

BRIAN M SCHILDER, PHD

Passionately pursuing transdisciplinary research to advance human health and knowledge.



Cold Spring Harbor Laboratory

Postdoctoral Research Scientist



EDUCATION

2024

Imperial College London / The Alan Turing Institute
PhD: Clinical Medical Research; Computational Genomics & Machine
Learning

Q London, UK

Thesis: Multi-omic medicine: dissecting the cell type-specific and pleiotropic mechanisms underlying disease genomics at scale

- Aim 1) Dissect the multi-scale mechanisms (e.g. genes, pathways, cell types, phenotypes) underlying all rare disease genomics.
- Aim 2) Decompose the phenome (all diseases and traits) into a unified latent genomic space to identify pleiotropy and disease trajectories at scale.
- Aim 3) Demonstrate and facilitate FAIR (Findable, Accessible, Interoperable, Reproducible) practices.

2017



The George Washington University / Georgetown University

MPhil: Human Evolution; Comparative Neuroscience & Genomics

Thesis: The evolution of the hippocampus and adult neurogenesis: novel insights into the origins of human memory

- Aim 1): Identify human-specific features of hippocampal subfield organization, adult neurogenesis, and their ecological correlates.
- Aim 2) Identify human-specific patterns of hippocampal subfield gene expression.
- Aim 3) Identify the genetic mechanisms mediating the evolution of human hippocampal neuroanatomy and gene expression.

2011



Brown University / Princeton University

ScB: Cognitive Neuroscience; Neurological Diseases & Disorders

Providence, RI, USA

CONTACT

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in LinkedIn

ORCID

G Google Scholar

GitHub

Twitter

YouTube

Personal Website

Lab Website

SUMMARY

11 16+ years of research

25 publications

> 7 preprints

41 software packages

11 databases & apps

i□ 24 talks

■ 14+ years of teaching & team management

TABLE OF CONTENTS

☎ Education

✓ Skills

✓ Expertise

Publications

Preprints

Acknowledgements

Reviewerships

Internal talks

Invited talks

†□ Conference talks

i□ Posters

(E) Experience

Packages

□ Websites

Databases

\$ Grants

Q Awards

Affiliations

▼ Data visualisation

Extracurricular



Updated Sep-09-2025



CORE SKILLS

Research

- 16+ years of deep expertise in genomics, AI, evolutionary biology and biomedicine. Strategically fuses concepts and methods across multiple domains.
- Publication record: 25 publications, 7 preprints and 14 awarded grants.
- Reproducibility: Global leader in promoting and enabling High-performance computing: Highly parallelised reproducible scientific practices. Writes 100% reproducible manuscripts programmatically.
- Bioinformatics: Created 45 Python and R packages to address key challenges in biological research.
 - analyses and AI model training (CPUs and GPUs).
 - · Web development: 6+ websites, web apps, and interactive reports.

AI & Machine Learning

Proficient in developing and deploying AI/ML models (PyTorch, tensorflow, Keras, sklearn and H2O) to solve complex biological problems. Applied examples include:

- predictions from DNA sequence models (DeepSEA, Basenji, IMPACT) to validate SNPs prioritised with Bayesian fine-mapping.
- Foundation models: Used transformer trained on >36M cells and protein sequence embeddings to uncover cell type-specific mechanisms of disease.
- LLM knowledge extraction: Developed framework to extract quantitative metrics of phenotype severity from GPT-4.
- · Causal variant effect prediction: Used functional impact · Disease genomics embeddings: Developed VAE/graph models to reveal joint latent representation of genomic signatures across all diseases and phenotypes.
 - NLP: Created a suite of proprietary Python packages for advanced topic modelling of the PubMed literature to provide business intelligence to the world's largest digital health, biotech, and pharma companies (as a consultant with 120/80 Group).
 - Tensor decomposition: Applied multi-condition factorisation to efficiently discover neurodegenerationrelevant trans-eQTLs

Project Management

Efficient management strategies to define objectives, track progress and coordinate diverse teams.

- **Documentation**: Defines objectives and tracks progress with GitHub Projects. Includes useful documentation in Issues, inline code and shareable reports.
- Version control: Extensive and daily use of GitHub, containers (Docker, Singularity, virtual machines),
- environments (conda) and pipelines (Nextflow).
- Team management: Led numerous collaborative research projects and supervised researchers at various career stages.

Soft Skills

Advances science through effective problem formulation, collaboration and communication.

- Problem formulation: Rapid hypothesis generation. project design, and creative problem solving.
- Collaboration: Diverse and global collaborative networking.
- · Communication: Clear and concise distillation of complex results to a variety of audiences. Presented 25 conference posters.

✓ EXPERTISE BY FIELD

The total height of each column represents my overall expertise in a given domain. The color and height of each rectangle are scaled to my relative level of expertise within each subdomain.

Neuroscience Medicine Genomics ΑI Computer Science Evolution [EHR] MLOps HTML / CSS / JS / Palaeontology Development Ethnic diversity Prognosis Shiny Imaging Human origins Databases Drug discovery Histology Systems biology NLP Diagnosis Phylogenetics Containers Memory Gene therapy Comparative biology Causal inference High-performance Neurodegeneration Translation Proteomics computing LLMs CI/CD Target discovery Single-cell Deep learning GitHub transcriptomics / Biomedical epigenomics / ontologies multi-omics Python Foundation models GWAS / QTL Knowledge graphs R Rare diseases Embeddings Common diseases

PUBLICATIONS CUT&Tag recovers u Nature Communications L Abbasova, P Urbanavic Chromatin Interactio

2023

2023

2023

2023

2023

2022

2022

2021

CUT&Tag recovers up to half of ENCODE ChIP-seq peaks

Nature Communications (2025) (16):2993; https://doi.org/10.1038/s41467-025-58137-2

L Abbasova, P Urbanaviciute, D Hu, JN Ismail, BM Schilder, A Nott, NG Skene, SJ Marzi

Chromatin Interaction and Histone Mark Signatures Associated With TBXT Expression in Metastatic Lung Cancer

Genes Chromosomes Cancer, (2025) (64):e70041; https://doi.org/10.1002/gcc.70041 RM Yaa, BM Schilder, RD Acemel, FC Wardle

rworkflows: automating reproducible practices for the R community

Nature Communications (2023) 15(149); https://doi.org/10.1038/s41467-023-44484-5 BM Schilder, AE Murphy, NG Skene

■ News

- Featured in Nature Communications Editors' Highlights

Artificial intelligence for neurodegenerative experimental models

Alzheimer's & Dementia (2023) http://doi.org/10.1002/alz.13479

SJ Marzi, **BM Schilder**, A Nott, C Sala Frigerio, S Willaime-Morawek, M Bucholc, DP Hanger, C James, PA Lewis, I Lourida, W Noble, F Rodriguez-Algarra, JA Sharif, M Tsalenchuk, LM Winchester, U Yaman, Z Yao, DEMON Network, JM Ranson, DJ Llewellyn

Artificial intelligence for dementia genetics and omics

Alzheimer's & Dementia (2023) http://doi.org/10.1002/alz.13427

C Bettencourt, NG Skene, S Bandres-Ciga, E Anderson, LM Winchester, IF Foote, J Schwartzentruber, JA Botia, M Nalls, A Singleton, **BM Schilder**, J Humphrey, SJ Marzi, CE Toomey, A Al Kleifat, EL Harshfield, V Garfield, C Sandor, S Keat, S Tamburin, C Sala Frigerio, I Lourida, DEMON Network, JM Ranson, DJ Llewellyn

Artificial intelligence for dementia research methods optimization

Alzheimer's & Dementia (2023) http://doi.org/10.1002/alz.13441

M Bucholc, C James, A Al Khleifat, A Badhwar, N Clarke, A Dehsarvi, CR Madan, SJ Marzi, C Shand, **BM Schilder**, S Tamburin, HM Tantiangco, I Lourida, DJ Llewellyn, JM Ranson

EpiCompare: R package for the comparison and quality control of epigenomic peak files

Bioinformatics Advances (2023) 13(1):vbad049; https://doi.org/10.1093/bioadv/vbad049

S Choi, BM Schilder, L Abbasova, AE Murphy, NG Skene

 Dissecting the Shared Genetic Architecture of Suicide Attempt, Psychiatric Disorders, and Known Risk Factors

Biological Psychiatry (2022) 91(3):313-327; https://doi.org/10.1016/j.biopsych.2021.05.029 N Mullins, J Kang, AI Campos,...BM Schilder, et al.

