Natasha Bonds

169 Clinton Avenue

Newark, New Jersey 07108

Email: [bonds.portfolio@gmail.com](mailto:bonds.portfolio@gmail.com)

Phone: 908.764.2432

Objective

To obtain a position that will allow me to utilize my analytical skills for the management of information and information systems.

Education

Degree

Hampton University, Bachelor of Science, Finance (Cum Laude), Hampton, Virginia

Certifications

Oracle Certified Associate 11g (OCA), currently pursuing. Expected by August 2016

Languages, Databases, Database Application, OS, and Developer Tools

*Languages*

SQL Implementations: MySQL, SQL\*Plus, T-SQL, Oracle Database SQL (Advanced Level)

PHP, Java, and C# (Beginners Level)

*Databases (Command-line and UI)*

Oracle Database 11g, Oracle Clinical, MySQL Workbench, SQL Server Express, and Oracle Application Express (APEX)

*Operating System Administration*

Linux (edX) - currently evaluating the course

*Misc Developer Tools*

Netbeans, Eclipse, Git bash/Github, Filezilla, Composer, and Oracle Virtualbox

Abbott Laboratories Abbott Park, Illinois Apr 2005 - Jan 2006

Experience

**Clinical Data Manager Database: Oracle Clinical**

Responsible for the overall quality and accuracy of clinical research databases. Worked closely with team members (DB Programmer, Statistics, and Safety) to perform the following responsibilities:

•Database Setup, Testing (including electronic validations), and Entry Conventions.

•Electronic Discrepancy/Validation Management (QA).

•Database Freezing/Locking.

Advanced Clinical Research Division Lincolnshire, Illinois Sep 2002 – Apr 2005

**Clinical Data Manager Database: Oracle Clinical**

Responsible for the overall quality and accuracy of databases for outsourced research. Worked closely

with client’s Data Management and Clinical Team to communicate to resolve issues in a timely manner.

Responsibilities included the following:

•Protocol Review and Study Setup (review of database, validations, and the development of Data

Entry Conventions for internal data entry team).

• Electronic Discrepancy Management (QA) and Database Audits (QC).

• Database Freezing and Locking.

TAP Pharmaceutical Products Lake Forest, Illinois Oct 2001 - Sep 2002

***R&D Paralegal***

Worked with corporate legal counsel functioning as legal support for R&D related projects.

Responsibilities included the following:

•Worked with clinical to execute contracts that would best meet the needs of project goals while

minimizing corporate risk exposure.

•Drafted and negotiated agreements related to R&D projects. Ensured that agreements were in

compliance with corporate policies (including policies addressing confidentiality, the indemnification

of researchers, and IP rights).

***Planning and Development, Resource Management***

Served as vendor liaison for matters relating to the management of payables for various clinical trials. Responsibilities included the following:

•Reviewed service contracts to gain an understanding of the services and rates agreed upon.

•Audited the company’s payable receipts from CROs.

•Worked with Clinical teams to validate labor and pass-thru expenses and to approve final

invoices for payment.

Kendle International Northbrook, Illinois Nov 1999 – Oct 2001

***Clinical Data Associate* Database: SAS-based**

Managed clinical databases and ensured the overall accuracy and quality of client databases. Worked

closely with client’s Data Management and Clinical teams to communicate and resolve issues in a timely

manner. Responsibilities included the following:

•Protocol Review and Study Setup (review of database, validations, and conventions).

•Electronic Discrepancy/Validation Management (QA) and Database Audits (QC).

•Study Closeout (QC and preparation for Database Freezing and Locking).

The University of Chicago, IRB Chicago, Illinois, May 1996 – Nov 1999

***Administrator/Supervisor of Regulatory Compliance, Aug 1997 – Nov 1999* Database: MS Access**

Responsible for the administration of the institution’s ethical review program for biomedical research.

Ensured that clinical research conducted at the institution were IRB approved as required by federal

regulations. Responsibilities included the following:

•Supervised Human Subject Regulatory Staff.

•Developed Standard Operating Procedures (SOPs) and IRB forms and reviewed all operational

documents annually.

•Organized special subcommittees to address emergent issues and advised Associate Director on

implementation strategies of new procedures.

***Administrator of Regulatory Compliance, May 1996 –Aug 1997***

Responsibilities included the following:

•Responsible for the pre-review of protocols, consent forms, and recruitment materials.

•Coordinated monthly committee meetings for the review of new and amended protocols and

drafted correspondence to investigators detailing the results of the committee’s review.

•Educated researchers and clinical research coordinators on human subject regulations and the IRB

submission process.

Teaching Experience

Urban Renewal Corp. Kearny, New Jersey July 2011 – Jan 2012

***GED Math Instructor***

Serviced Essex County residents through the teaching of basic and intermediate level math skills. Administered GED practice exams. Created lesson plans and training materials. Organized and held small group teaching sessions. Tracked student progress and adjusted lesson plans accordingly.

Volunteerism

Missionaries of Charity Newark, NJ, Summer 2012/April 2016

***Interim Housemother***

Assisted in the supervision of a small night shelter operated by the Missionaries of Charity (an

organization established by Mother Teresa, a Roman Catholic nun). Oversaw the night

operations for approximately 10 homeless women.