

# Xenofon Giannoulis

✉ (+49) 15772182145 | ✉ xgiannoulis@proton.me | ✉ xenophong.github.io/web/

## Work Experience

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### Statistical Geneticist

Pheiron GmbH, DE

July 2025 – now

PLATFORM DEVELOPMENT & SCIENTIFIC DISCOVERY

- Developed predictive genetic models scoring 36M+ target-indication pairs to de-risk early-stage drug development.
- Automated MR and Coloc pipelines to validate mechanism chains and predict clinical trial success.
- Architected a unified GWAS catalog using out-of-core DuckDB/Polars ETL pipeline to harmonize FinnGen, PAN-UKB, MVP.
- Designed scalable precomputation pipelines (GWAS, PGS, burden tests) with fast query interfaces for rapid association lookups.
- Built an automated post-ETL QC and validation framework with strict statistical checks to ensure decision-grade data quality.
- Authored end-to-end documentation (architecture, workflows, benchmarks, troubleshooting) to enable team communication.

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### Postdoctoral Researcher

Helmholtz Munich, DE

INSTITUTE OF COMPUTATIONAL NEUROBIOLOGY

January 2025 – June 2025

- Prioritized Alzheimer's therapeutic targets by integrating multi-omics evidence data into knowledge graphs and ML models.
- Expanded the AD Atlas (R Shiny, Neo4j), an interactive network platform enabling rapid disease pathway exploration.
- Engineered graph abstraction layers to translate heterogeneous data into interpretable network topologies.
- Deployed reproducible harmonization pipelines processing high-dimensional omics to extract robust molecular signatures.

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### Scientific Researcher

Helmholtz Pioneer Campus, DE

INSTITUTE OF TRANSLATIONAL GENETICS

June 2020 – December 2024

- Built a multi-tissue atlas of mitonuclear gene regulation in GTEx v8 (684 donors, 48 tissues) using WGS + RNA-seq.
- Implemented LMM eQTL workflows with rigorous QC, mtDNA haplogroup inference, and tissue-specific covariate control.
- Extended discovery to trans effects and causal follow-ups via colocalization, mediation testing, and Mendelian randomization.
- Identified genome-wide trans mito-nuclear regulatory links, including CNS-enriched cell-type interaction effects.
- Integrated eQTL results into network-based gene prioritization using SPEOS (GNN/MLP; TAG graph convolution).
- Built an RNA-seq pipeline to quantify mtDNA heteroplasmy and mtRNA modifications (12,577 samples, 49 tissues).
- Presented consortia findings and authored two manuscripts currently under revision at Nature Communications.

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### Research Fellow

LMU, Munich, DE

DATA SCIENCE FOR SOCIAL GOOD | MUNICH CENTER FOR MACHINE LEARNING

August 2023 – September 2023

- Built a geospatial planning tool for fire brigades to model hose reachability under real-world obstacles (stairs, rails, buildings).
- Computed reverse-planning polygons for where a water point must lie to cover a selected building/site.
- Developed a full-stack AWS web app for interactive mapping, real-time routing, and resource planning in the field.
- Delivered system to the Federal Office for Disaster Control & Munich Fire Brigade; presented to the German federal parliament.

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### Instructor, Human Genetics of Complex Traits

TUM Munich, DE

FACULTY OF MEDICINE (ME1660)

October 2022 – March 2023

- Instructed hands-on computational genomics and mentored students in complex trait analysis.
- Developed and refined interactive JupyterLab course materials to optimize practical workshop flow.
- Taught UNIX, GWAS QC, association testing, meta-analysis, and polygenic risk score methods.

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### Genomic Data Science Intern

Helmholtz Munich, DE

INSTITUTE OF TRANSLATIONAL GENOMICS

October 2019 – April 2020

- Analyzed multi-center GWAS data from joint replacement cohorts to identify novel genetic risk factors.
- Harmonized Illumina exome-array datasets, resolving cross-center batch effects via QC workflows.
- Engineered GWAS sample- and variant-level QC pipeline in PLINK to eliminate technical biases.
- Executed large-scale genotype imputation via the Michigan Imputation Server using the HRC reference panel.