Genetic analysis of the human microglial transcriptome across brain regions, aging and disease pathologies

Nature Genetics (2022) https://doi.org/10.1038/s41588-021-00976-y

K de Paiva Lopes, G JL Snijders, J Humphrey, A Allan, M Sneeboer, E Navarro, **BM Schilder**...T Raj

- Microglial transcriptomics meets genetics: new disease leads (Nature Reviews Neurology, 2022)
- Mighty MiGA: Microglial Genomic Atlas Zeros in on Causal AD Risk Variants (ALZFORUM, 2022)
- Can a Human Microglial Atlas Guide Brain Disorder Research? (Mount Sinai Health System, 2022)
- Polygenic Scores Paint Microglia as Culprits in Alzheimer's (ALZFORUM, 2021)

Multi-omic insights into Parkinson's Disease: From genetic associations to functional mechanisms

Neurobiology of Disease (2021) 105580; https://doi.org/10.1016/j.nbd.2021.105580 BM Schilder, E Navarro, T Raj

2021	•	Fine-Mapping of Parkinson's Disease Susceptibility Loci Identifies Putative Causal Variants Human Molecular Genetics (2021) ddab294; https://doi.org/10.1093/hmg/ddab294 BM Schilder, T Raj
2021		echolocatoR: An Automated End-to-End Statistical and Functional Genomic Fine-Mapping Pipeline Bioinformatics (2021) btab658; https://doi.org/10.1093/bioinformatics/btab658 BM Schilder, J Humphrey, T Raj
2021		MungeSumstats: A Bioconductor Package for the Standardisation and Quality Control of Many GWAS Summary Statistics Bioinformatics (2021) 37(23):4593-4596; https://doi.org/10.1093/bioinformatics/btab665 A Murphy, BM Schilder, NG Skene
2021		Dysregulation of mitochondrial and proteo-lysosomal genes in Parkinson's disease myeloid cells Nature Genetics (2021) https://doi.org/10.1101/2020.07.20.212407 E Navarro, E Udine, K de Paiva Lopes, M Parks, G Riboldi, BM SchilderT Raj B News - Mount Sinai: Fighting Neurodegenerative Disorders (Mount Sinai Health System, 2019)
2021		Phenome-wide and eQTL Associations of COVID-19 Genetic Risk Loci iScience (2021) https://doi.org/10.1016/j.isci.2021.102550 C Moon, BM Schilder, T Raj, K-I Huang
2021		Genome-Wide Association Study of over 40,000 Bipolar Disorder Cases Provides Novel Biological Insights Nature Genetics (2021) 53:817-829; https://doi.org/10.1038/s41588-021-00857-4 N Mullins, AJ Forstner, KS O'Connell, B Coombes, JRI ColemanBM Schilder et al. News Researchers identify 64 regions of the genome that increase risk for bipolar disorder (EurekAlert, 2021) Largest Bipolar Disorder Genetics Study Doubles Genetic Risk Factors (Nordic Society of Human Genetics and Precision Medicine, 2021)
2020		Tensor decomposition of stimulated monocyte and macrophage gene expression profiles identifies neurodegenerative disease-specific trans-eQTLs PLOS Genetics (2020) 16(9):e1008549; https://doi.org/10.1371/journal.pgen.1008549 S Ramdhani, E Navarro, E Udine, AG Efthymiou, BM Schilder, M Parks, A Goate, T Raj
2019		Evolutionary shifts dramatically reorganized the human hippocampal complex Journal of Comparative Neurology (2019) 528(17):3143-3170; https://doi.org/10.1002/cne.24822 BM Schilder, HM Petry, PR Hof
2019		FAIRshake: Toolkit to Evaluate the Findability, Accessibility, Interoperability, and Reusability of Research Digital Resources Cell Systems (2019) 9; https://doi.org/10.1016/j.cels.2019.09.011 D Clarke, L Wang, A Jones, M Wojciechowicz, D Torre, K Jagodnik, S Jenkins, P McQuilton, Z Flamholz, M Silverstein, BM SchilderA Ma'ayan B News - Chosen as 'Featured Frontmatter' article in Cell Systems
2019		Geneshot: search engine for ranking genes from arbitrary text queries Nucleic Acids Research (2019) 47(W1):W571-W577; https://doi.org/10.1093/nar/gkz393 A Lachmann, BM Schilder, ML Wojciechowicz, D Torre, MV Kuleshov, AB Keenan, A Ma'ayan ■ News - Geneshot: Piercing the Literature to Identify and Predict Relevant Genes (University of Pittsburgh Health Sciences Library System Update, 2019) - The Future of AI at the Hasso Plattner Institute for Digital Health at Mount Sinai (Mount Sinai Health System, 2020)

eXpression2Kinases (X2K) Web: linking expression signatures to upstream cell signaling networks

Nucleic Acids Research (2018) 46(W1):W171-W179; https://doi.org/10.1093/nar/gky458

DJB Clarke, MV Kuleshov, BM Schilder, D Torre, ME Duffy, AB Keenan, A Lachmann, AS Feldmann, GW Gundersen, MC Silverstein, Z Wang

News

- Mount Sinai Faculty Spotlight: Ma'ayan Lab (Mount Sinai Health System, 2018)

 Defining elemental imitation mechanisms: A comparison of cognitive and motor-spatial imitation learning across object- and computer-based tasks

Journal of Cognition and Development (2015) 17(2):221-243; https://doi.org/10.1080/15248372.2015.1053483 F Subiaul, L Zimmerman, E Renner, BM Schilder, R Barr

Take the monkey and run

2015

2015

2014

2022

2025

2024

2024

2024

Journal of Neuroscience Methods (2015) 248:28-31; http://doi.org/10.1016/j.jneumeth.2015.03.023 KA Phillips, MK Hambright, K Hewes, BM Schilder, CN Ross, SD Tardif

- Monkeys on a Treadmill? A Conversation with Dr. Kimberley Phillips (Why Social Science?)

Becoming a high-fidelity - super - imitator: what are the contributions of social and individual learning?

