# Anika Gupta

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## **EDUCATION**

**Harvard University** 

2018 – 2023 (Expected)

Doctor of Philosophy, Bioinformatics and Integrative Genomics

Cambridge, MA

Massachusetts Institute of Technology

June 2018

Bachelor of Science, Computer Science and Molecular Biology

Cambridge, MA

**Relevant Coursework:** Machine Learning, Human Pathology, Deep Learning for Biomedical Data, Computational and Functional Genomics, Design and Analysis of Algorithms, Genetics, Concepts in Genome Analysis, Healthcare Ventures

## RESEARCH EXPERIENCE

## **Broad Institute of Harvard and MIT, PI: Eric Lander**

Jan 2019 – Present

Deep Learning Genomics Researcher

Cambridge, MA

Applying Variational Autoencoders and Latent Dirichlet Allocation to single cell- RNA-sequencing and ATAC-seq data (250k cells X 20k genes/enhancers) to systematically learn universal and cell type-specific regulatory pathway signatures.

#### Stanford University, PI: Dennis Wall

June 2017 – June 2018

Autism Data Science Researcher

Palo Alto, CA

Implemented coalitional game theory and unsupervised machine learning techniques on 4,610 autism spectrum disorder genomes to map the genotype-phenotype bridge and enable a causative, molecular understanding for more precise treatments.

# Broad Institute of Harvard and MIT, Pls: Aviv Regev and Kasper Lage

Sep 2015 - June 2018

Statistical Genomics Researcher

Cambridge, MA

Detected cancer vulnerabilities through a robust statistical framework that capitalizes on the identification of protein interaction networks under purifying selection in 4,700 cancer genomes.

#### **Foundation Medicine**

May 2016 – May 2017

Cancer Genomics Researcher

Cambridge, MA

Characterized the therapeutically targetable gene alteration landscapes of lung cancers through analysis of >20,000 patient tumors to help make personalized medicine a reality.

Merck Bioinformatics Researcher May – Aug 2015

Palo Alto, CA

Analyzed gene expression and signaling pathway data for 416 genes to develop immunotherapy combination signature-predicting algorithms for synergistic therapeutic delivery against melanoma.

## **PROJECTS**

**5AM Ventures,** Fellow

Jul – Dec 2019

One of two fellows, guiding strategic decisions in the venture creation arm. Designed the bioinformatics platform for an immuno-oncology company, outlined the translational biomarker strategy for a neurodegenerative disease company, and conducted scientific diligence in the regenerative medicine and RNA splicing domains for two potential new company builds.

## RA Capital, Business of Biotechnology Advanced Concepts Short Course

June 2019

Chosen from >100 applicants to learn and debate principles of drug pricing, company formation, and biotech investment landscapes. Developed frameworks for thinking about value creation and financing strategies.

## Undiagnosed Diseases Network, Bioinformatics Case Analysis Team

Oct - Dec 2018

Standardized and automated (for variant classification) a proof of concept genomic analysis pipeline for the Brigham Genomic Medicine team's efforts to uncover the genetic causes of four previously intractable, rare patient disease cases.

#### Healthcare Innovation and Commercialization, Pathology AI Team

Sep - Nov 2018

Developed a business model to translate a new deep learning framework for pathology-based lung cancer diagnosis from academia to startup. Incorporated feedback from seasoned pathologists and pitched proposal to biotech venture capitalists.

#### Flagship Pioneering, VentureLabs Innovation Intern

Jan - May 2017

Built quantitative models to enable key strategic decisions that could maximize value creation for a mitochondrial-based Series A biotech company being built within this venture firm.

# **PUBLICATIONS**

Sun M\*, **Gupta A**\*, Varma M, Paskov K, Jung J, Stockham N, Wall D (2019). Coalitional game theory facilitates identification of non-coding variants associated with autism. *Journal of Biomedical Informatics Insights*.

**Gupta** A\*, Sun M\*, Paskov K, Stockham N, Jung J, Wall D (2018). Coalitional game theory as a promising approach to identify candidate autism genes. *Pac Symp Biocomputing* 2018. pp. 436-447 (https://doi.org/10.1142/9789813235533\_0040).

**Gupta** A\*, Horn H\*, Razaz P, Kim A, Lawrence M, Getz G, Lage K (2017). Detecting cancer vulnerabilities through gene networks under purifying selection in 4,700 cancer genomes (https://www.biorxiv.org/content/early/2017/11/21/222687.1).

