# Keiana Dunn DATA SCIENTIST

## CONTACT

**(718) 930-8517** 

keianadunn@gmail.com

Ounion, New Jersey

in linkedin.com/in/keiana-dunn

github/ktd2001

# **SKILLS**

## Personal

Stakeholder engagement Public Speaking Project management

#### Languages

Python RStudio Linux

#### **Database Management**

SQL Server

### Visualization

Matplotlib ggplot

#### Frameworks/Packages

Pandas NumPy Tidyverse R markdown

Scikit-learn

Keras

Open CV

Excel

## **Machine Learning**

Regression Classification Feature engineering Model selection

## **EDUCATION**

## **MS Data Science and Strategic Analytics**

Stockton University 2019 Galloway, New Jersey Concentrations in Machine Learning, Data Analysis, Data Gathering & Warehousing, Data Visualization.

# **BS Animal Science and Management**

University of California, Davis 2002 Davis, California Fundamental courses Mathematics, Animal Modeling, Biostatistics, Business Organization and Animal Science.

## **EXPERIENCE**

## **Program Director**

Super Smart Enrichment, Inc. 2016 - 2018 Brooklyn, New York

- Translated year-long growth ambitions into short-term day to day operations.
- Developed and implemented educational, enrichment, and skill-based programs for elementary students in collaboration with teachers and stakeholders.
- Optimized networking capacity to attract clients and stakeholders.
- Prepared and monitored program's budget.
- Developed SMART goals and tracked metrics shrinking the revenue gap by 18% year over year.

#### **Project Coordinator**

Center for Law & Social Justice 2016 - 2017 Brooklyn, New York

- Implemented data-driven project management strategies to grow community engagement volumes by 61% and surpassed funding target by 25%.
- Collected, tabulated and reported research data for advocacy programs resulting in \$100K grant funding for succeeding year.

#### **Associate Scientist**

Merck & Co. 2010 - 2016 Kenilworth, New Jersey

- Produced early stage invivo modeling results for Diabetes, Thrombosis, and Infectious Disease therapeutic areas.
- Lead projects from inception, planning, execution through completion.
- Authored standard operating procedures, scientific study protocols and clinical study reports.
- · Conducted monitoring of ongoing studies.
- Collaborated with key stakeholders, investigators, operational teams, regulatory agencies, and ethical committees.

## **Assistant Scientist (formerly Schering-Plough Research Institute)**

Merck & Co. 2006 - 2010 Summit, New Jersey

- Implemented pre-clinical development for invivo investigative research in Diabetes, Obesity, Respiratory, Immunology and Cardiovascular therapeutic areas.
- Co-developed and co-managed toxicology trials working closely with clinical monitors, investigators, regulatory agencies, ethical committees, and key stakeholders.
- Co-authored scientific and clinical sections of study protocols, study abstracts, publications and presentations for scientific community platforms for early drug development.