Developmental Science (2014) 18(6):1025-1035; http://doi.org/10.1111/desc.12276 F Subiaul, EM Patterson, BM Schilder, E Renner, R Barr

2014 • Working memory constraints on imitation and emulation

Journal of Experimental Child Psychology (2014) 128:190-200; http://doi.org/10.1016/j.jecp.2014.07.005 F Subiaul, BM Schilder

▶ PREPRINTS

Mondo: Integrating Disease Terminology Across Communities

medRxiv (2022) https://doi.org/10.1101/2022.04.13.22273750

NA Vasilevsky, ... BM Schilder, ..., PN Robinson, CJ Mungall, A Hamosh, MA Haendel

2025 Cell type-specific contextualisation of the human phenome: towards the systematic treatment of all rare diseases

medRxiv (2025) https://doi.org/10.1101/2023.02.13.23285820

BM Schilder, KB Murphy, H Dash, Y Zhang, R Gordon-Smith, J Chapman, M Otani, NG Skene

Gene expression patterns of the developing human face at single cell resolution reveal cell type contributions to normal facial variation and disease risk

bioRxiv (2025) https://www.biorxiv.org/content/10.1101/2025.01.18.633396v1

N Khouri-Farah, EW Winchester, BM Schilder, K Robinson, SW Curtis, NM Skene, E Leslie-Clarkson, J Cotney

Harnessing generative AI to annotate the severity of all phenotypic abnormalities within the Human Phenotype Ontology

medRxiv (2024) https://doi.org/10.1101/2024.06.10.24308475

KB Murphy, BM Schilder, NG Skene

 Navigating the Phenomic Landscape: systematic characterisation of the latent genomic space underlying all traits and diseases

bioRxiv (2024) http://dx.doi.org/10.13140/RG.2.2.12144.26880

BM Schilder, NG Skene

Integrative multi-omics analysis of glial signatures associated with accelerated cognitive decline in Alzheimer's disease

bioRxiv (2024) https://doi.org/10.1101/2024.08.27.24312641

E Schneegans, N Fancy, V Chau, TKD Cheung, E Adair, M Papageorgopoulou, **BM Schilder**, PM Matthews, JS Jackson

2023		Fine-mapping genomic loci refines bipolar disorder risk genes medRxiv (2023) https://www.medrxiv.org/content/10.1101/2024.02.12.24302716v1 M Koromina, A Ravi, G Panagiotaropoulou, BM Schilder, S Ripke, T Raj, JRI Coleman, N Mullins BNews - Currently under journal review
		ACKNOWLEDGEMENTS
2021		eQTL Catalogue: a compendium of uniformly processed human gene expression and splicing QTLs. Nature Genetics (2021) 53:1290-1299; https://doi.org/10.1038/s41588-021-00924-w N Kerimov, JD Hayhurst, K Peikova et al.
2020		Functionally-informed fine-mapping and polygenic localization of complex trait heritability Nature Genetics (2020) https://doi.org/10.1038/s41588-020-00735-5 O WeissbrodAL Price
2019		Wayfinding: The science and mystery of how humans navigate the world. St. Martin's Press (2019) ISBN-13: 978-1250096968; https://www.amazon.co.uk/Wayfinding-Science-Mystery-Humans-Navigate/dp/1250096960 MR O'Connor
2012		EEG oscillations reveal neural correlates of evidence accumulation Frontiers in Decision Neuroscience (2012) 6(106):13-Jan; https://doi.org/10.3389/fnins.2012.00106 M van Vugt, P Simen, L Nystrom, P Holmes, J Cohen
2011		Trial-by-trial adaptation of decision making performance: a model-based EEG analysis Interdisciplinary Perspectives on Cognition, Education, and the Brain (2011) 7; https://www.semanticscholar.org /paper/Trial-by-trial-adaptation-of-decision-making-a-EEG-Vugt-Simen /330371d08842ecd1bda332dd22351a7135b5cb1f M van Vugt, P Simen, J Cohen
		REVIEWERSHIPS
2025		Mixture of Experts Enable Efficient and Effective Protein Understanding and Design Nature Methods (2025) N Sun, S Zou, T Tao, S Mahbub, D Li, Y Zhuang, H Wang, X Cheng, L Song, EP Xing
2025		Compressing the collective knowledge of ESM into a single protein language model Nature Methods (2025) T Dinh, S Jang, N Zaitlen, V Ntranos
2025	•	Large language models identify causal genes in complex trait GWAS Nature Communications (2025) SS Shringarpure, W Wang, S Karagounis, X Wang, AC Reisetter, A Auton, AA Khan

ENCORE: a practical implementation to improve reproducibility and transparency of computational research

Nature Communications (2025)

2025

2025

AHC van Kampen, U Mahamune, A Jongejan, BDC van Schaik, D Balashova, D Lashgari, M Pras-Raves, EJM Wever, AD Dane, R García-Valiente, PD Moerland

L2G: Repurposing Language Models for Genomics Tasks *ISMB/ECCB* (2025) https://doi.org/10.1101/2024.12.09.627422

1011B/200B (2020) 11ttp0://doi.org/10.1101/2021.12.00.02712

W Cheng, J Shen, M Khodak, J Ma, A Talwalkar

Enhancing regulatory variant prioritization via long-range DNA sequences and multi-task 2025 learning The American Journal of Human Genetics (2025) Y Takahashi, QS Wang, T Hasegawa, H Namkoong, F Inoue, K Fukunaga, S Imoto, S Miyano, Y Okada, Japan COVID-19 Task Force Deciphering the tissue-specific functional effect of Alzheimer risk SNPs with deep genome 2024 BioData Mining (2024) https://doi.org/10.1186/s13040-024-00400-1 PV Pugalenthi, B He, L Xie, K Nho, AJ Saykin, J Yan A genetic and transcriptomic assessment of the KTN1 gene in Parkinson's disease risk 2024 Neurobiology of Aging (2024) https://doi.org/10.1016/j.neurobiolaging.2023.11.001 A Moore, PW Crea, M Makarious, S Bandres-Ciga, C Blauwendraat, M Diez-Fairen [Unpublished article] 2024 BMJ Open (2024) Multi-region brain transcriptomes uncover two subtypes of aging individuals with differences in 2023 Alzheimer risk and the impact of APOE₂4 Neuron (2023) https://doi.org/10.1101/2023.01.25.524961 AJ Lee, Y Ma, L Yu, RJ Dawe, C McCabe, K Arfanakis, R Mayeux, DA Bennett, HU Klein, PL De Jager 2023 Summary statistics-based association test for identifying the pleiotropic effects with set of genetic variants Bioinformatics (2023) https://doi-org.iclibezp1.cc.ic.ac.uk/10.1093/bioinformatics/btad182 D Bu, X Wang, Q Li Most pathways can be related to the pathogenesis of Alzheimer's Disease 2021 Alzheimer's Research & Therapy (2021) https://doi.org/10.3389/fnagi.2022.846902 SL Morgan, P Naderi, K Koler, Y Pita-Juarez, D Prokopenko, IS Vlachos, RE Tanzi, L Bertram, WA Hide CLIP: accurate prediction of disordered linear interacting peptides from protein sequences using 2021 co-evolutionary information Bioinformatics (2021) https://doi.org/10.1093/bib/bbac502 Z Peng, Z Li, Q Meng, B Zhao, L Kurgan Single-cell transcriptomics and in situ morphological analyses reveal microglia heterogeneity 2020 across the nigrostriatal pathway Neurobiology of Disease (2020) https://doi.org/10.3389/fimmu.2021.639613 O Uriarte Huarte, D Kyriakis, T Heurtaux, Y Pires-Afonso, K Grzyb, R Halder, M Buttini, A Skupin, M Mittelbronn, A Michelucci Deconstructing cerebellar development cell by cell 2019 PLOS Genetics (2019) https://doi.org/10.1371/journal.pgen.1008630 MJ van Essen, S Nayler, EBE Becker, J Jacob Partitioning the genetic architecture of amyotrophic lateral sclerosis 2019 Nature Neuroscience (2019) https://doi.org/10.1101/505693 IJ Broce,... RS Desikan **忙** INTERNAL TALKS Drug (re)discovery in the age of genomics: multi-omic strategies for identifying disease 2022

treatments

Seminar

Drug Discovery and Trials Optimisation Working Group Deep Dementia Phenotyping Network (DEMON)

