, write topical papers and drug safety news stories, help to keep our product up-to-date with new, real-time information relating to drug safety, and interact with clients on an as-needed basis. Occasional / light travel to industry conferences or client meetings is required.

Qualifications:

- PharmD degree
- Expertise or background in drug safety preferred
- Expertise in MS Excel preferred and proficient in MS Office products required
- Excellent Internet research, analytical, and writing skills
- Detail oriented
- Perform under time pressure while preserving data integrity
- Ability to work independently or in a team setting

Compensation/Benefits:

- Salary commensurate with experience, education, and background
- Equity participation through stock options.
- Paid sick and vacation days*
- Standard benefits including health and dental plans*

* Eligible after 90 days of employment.

Adverse Events, Inc. is an equal opportunity employer. However, the company prefers candidates already located in the Santa Rosa or North Bay area to minimize transition time. The company will not pay moving expenses for any candidates. The company will not consider candidates from outside the U.S. for this position.

About AdverseEvents, Inc.

AdverseEvents is a healthcare informatics company that improves patient safety and reduces systemic healthcare costs through the comprehensive analysis of post-marketing drug side effect data. Utilizing data-mining and analysis technology, through its proprietary RxSuite™ of analytics, AEI makes post-marketing drug safety data accessible, actionable, and predictable.

AEI provides services to enterprise markets including managed care organizations, the pharmaceutical industry, and financial institutions, all while keeping the patient as the key stakeholder.

To apply, please submit cover letter, resume, and compensation expectations to brian@adverseevents.com.