## Education

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Technical University of Munich (TUM) - Experimental Medicine

Munich, DE

PH.D. / DR.RER.NAT. IN SCHOOL OF MEDICINE (MGC)

Dec 2024

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Thessaly University - Computer Science and Biomedical Informatics

Lamia, GR

MSC. IN INFORMATICS AND COMPUTATIONAL BIOMEDICINE

April 2020

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Piraeus University - Department of Statistics and Insurance Science

Athens, GR

PTYCHION (4-YEAR BSC.) IN STATISTICS AND ACTUARIAL SCIENCE

February 2018

## Technical Experience

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Platform/API	Queryable data catalogs and analytical storage: <a href="#">Apache Iceberg</a> , <a href="#">Parquet</a> , partitioning, schema evolution. Analytical SQL and in-process analytics: <a href="#">SQL</a> , <a href="#">DuckDB</a> . Interactive apps with <a href="#">RShiny</a> + <a href="#">Neo4j</a> for graph exploration and hypothesis generation. REST query APIs for gene/variant/phenotype lookups and catalog search/querying. REST services with <a href="#">Flask</a> for geospatial routing and mapping workflows. Data validation and monitoring for production pipelines (QC modules, checks, runbooks).
Engineering	<a href="#">Python</a> , <a href="#">R</a> , <a href="#">Bash</a> , <a href="#">Linux</a> for large-scale data processing and workflow automation. Reproducible workflows, containerization, and cloud execution. Knowledge graph / semantic web tooling: <a href="#">Protégé</a> , <a href="#">RDF</a> , <a href="#">SPARQL</a> . Certified Oracle MySQL Developer (1z0-882). Version control with <a href="#">GitHub</a> ; documentation with <a href="#">LaTeX</a> and <a href="#">RMarkdown</a> . AI-assisted development with <a href="#">Cursor</a> , <a href="#">Copilot</a> , and LLM-based tooling.
Genetics	GWAS, PheWAS, eQTL/ct-eQTL mapping, WGS, RNA-seq, variant calling, Illumina array processing. Tools: <a href="#">Regenie</a> , <a href="#">PLINK</a> , <a href="#">LDAK</a> , <a href="#">Bioconductor</a> , <a href="#">MendelianRandomization</a> , <a href="#">Coloc</a> . PRS construction, burden testing, fine-mapping, conditional analysis, annotation integration. Multi-omics integration (genomic, transcriptomic, proteomic, regulatory). Biobanks: GTEx, PsychEncode, UK Biobank, All of Us, FinnGen, MVP.
Logic	Linear mixed models (GRM/variance components), GLMs; heteroskedasticity-aware modeling. Empirical inference: genotype permutations, simulations, likelihood-ratio tests; model evaluation. Multiple testing control: BH/q-value FDR and Bonferroni at tissue/study-wide levels. Causal follow-up: Bayesian colocalization, conditional/mediation analysis, Mendelian randomization. Feature engineering, PCA, ensemble learning for gene prioritization.
Cloud	Cloud-based genomic workflows with <a href="#">DNAAnexus</a> , <a href="#">AWS/S3</a> , and <a href="#">Docker</a> . Parallel execution, resumable runs, and cost-aware scaling for biobank-scale analyses.

## Scientific Publications

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Interplay between mitochondrial and nuclear DNA in gene expression regulation. [Biorxiv](#)

Xenofon Giannoulis, SIMON WENGERT, FLORIN RATAJCZAK, MATTHIAS HEINIG, NA CAI.

UNDER REVIEW AT NATURE COMMUNICATIONS.

Tissue-Specific mtDNA Heteroplasmy Linked to Aging and Gene Expression. [Biorxiv](#)

SIMON WENGERT, Xenofon Giannoulis, PETER KREITMAIER, HOLGER PROKISCH, PAOLO CASALE, MATTHIAS HEINIG, NA CAI.

UNDER REVIEW AT NATURE COMMUNICATIONS.