2021	ļ	We've tagged a lot of cells, and sorted them in wells, some of the reads were double, So we
		looked into the trouble Seminar
		UK Dementia Research Institute Imperial College London
2021		Beyond GWAS: getting more out of genomic data in the age of machine learning
		Methods Optimisation Working Group Deep Dementia Phenotyping Network (DEMON)
2021		Interspecies translation of single-cell transcriptomic signatures Seminar
		Experimental Models Working Group Deep Dementia Phenotyping Network (DEMON)
2020		Automated consensus fine-mapping of neurological disorder genomics Seminar
		UK Dementia Research Institute (UK DRI)
	ήE	INVITED TALKS
2024		Navigating the Phenomic Landscape: Systematic characterisation of the latent genomic space underlying all traits and diseases Mondo Outreach Call
		Monarch Initiative
2024		Harnessing AI to annotate the severity of all phenotypic abnormalities within the Human Phenotype Ontology Turing Omics Meeting Omics Data Generation & Analysis Group
		The Alan Turing Institute
2023		Multi-omics medicine: investigating shared genetic risk factors to better understand neurodegenerative disease Turing Omics Meeting Omics Data Generation & Analysis Group The Alan Turing Institute
2022		Decomposing the phenome: learning the latent genomic structure underlying thousands of
		diseases and traits Neuroepidemiology of Aging Webinar RUSH Alzheimer's Disease Center (RADC) RUSH University
2022		Drug (re)discovery in the age of genomics: multi-omic strategies for identifying disease treatments Department Seminar 3D (Drug, Disease, Delivery) Center / Department of Pharmaceutical Sciences University of South Dakota
2020		Statistical and functional genetic fine-mapping across multiple disease
		Seminar Alzheimer's Disease Sequencing Project Columbia University / Icahn School of Medicine at Mount Sinai
2020		Statistical and functional genetic fine-mapping across multiple disease Laboratory of Neurogenetics Friday Workshop National Institute on Aging National Institutes of Health

	∱ E	CONFERENCE TALKS
2025	•	Evaluating the sensitivity of genomic foundation models to interindividual variation using biobank-scale personalized genomes Al x Bio
		Wellcome Genome Campus
2023		rworkflows: taming the Wild West of R packages EuroBioc2023
		Bioconductor 45-minute workshop.
2023		rworkflows: taming the Wild West of R packages BioC2023
		Bioconductor 10-minute talk within the Infrastructure Track.
2023		Navigating the rare diseases landscape: A comprehensive approach to identify gene therapy targets based on cell type-phenotype associations Intelligent Systems For Molecular Biology / European Conference on Computational Biology (ISMB/ECCB)
		International Society for Computational Biology (ISMB) 20-minute talk within the Bio-Ontologies COSI Track.
2022		Systematic quantification of animal model viability across human diseases Informatics-Synapse Joint Early Career Researcher Meeting
		UK Dementia Research Institute (UK DRI)
2020		Automated genetic fine-mapping of neurological disorders London Genetics Network
		The Genetics Society 6-minute talk
2019		Parkinson's disease derived monocytes show alteration in the phago-lysosomal pathway American Society of Human Genetics (ASHG) Annual Meeting
		American Society of Human Genetics (ASHG) Co-contributor
2017		Comparative neuroanatomy of navigational maps in primates JB Johnston Club for Evolutionary Neuroscience
		Society for Neuroscience (SfN) Co-contributor
2016		The evolution of human hippocampal gene expression JB Johnston Club for Evolutionary Neuroscience
		Society for Neuroscience (SfN)
2015		The neurobiological effects of exercise on marmoset models of Multiple Sclerosis Marmoset Social
		Society for Neuroscience (SfN)
2015		The neurobiological effects of exercise on marmoset models of Multiple Sclerosis JB Johnston Club for Evolutionary Neuroscience

Society for Neuroscience (SfN)

	f □	CONFERENCE POSTERS
2024		Navigating the Phenomic Landscape: systematic characterisation of the latent genomic space underlying all traits and diseases Target to Patient (2024) https://www.ebi.ac.uk/industry/targettopatient/ BM Schilder, NG Skene
2023		CUT&Tag recovers up to half of ENCODE ChIP-seq peaks Connectome (UK Dementia Research Institute) (2023) https://ukdri.ac.uk/ J Ismail, D Hu, L Abbasova, BM Schilder, A Nott, NG Skene, SJ Marzi
2023		Navigating the rare diseases landscape: A comprehensive approach to identify gene therapy targets based on cell type-phenotype associations Intelligent Systems For Molecular Biology / European Conference on Computational Biology (ISMB/ECCB) (2023) https://www.iscb.org/ismbeccb2023 BM Schilder, KB Murphy, R Gordon-Smith, J Chapman, M Otani, NG Skene
2023		Identification of cell type-specific gene targets underlying thousands of rare diseases and clinical phenotypes Genomics of Rare Diseases (2023) https://coursesandconferences.wellcomeconnectingscience.org/event //genomics-of-rare-disease-20230424/ BM Schilder, KB Murphy, R Gordon-Smith, J Chapman, M Otani, NG Skene
2023		Statistical and Functional Fine-Mapping as a Powerful Tool to Unravel the Biological Etiology of Bipolar Disorder Biological Psychiatry (2023) 93(9):S18; https://doi.org/10.1016/j.biopsych.2023.02.063 M Koromina, A Ravi, BM Schilder, B Muller, J Coleman, T Raj
2023		Systematic quantification of animal model viability across human disease UK Dementia Research Institute Scientific Advisory Board (2023) BM Schilder, NG Skene
2022		Systematic quantification of animal model viability across human disease Rising Scientist Day at Imperial College London (2022) BM Schilder, NG Skene
2022		A compehensive statistical and functional fine-mapping pipeline applied to Bipolar Disorder GWAS risk loci European Neuropsychopharmacology (2022) 63:e14; http://dx.doi.org/10.1016/j.euroneuro.2022.07.037 M Koromina, A Ravi, BM Schilder, B Muller, J Coleman, T Raj, N Mullins
2021		Genetic Effects on Human Microglia Transcriptome in Neuropsychiatric Diseases Biological Psychiatry (2021) 89(9):S84-S85; https://doi.org/10.1016/j.biopsych.2021.02.225 G Snijders, K de Paiva Lopes, J Humphrey, S Allan, M Sneeboer, R Navarro, BM Schilder, R Vialle, M Parks, R Missall, W van Zuiden, F Gigase, R Kubler, AB van Berlekom, C Bottcher, J Priller, R Kahn, L de Witte, T Raj
2020		Cell-type-specific reconstruction of primate evolution from genomic positive selection Rising Scientist Day at Imperial College London (2020) K Murphy, BM Schilder, NG Skene
2019		Automated genetic and functional fine-mapping of Parkinson's Disease Loci American Society of Human Genetics (2019) BM Schilder, T Raj
2019		Parkinson's disease derived monocytes show alteration in the phago-lysosomal pathway American Society of Human Genetics (2019) E Udine, E Navarro,BM Schilder,T Raj