**Gupta** A, Connelly C, Frampton G, Chmielecki J, Ali S, Suh J, Schrock A, Ross J, Stephens P, Miller V (2017). The druggable mutation landscape of lung adenocarcinoma. *Journal of Thoracic Oncology*. Volume 12, Issue 1, S977.

## **CONFERENCE PRESENTATIONS**

**Gupta A\***, Siraj L\*, Jones TR, Bloemendal A, Subramanian V, Lander ES (2019). Systematically learning cellular programs from single-cell transcriptional and chromatin accessibility data. *33<sup>rd</sup> Conference on Neural Information Processing Systems: Learning Meaningful Representations for Life workshop.* Vancouver, CA.

**Gupta A\***, Siraj L\*, Jones TR, Bloemendal A, Subramanian V, Lander ES (2019). Systematically learning biological pathway signatures from single cell gene expression data. *3<sup>rd</sup> Annual Women in Data Science Conference*. Cambridge, MA.

**Gupta A**, Horn H, Razaz P, Kim A, Lawrence M, Getz G, Lage K (2018). Detecting cancer vulnerabilities through gene networks under purifying selection. *American Society of Human Genetics Annual Meeting*. San Diego, CA.

**Gupta A\***, Sun M\*, Paskov K, Stockham N, Jung J, Wall D (2018). Identifying candidate autism genes via coalitional game theory. *American Medical Informatics Association Informatics Summit*. San Francisco, CA.

**Gupta** A\*, Sun M\*, Paskov K, Stockham N, Jung J, Wall D (2018). Coalitional game theory as a promising approach to identify candidate autism genes. 23<sup>rd</sup> Annual Pacific Symposium on Biocomputing. Big Island, HI.

Frampton G, Hartmaier R, Sokol E, **Gupta A**, Greenbow J, Roels S, Gay L, Stephens P (2018). Novel CDH1 mutations in breast invasive lobular carcinoma. *American Association for Cancer Research Annual Meeting*. Chicago, IL.

**Gupta A**, Chalmers Z, Connelly C, Frampton G, Chmielecki J, Ali S, Suh J, Schrock A, Ross J, Stephens P, Miller V (2016). The druggable mutation landscape of lung cancer. *IASLC 17<sup>th</sup> World Conference on Lung Cancer*. Vienna, Austria.

**Gupta A**, Horn H, Lawrence M, Getz G, Lage K (2015). Identifying and targeting gene networks under purifying selection. *11<sup>th</sup> Annual Broad Institute Symposium*. Cambridge, MA.

#### **AWARDS**

| National Human Genome Research Institute, T32 Grant  | 2018 |
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| Pacific Symposium on Biocomputing, National Library of Medicine/National Institutes of Health Travel Award | 2017 |
| Grace Hopper Conference for Women in Computing, Microsoft Scholarship Recipient                            | 2016 |
| Intel Science Talent Search Competition, National Semifinalist   | 2014 |
| Siemens Competition in Math, Science, and Technology, National Semifinalist                                | 2012 |

#### **LEADERSHIP**

# MIT Biotechnology Group, Founder and Co-President

June 2015 – June 2017

Established the undergraduate chapter of MIT's first (now >1,000-member) biotech initiative as a liaison between students and the biotech industry. Organize pitch competitions, speaker panels, interactive workshops, mentorship programs, due diligence groups, and interactive symposiums to foster student entrepreneurship within the biotech industry.

**Residential Associate Adviser,** *to Professor Maria Zuber (MIT VP of Research)*Sep 2015 – June 2018 Mentored MIT freshmen as they navigate through their first year academic and personal pursuits at the institute.

## **SKILLS**

Computer Science Languages: Python (keras, scikit-learn, numpy, pandas, scipy), R, Java, HTML, Unix/Linux, LaTex Databases Familiar With Analyzing: The Cancer Genome Atlas, Human Protein Atlas, Project Achilles (cancer vulnerabilities), Genome Aggregation Consortium, Broad Mutation Signatures Database, ClinicalTrials.gov, genome-wide expression (bulk and single cell RNA-seq), chromatin accessibility (ATAC-seq), and sequencing data (UK Biobank and proprietary)

Biochemical: Mutagenesis, Plasmid Preparation (Design, Synthesis, Purification), Primer Design, Polymer Chain Reaction, Gel Electrophoresis, MTT Cell Viability Assays, Spectroscopy (<sup>13</sup>C- & <sup>1</sup>H-NMR, IR), Chromatography (Column, Paper, and Thin Layer), Flow Cytometry, Spectrophotometry, Protein Engineering via Directed Evolution, Western Blotting Languages: English (native), Spanish (proficient), Hindi (fluent)