2018	7	Learning X2K: Parameter Optimization via Genetic Algorithms to Calibrate the
		Expression2Kinases Pipeline Illuminating the Druggable Genome (2018)
		BM Schilder, A Lachmann, M Kuleshov, A Ma'ayan
2018		Learning X2K: Parameter Optimization via Genetic Algorithms to Calibrate the Expression2Kinases Pipeline
		Big Data 2 Knowledge - Library of Integrated Network-Based Cellular Signatures (LINCS) (2018) BM Schilder, A Lachmann, M Kuleshov, A Ma'ayan
2017		The evolution of the human hippocampus and neuroplasticity Association for American Physical Anthropologists (2017) https://www.abstractsonline.com/pp8/index.html#! /4071/presentation/4471 BM Schilder, BJ Bradley, CC Sherwood
2016		The molecular evolution of plasticity and the human hippocampus Society for Neuroscience (2016) https://www.abstractsonline.com/pp8/index.html#!/4071/presentation/4471 BM Schilder, BJ Bradley, CC Sherwood
2015		Effects of exercise on disease progression and cognition in the marmoset EAE model JB Johnston Club for Evolutionary Neuroscience (2015) KA Phillips, MK Hambright, K Hewes, BM Schilder, B Jagessar, B t'Hart, SD Tardif
2015		The effects of climatic trends, variability, and rates of change on mammalian brain evolution Association for American Physical Anthropologists (2015) BM Schilder, WA Barr, R Bobe, CC Sherwood
2015		Individual, Observational, and Imitation Learning in Orangutans and Children Association for American Physical Anthropologists (2015) E Renner, BM Schilder, F Subiaul
2014		The helper hinderer task revisited: an infant eye tracking study The George Washington University Research Day (2014) A Gokhale, BM Schilder, F Subiaul
2013		Dendritic morphology of pyramidal neurons across the visual stream: A direct comparison of chimpanzees and humans Society for Neuroscience (2013) BM Schilder, O Adeyo
2013		The striatum in the evolution of learned vocalizations: Understanding the neurobiological precursors to human speech using a chimpanzee model Society for Neuroscience (2013) S Bianchi, T Duka, G Muntane, BM Schilder, CD Stimpson, WD Hopkins
2013		Imitation & emulation in a novel box task Association for Psychological Science (2013) L Zimmerman, N Brito, C Mendelson, R Barr, E Renner, BM Schilder, F Subiaul
0010		
2013	Ĭ	A study of imitation and working memory in 2- to 4- year-olds Association for Psychological Science (2013) R Barr, F Subiaul, L Zimmerman, L Renner, BM Schilder, C Mendelson, L Golojuch
2013		The impact of wealth on sharing preferences in children Child Development Society (2013) J Miller, BM Schilder, L Peizer, F Subiaul

RESEARCH EXPERIENCE

Postdoctoral Research Scientist

Cold Spring Harbor Laboratory (Simons Center for Quantitative Biology)

Ocold Spring Harbor, NY, USA

- · Advancing deep learning applications in genomics and biomedicine in the laboratory of Dr. Peter Koo.
- Developing a genomic foundation model to map complex genome-phenome relationships and make highly accurate, personalized disease risk predictions.

Lead Data Scientist

120/80 Group

2024

2019

2018

2018

2017

2014

2014

2013

2011

New York, NY, USA

- Offers data-driven consultation services to a wide portfolio of high-profile digital healthcare, pharmaceutical and biotech companies.
- Developed a suite of proprietary softwares to extract customised business intelligence from the published literature to generate customised and interpretable reports to clients.
- · Provides clients guidance on strategic AI implementation, data analysis, publication and transparency.

2020 • Bioinformatician II

Icahn School of Medicine at Mount Sinai (Department of Neuroscience / Department of Neurology / Department of Genetics & Genomics / Ronald M. Loeb Center for Alzheimer's Disease)

New York, NY, USA

- Developed machine learning systems to integrate large-scale multi-omics datasets (e.g. whole-genome sequencing, bulk and single-cell RNA-seq, epigenomics, clinical data) to uncover the molecular mechanisms underlying neurodegenerative diseases (e.g. Alzheimer's, Parkinson's, ALS).
- Computationally identified specific disease-causal variants, pathways and cell-types for subsequent functional wet lab validation (e.g. CRISPR-cas9 editing in patient-derived cell cultures, iPSCs and cerebral organoids).

Bioinformatician II

Icahn School of Medicine at Mount Sinai (Department of Pharmacological Sciences) • New York, NY, USA

- Conducted computational systems biology research. Integrated and analyzed large-scale genomic and biomedical data (e.g. Python, R, JavaScript).
- Developed evolutionary algorithm to optimize gene network kinase regulator prediction (eXpression2Kinases).
- Developed and deployed computational tools, software, databases and web applications for basic and clinical research, resulting in 3 peer-reviewed publications.

2017 • Participant

Technische Universität Dresden / eMed (Summer School in Systems Medicine) Frauenchiemsee, Germany

- Attended lectures and extended skills in extraction and analysis of big data from biomedical and neurogenomic resources.
- · Developed, performed and wrote manuscript for collaborative bioinformatics research project in less than one week.

2016 Participant

Icahn School of Medicine at Mount Sinai (Scientific Computing & Data Science)

• New York, NY, USA

 Intensive summer school in high-performance computing, coding, genome database utilization and bioinformatics methods including transcriptomics and genetic association testing.

2017 • Collaborator

Trinity University / Southwestern National Primate Research Center (Department of Neuroscience)

San Antonio, TX, USA

 Investigated the neurobiological mechanisms underlying the ameliorating effects of exercise on relapse-remitting Multiple Sclerosis.

Teaching Assistant / Project Leader

The George Washington University / Rutgers University (Department of Anthropology)

lleret, Kenya

- Served as Teaching Assistant while excavating Lower Paleolithic hominin sites (Homo, Paranthropus).
- As Project Leader, investigated the running biomechanics of local Daasanach tribespeople while mentoring undergraduate students.

Research Assistant

The George Washington University (Department of Anthropology)

Washington, DC, USA

- · Performed dissection, histology, microscopy and quantitative stereology in post-mortem primate brain tissues.
- · Trained junior and senior personnel on lab protocols.

13

2013 1 2011

Senior Lab Manager

The George Washington University (Department of Speech, Language & Hearing Sciences)

Washington, DC, USA

 Organized and trained dozens of undergraduates to conduct weekly cognitive development research; designed and/or directly contributed to over 15 research projects in two years.

2012

Volunteer Researcher

University of Winnipeg / University of Belgrade (Department of Anthropology / Department of Archaeology)

Sićevo, Serbia

· Excavated Paleolithic fossils and tools (H. heidelbergensis, H. neanderthalensis) at Mala Balanica, Velika Balanica, and Pešturina sites.

2011

Volunteer Researcher

Universidad de Murcia (Department of Zoology & Physical Anthropology)

Murcia, Spain

· Excavated Paleolithic fossils and tools from Cueva Negra (H. heidelbergensis) and Sima de las Palomas (H. neanderthalensis) with an international research team.

2011

Volunteer Research Intern

American Museum of Natural History (Division of Anthropology)

New York, NY, USA

· Contributed to paleoanthropological research on primate fossils using 3D morphometry imaging equipment including Minolta, Microscribe and CT.

2010

Paid Research Intern

Princeton University (Princeton Neuroscience Institute)

Princeton, NJ, USA

- · Investigated the neural basis of decision-making in humans.
- Recruited participants, recorded EEG and analyzed data in MATLAB.

2010 2009

Student Researcher

Brown University (Department of Cognitive, Linguistic & Psychological Sciences)

Providence, RI, USA

- · Experimental Analysis of Animal Behavior & Cognition: Conducted various operant conditioning experiments on rats. Gained experience in animal behavioral training, data collection, and data analysis in MATLAB.
- · Laboratory in Genes and Behavior: Tested transgenic mice with modified N-type voltage-gated calcium channel subunits in a battery of cognitive and sensorimotor tasks. Results were published.

TEACHING / MENTORING EXPERIENCE

1 2020

Research Mentor

Imperial College London (Department of Brain Sciences / Department of Life Sciences)

◆ London, UK

- · Mentored students and affiliated projects:
- · Kitty Murphy (PhD): 'Evolutionary pressures on cell types: leveraging species differences to gain insight into neurodegenerative disease risk'
- Sheen Lei (BSc): 'Benchmarking cell-type-specific enrichment of genome-wide disease signatures'
- Ted Reese (MSc): 'Computational cell-type annotation of single-cell epigenomics data'
- · Xindong Sun (MSc): 'Benchmark of Targeted insertion of promoters sequencing (TIP-seq) on histone modification H3K27ac and H3K27me3 in K562 cell line'
- · Shuhan Shen (MSc): 'Evaluation and optimisation of methods for identifying the cell types underlying genetic disease signatures'
- Lusheng Li (MSc): 'Genetic identification of cell types underlying mammalian phenotypes'
- Sera Choi (BSc): 'EpiCompare: R package for QC and benchmarking epigenetic datasets'
- Emilie Cottard (MSc) & Will Lunt (BSc): 'A meta-analysis of selective cell-type vulnerability in Parkinson's Disease neuropathology'
- · Jai Chapman (BSc): 'Expression Weighted Cell Type Enrichment as a Tool for Identifying Cell Types Underlying Rare Disease Phenotypes'
- · Bobby Gordon-Smith (MSc): 'Identification of cell types involved in rare disease-associated human phenotypes'
- · Leyla Abbasova (MSc): 'Analysis and optimisation of CUT&Tag for epigenomic profiling of the brain'
- · Barney Hill (BSc): 'Identification of cell-types associated with latent factors inferred from phenome-wide GWAS summary statistics'

2020 I 2019

Research Mentor

Icahn School of Medicine at Mount Sinai (Department of Neuroscience / Department of Neurology / Department of Genetics & Genomics / Ronald M. Loeb Center for Alzheimer's Disease)

New York, NY, USA

• Mentored MS, MD, and PhD students in projects focused on computational exploration of phenotype clustering and genomic regulation of neurodegenerative diseases.

2018

Research Co-mentor

Icahn School of Medicine at Mount Sinai (Department of Pharmacological Sciences)

New York, NY, USA

- · Mentored students and affiliated projects:
- Vivian Utti (BSc): 'ChEA3: Transcription Factor Enrichment Analysis' as part of the Summer Research Training Program in Biomedical Big Data Science.
- · Mary Duffy (PhD): 'Predicting upstream kinase regulators from interaction network databases'
- · Zach Flamholz (BSc): 'modEnrichr: a suite of gene set enrichment analysis tools for model organisms'

2018

Guest Lecturer

Icahn School of Medicine at Mount Sinai (Department of Pharmacological Sciences)

New York, NY, USA

 Lectured on data visualization in Python and Jupyter notebooks in the PhD/MD course 'Programming for Big Data Biomedicine'.

2017 l 2016

Research Mentor

The George Washington University (Department of Anthropology)

- · Mentored students and affiliated projects:
- · Jamie Kleiner (BSc): 'Animal model simulating MS and exercise's impact on adult hippocampal neurogenesis'

2015

Teaching Assistant

The George Washington University (Department of Anthropology)

- · Course: 'Human Brain Evolution'
- · Guest lectured, graded all assignments and exams, and provided additional educational support during office hours.

2014

Teaching Assistant

The George Washington University (Department of Psychology)

♥ Washington, DC, USA

- · Course: 'Biological Psychology'
- Led undergraduates in article discussions, graded all assignments and exams, and provided additional educational support during office hours.

2014 l 2013

Teaching Assistant

The George Washington University (Department of Anthropology)

♥ Washington, DC, USA

- · Course: 'Biological Anthropology'
- Led undergraduate students in two, 2-hour lab sessions per week, graded lab assignments and exams, and provided additional educational support during office hours.

2013

2012

Research Mentor

The George Washington University (Department of Psychology)

Washington, DC, USA

- Mentored students and affiliated projects:
- · Anushka Gokhale (BSc): 'Infants' Social Assessment of Characters Through Eye Gaze'

2013 | 2011

Lab Protocol Trainer

The George Washington University (Department of Anthropology)

Washington, DC, USA

 Trained undergraduate, graduate, and post-doctoral researchers in Social Cognition Lab and Lab for Evolutionary Neuroscience in a variety of methodological research protocols.



SOFTWARE PACKAGES

1.

MSTExplorer R

Multi-Scale Targets Explorer: Systematically identify, prioritise and visualise cell-type-specific gene therapy targets across the phenome.

- https://github.com/neurogenomics/MSTExplorer
- https://doi.org/10.1101/2023.02.13.23285820

2. **HPOExplorer** \mathbf{R}

> Import, annotate and visualise the 18k+ hierarchically structured clinical phenotypes across the Human Phenotype Ontology.

- https://github.com/neurogenomics/HPOExplorer
- https://doi.org/10.1101/2023.02.13.23285820

KGExplorer 🖫 🕏 3.

Query, construct, and analyse large-scale biomedical knowledge graphs and ontologies.

https://github.com/neurogenomics/KGExplorer

autoCV 😱 😇 👸 🧓 4.

Automatically generate and style your CV from tables.

https://github.com/bschilder/autoCV

anndataR 😱 🕏 5.

> Bring the power and flexibility of AnnData to the R ecosystem, allowing you to effortlessly manipulate and analyze your single-cell data.

https://github.com/scverse/anndataR

gptPhD R 6.

Query Large Language Models for the purposes of systematically extracting biomedical knowledge.

https://github.com/neurogenomics/gptPhD

ThreeWayTest R 7.

Summary statistics-based association test for identifying the pleiotropic effects with set of genetic variants.

https://github.com/bschilder/ThreeWayTest

SCAVENGE 🖳 8.

Variant to function mapping at single-cell resolution through network propagation.

- https://github.com/sankaranlab/SCAVENGE
- https://doi.org/10.1038/s41587-022-01341-y

rworkflows R 🥞 9.

> Continuous integration for R packages. Automates testing, documentation website building, and containerised deployment.

- https://github.com/neurogenomics/rworkflows
- https://doi.org/10.21203/rs.3.rs-2399015/v1

TIPseeker R 10.

R package for post-processing [single-cell] TIP-seq data.

https://github.com/neurogenomics/TIPseeker

PeakvFinders R 11.

R package for mining, calling, and importing epigenomic peaks.

https://github.com/neurogenomics/PeakyFinders

12. graphiti 🖳

14

16

Extract colour palettes from graffiti artworks.

nttps://github.com/bschilder/graphiti

SkillNet 🖳 13. Creates user-specific contribution networks from GitHub Organization repositories.

https://github.com/neurogenomics/SkillNet

phenoRx 🖳

Make cell type-specific predictions for patients based on clinical phenotypes and/or risk genes. https://github.com/neurogenomics/phenoRx

phenomix 😱 🕏 15. R package for the exploration and analysis of many genotype-phenotype datasets at once.

https://github.com/neurogenomics/phenomix

MAGMA.Celltyping R Identify cell types underlying the associations found in GWAS summary statistics.

https://github.com/neurogenomics/MAGMA_Celltyping

EWCE 17. Expression Weighted Celltype Enrichment. https://github.com/NathanSkene/EWCE EpiCompare R 18. 19. orthogene R Interspecies gene mapping.

R package for QC and benchmarking epigenetic datasets.

https://github.com/neurogenomics/EpiCompare

https://doi.org/10.1101/2022.07.22.501149

https://github.com/neurogenomics/orthogene

20. MungeSumstats R

Standardise the format of summary statistics from GWAS.

https://github.com/neurogenomics/MungeSumstats

https://doi.org/10.1093/bioinformatics/btab665

scNLP 😱 21.

Tools for applying natural language processing (NLP) techniques to single-cell (sc) omics data.

https://github.com/neurogenomics/scNLP

scKirby \mathbf{R} 🕏 22.

Automated ingestion and conversion of various single-cell data formats.

https://github.com/neurogenomics/scKirby

geneshotR R 23.

R package for querying and processing results from Geneshot.

https://github.com/bschilder/geneshotR

templateR R 24

Self-updating template for developing R packages.

https://github.com/neurogenomics/templateR

https://doi.org/10.21203/rs.3.rs-2399015/v1

25. echoverseTemplate R

Self-updating template for creating echoverse R packages.

https://github.com/RajLabMSSM/echoverseTemplate/

https://doi.org/10.1093/bioinformatics/btab658

echolocatoR 🖫 🕏 26.

R package for end-to-end statistical and functional fine-mapping with extensive dataset access.

https://github.com/RajLabMSSM/echolocatoR

https://doi.org/10.1093/bioinformatics/btab658

echodata 😱 27.

> Examples of fine-mapped GWAS summary statistics, data formatting functions, and API access to the echolocatoR Fine-mapping Portal.

https://github.com/RajLabMSSM/echodata

https://doi.org/10.1093/bioinformatics/btab658

echoannot R 28

Functions for annotating genomic data with annotations and epigenomic data.

https://github.com/RajLabMSSM/echoannot

https://doi.org/10.1093/bioinformatics/btab658

29. echoplot R

R package for LocusZoom-inspired GWAS/QTL visualization, with API access to LD panels.

https://github.com/RajLabMSSM/echoplot

https://doi.org/10.1093/bioinformatics/btab658

echoconda 😱 🕏 30.

Various utility functions to find, build, and use conda environments from within R.

https://github.com/RajLabMSSM/echoconda

https://doi.org/10.1093/bioinformatics/btab658



Tabix indexing and querying.

- https://github.com/RajLabMSSM/echotabix
- https://doi.org/10.1093/bioinformatics/btab658
- 32. echoLD 🕝 🕏

LD downloading and processing.

- https://github.com/RajLabMSSM/echoLD
- https://doi.org/10.1093/bioinformatics/btab658
- 33. echofinemap \mathbb{R}

Statistical and functional fine-mapping functions.

- https://github.com/RajLabMSSM/echofinemap
- https://doi.org/10.1093/bioinformatics/btab658
- 34. echodeps R

Creates interactive dependency networks for R packages.

- https://github.com/RajLabMSSM/echodeps
- https://doi.org/10.1093/bioinformatics/btab658
- 35. echogithub 😱

Access and process metadata from GitHub.

- https://github.com/RajLabMSSM/echogithub
- https://doi.org/10.1093/bioinformatics/btab658
- 36. devoptera 😱

Practical tools for R developers.

- https://github.com/RajLabMSSM/devoptera
- https://doi.org/10.1093/bioinformatics/btab658
- 37. downloadR 😱

Single- and multi-threaded downloading functions.

- https://github.com/RajLabMSSM/downloadR
- https://doi.org/10.1093/bioinformatics/btab658
- 38. catalogueR 😱

R package for rapid API-access and colocalization of summary statistics from eQTL Catalogue.

- https://github.com/RajLabMSSM/catalogueR
- https://doi.org/10.1093/bioinformatics/btab658
- 39. TopicModeler
 - Proprietary Python package to run advanced topic modeling on text corpuses.
- 40. **LinkReporter** Proprietary Python package to extract job postings and company employee listings from LinkedIn and generate interactive business intelligence reports.
- PubReporter Proprietary Python package to extract relevant scientific literature, gather citations, and generate interactive business intelligence reports.
 - DATABASES / WEB APPS
- 1. EpiArchives 😱

Public archive for EpiCompare reports.

- https://github.com/neurogenomics/EpiArchives
- https://doi.org/10.1101/2022.07.22.501149

2. Rare Disease Celltyping Portal 😱

Web portal connecting to multiple R Shiny apps to explore, visualize, and download cell type-specific enrichment results and systematically prioritised gene targets for over 6,000 rare disease phenotypes.

https://github.com/neurogenomics/rare_disease_celltyping_apps

https://neurogenomics.github.io/rare_disease_celltyping_apps/home

https://doi.org/10.1101/2023.02.13.23285820

3. Parkinson's Disease Omics Review 😱

Data and code associated with the Parkinson's Disease review paper by Schilder, Navarro & Raj (Neurobiology of Disease, 2021).

https://github.com/RajLabMSSM/PD_omics_review

https://rajlabmssm.github.io/PD_omics_review/

https://doi.org/10.1016/j.nbd.2021.105580

4. Selective Vulnerability Meta-analysis 😱

Selective Vulnerability Meta-analysis: Shiny app dedicated to the exploration and dissemination of metaanalysed cell counts manually curated and harmonised from the Parkinson's Disease literature.

https://github.com/neurogenomics/SelectiveVulnerabilityMetaAnalysis

5. MAGMA Files Public R

Gene enrichment files for hundreds of GWAS generated with Multi-marker Analysis of GenoMic Annotation (MAGMA) for use in downstream analyses.

https://github.com/neurogenomics/MAGMA Files Public

6. echolocatoR Fine-mapping Portal R

Access to interactive plots and fine-mapping results across many GWAS/QTL datasets using echolocatoR.

https://github.com/RajLabMSSM/Fine_Mapping_Shiny

https://rajlab.shinyapps.io/Fine_Mapping_Shiny

https://doi.org/10.1093/bioinformatics/btab658

7. COVID-19 Patient Tracker

Web app for summarizing and visualizing real-time EHR data of COVID-19 patients within the Mount Sinai Health System.

8. Tensor Decomposition Shiny App 😱

Interactive application to explore and download all results and plots from Ramdhani et al. (PLOS Genetics, 2020).

https://github.com/RajLabMSSM/Tensor_myeloid

https://rajlab.shinyapps.io/Tensor_myeloid

https://doi.org/10.1101/499509

9. Hippocampal Evolution R

Interactive code, results and visualization for the manuscript "Evolutionary selective pressures dramatically expanded and reorganized the human hippocampal complex".

https://github.com/bschilder/Hippo Eco

https://bschilder.github.io/Hippo_Eco/HPsubfield_eco

https://doi.org/10.1002/cne.24822

10. **Geneshot** 👙 🖑 🗓 🗒

Flexible tool to identify genes associated with any biomedical term and to predict novel target genes.

http://amp.pharm.mssm.edu/geneshot

https://doi.org/10.1093/nar/gkz393

11. **X2K** 👙 👶 🗓 💆

eXpression 2 Kinases (X2K) Web: Automated computational pipeline to infer kinase regulators from weighted or unweighted gene lists.

http://amp.pharm.mssm.edu/X2K

https://doi.org/10.1093/nar/gky458

WEBSITES

1. Personal Website 😇 😇 🧓

nttps://github.com/bschilder/BMSchilder
https://bschilder.github.io/BMSchilder

2. Official Raj Lab Website 😇 💆 🗓

https://github.com/RajLabMSSM/RajLab_website thtps://www.rajlab.org

\$ GRANTS

Total (all grants): \$3,049,872

Total (as primary applicant): \$311,382

Cancer Center Pilot Awards Program, Cold Spring Harbor Laboratory Cancer Center

Project: 'Exploiting interpretable AI to uncover the genetic basis of cellular reprogramming in cancer'

• Role: Co-applicant • Amount: \$100,000

• PI: DM McCandlish

2023 • EuroBioc2023 Scholarship,

Bioconductor

Project: 'rworkflows: taming the Wild West of R packages'

News

- Awarded to support attending the EuroBioc2023 meeting.

• Role: Primary applicant • Amount: \$250

· PI: BM Schilder

2023 • BioC2023 Scholarship,

Bioconductor

Project: 'rworkflows: taming the Wild West of R packages'

News

- Awarded to support attending the BioC2023 meeting. Additionally included free lodging.

• Role: Primary applicant • Amount: \$1,500

· PI: BM Schilder

2023 • Junior Scientist Conference Grant,

The Genetics Society

Project: 'Identification of cell type-specific gene targets underlying thousands of rare diseases and subtraits'

• Role: Primary applicant • Amount: £750

• PI: BM Schilder

2023

2024

2022

Imperial UK Research Institute Impact Acceleration Account,

Imperial College London

Project: 'Creating commercial kit solutions for single cell epigenetic profiling of histone marks and transcription factors'

• Role: Co-applicant • Amount: £80,000

• PI: NG Skene

Turing Community Award,

Alan Turing Institute

Project: 'Multi-omic medicine: dissecting the cell-type-specific molecular mechanisms underlying neurodegenerative disease genomics'

• Role: Primary applicant • Amount: £1,500

• PI: BM Schilder

National Institutes of Health 2022 Project: 'Statistical and functional fine-mapping of bipolar disorder genetic risk loci' · Role: Co-applicant · PI: N Mullins Collaborative Single Cell and Spatial Transcriptomics Studies Award Programme, 2021 **UK Dementia Research Institute** Project: 'Amplifying genome coverage of single cell epigenetic profiling of the human brain' · Role: Co-applicant · Amount: £12.790 • PI: D Hu. NG Skene **National Institutes of Health** 2020 Project: "Cognitive Systems Analysis of Alzheimer's Disease Genetic and Phenotypic Data" · Role: Co-applicant • Amount: \$2,523,431 · PI: T Raj, D Knowles **UK Dementia Research Institute** 2024 Project: 'UK DRI at Imperial Distinguished Studentship' 2020 · Role: Primary applicant • Amount: £217,000 · PI: BM Schilder The Michael J. Fox Foundation 2019 Project: "The Role of Peripheral Myeloid Cells in Parkinson's Disease" 2017 · Role: Fundee • **PI**: T Raj The Michael J. Fox Foundation 2020 Project: 'Functional Fine-Mapping of LRRK2 Locus' 2017 · Role: Fundee PI: T Raj **National Science Foundation** 2017 Project: 'The evolution of the hippocampus and adult neurogenesis: novel insights into the origins of human memory' · Role: Primary applicant · Amount: \$31 543 · PI: BM Schilder Wenner-Gren 2017 Project: 'The evolution of the hippocampus and adult neurogenesis: insights into the origins of human memory' · Role: Primary applicant • Amount: \$19,512 · PI: BM Schilder **Leakey Foundation** 2016 Project: 'The evolution of the hippocampus and adult neurogenesis: Novel insights into the origins of human memory' · Role: Primary applicant · Amount: \$15,000 · PI: BM Schilder **COSMOS Club** 2016 **Project**: 'The evolution of adult neurogenesis across primates' · Role: Primary applicant • Amount: \$3.250 · PI: BM Schilder AWARDS Prize for Computational Reproducibility in Dementia Research, 2023 **UK Dementia Research Institute** Project: 'rworkflows: taming the Wild West of R packages' ■ News - Awarded honourable mention.

2022

Prize for Computational Reproducibility in Dementia Research,

UK Dementia Research Institute

Project: 'MungeSumstats: A Bioconductor package for the standardisation and quality control of many GWAS summary statistics'

News

- Awarded honourable mention.

2022

Poster Competition,

Rising Scientist Day

Project: 'Systematic quantification of animal model viability across human diseases'

News

- Awarded prize for research poster competition.

2022

Award for Outstanding Contribution,

NEUROHACK, Deep Dementia Phenotyping Network (DEMON)

Project: 'Predicting ALS drug targets using integrative multi-modal deep learning'

News

- Individually awarded for outstanding contributions during the NEUROHACK 2022, a competitive 4-day hackathon to apply AI in finding ALS therapeutic solutions.

2021

Prize for Computational Reproducibility in Dementia Research,

UK Dementia Research Institute

Project: 'echolocatoR: an automated end-to-end statistical and functional genomic fine-mapping pipeline' ■ News

- Winners announced for UK DRI's first 'Prize for Computational Reproducibility in Dementia Research (UK Dementia Research Institute, 2021)
- Jointly awarded inaugural prize with Kitty Murphy.

2021

Centre Photography Competition,

UK Dementia Research Institute

Project: 'Wildfire Circle, Golden Brain, Wildfire, Geneshot, Geology of Biology, Neon Brain'

News

- One of the winners of the scientific image competition.

2019

Art of the Brain,

Friedman Brain Institute, Icahn School of Medicine

Project: 'Wildfire'

News

- Awarded 2nd place and Featured on the cover of Biological Psychiatry: Volume 87, Issue 12 (2020). Exhibited and auctioned at the Grady Alexis Gallery (New York City), where all proceeds were voluntarily donated to the Diversity in Neuroscience Initiative.



AFFILIATIONS



Fellow



Member



· Society of Technological Advancement Member



- Synapse Working Group
- · Informatics Working Group



DEMON Network

Data science and Al for dementia

- · Genetics & Omics Working Group
- Experimental Models Working Group
- Drug Discovery & Trials Optimisation Working Group

The Alan Turing Institute

- Turing Enrichment Scheme
- Turing-Roche Strategic Partnership
- Turing Omics Data Generation & Analysis Interest Group
- Turing Clinical Al Interest Group



· Bipolar Disorder Working Group



- Chair of the Bioconductor Cloud Methods Working Group
- · Lead of the Bioconductor GitHub Actions Subgroup



Member

▼ DATA VISUALISATION / ARTWORK PORTFOLIO

2022 • echoverse Dependency Graph

♀ London, UK

• Interactive graph showing the dependency structure of all packages within the echoverse suite.

Hex stickers

2020

2023

2023

2023

2021

2021

Q London, UK

· All hex stickers for R packages I've helped develop.

3D Human Phenotype Ontology

Q London, UK

- 3D force-directed graph of the Human Phenotype Ontology (clouds above) with kernel density estimation projected from the x/y planes (mountains below). Connections represent the hierarchical relationships between rare diseases and their associated symptoms/phenotypes.
- Associated preprint

Multi-scale Rare Disease Mechanisms

♠ London, UK

- Network of systematically prioritised gene therapy targets for rare diseases
- Associated preprint

Curriculum Vitae Connexa

♠ London, UK

• Term co-occurrence network generated by analysing all data that went into this CV.

Experiments with Generative Al

♀ London, UK

- · wombo.art: 'Multi-omic medicine: dissecting the cell-type-specific mechanisms in neurodegeneration genomics'
- wombo.art: 'Multi-omic medicine: neurodegenerative disease genomics'
- wombo.art: 'Multi-omic medicine: neurodegeneration'
- wombo.art: 'Neurodegeneration'

Lights in the dark genome: the current state of Parkinson's research

Q London, UK

- The majority of PD genetics research has focused on a relatively small number of genes. Above, are the top 75 most commonly mentioned genes in the PD literature, extracted using Geneshot.
- · Associated study

2020 Pacrophage

Q London, UK

- · Colocalised genetic loci ...but shaped as Pac-Man!
- · Associated study

2020 Circos

· Colocalised genetic loci across a variety of neurological disease GWAS and cell-type-specific QTLs.

· Associated study

2019 • Wildfire Circle

New York, NY, USA

Q London, UK

- · Awarded 2nd place in the 2019 Art of the Brain competition, put on by the Mount Sinai's Friedman Brain Institute.
- Exhibited and auctioned at the Grady Alexis Gallery (New York City), where all proceeds were donated to the Diversity in Neuroscience Initiative .
- · Featured on cover of Biological Psychiatry.

2019 • Wildfire

2003

1

1995

New York, NY, USA

• Transcriptomic data from 16k+ individual brain cells (shown as points) after reducing the dimensionality with an autoencoder and UMAP. 5 million tracts are shown interconnecting these cells, where shorter tract length represents greater similarity in their molecular profiles.

2019 • 3D Brain Model

New York, NY, USA

• 3D model of my brain generated from MRI scans.

S EXTRACURRICULAR EXPERIENCE

Competitive Running Career

- 8+ years of varsity and Division I cross country, winter track, and spring track throughout high school and college.
- Year-round, daily training and travel to weekly competitions necessitated a dedicated and regimented lifestyle
 in order to succeed as a student-athlete
- Running remains a passion of mine and I enjoy training for ultra-marathons in my free time.
- This passion, work ethic and self-insight have carried over to all aspects of my life, including my career as a researcher.

Earth

Music Production

• Writes, records, produces and performs original music.

Earth

· Instruments: keys, percussion, vocals, etc.

- Proficient in Logic Pro X Digital Audio Workstation (DAW).
- Experiments with generative AI.

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