Niek de Klein

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- Scholar.google.com/citations?user=QNwjBnwAAAAJ&hl

Education

May 2015 - Sep 2021

■ PhD Functional Genomics

Genetics Department of University Medical Center Groningen, Groningen, the Netherlands.

Supervisor Prof. Lude Franke

Thesis title Genetic regulation of gene expression in brain and blood DOI 10.33612/diss.178049845

Sep 2012 - Aug 2014

M. Sc. Bioinformatics

Vrije Universiteit, Amsterdam, Netherlands **grade** 8.6/10 (Cum laude)

Sep 2008 – Aug 2012

■ B. Sc. Bioinformatics

HAN University of Applied Sciences, Nijmegen, Netherlands grade 8.2/10

Employment

Sep 2020 - Aug 2023

■ Postdoctoral researcher

Company Wellcome Sanger, Cambridge, United Kingdom Lab Joint post-doc in labs of Dr. Emma Davenport and Dr. Gosia Trynka Project For my postdoctoral research I use genotypes, single cell RNA-seq, CITE-seq, and vdj-seq data from individuals with Systemic Lupus Erythematosus to study clinical heterogeneity and the role of GWAS variants in regulating the immune system.

Skills

refer to publications, see Research Publications section.

Genomics/Genetics

Single cell

■ Experience with computational analysis of single cell RNA-seq, CITE-seq, and vdj-seq.

QTLs

Experience with downstream analysis of eQTLs (2, 9, 12) as well as developing new software for calculating interaction eQTLs (8 and github.com/npklein/systemsgenetics/tree/master/Decon2)

Differential expression

Experience running differential expression analysis (10).

(G/E)WAS

Experience running EWAS as part of large consortium (4), 5).

Method development

Experience developing new methods for gene identification (17 and 18), gene ID mapping between species (16), improving protein identification from mass spectrometry (See B.Sc. and M.sc. internships), and identifying cell type specific eQTL effects in bulk data (8).

Coding

Coding

■ Most experienced in R Bash, Python. Experience in Java. Code on github.com/npklein. Code written during current employment is in a private repository, can share recent examples on request.

Skills (continued)

Databases

Experience with MysqL and sqLite

General

Writing

Experience writing scientific articles (See Research Publications below).

Teaching and supervision

Experience giving workshops and lectures (See Teaching below). Experience supervising students (See Teaching below for details.)

Presenting work

Regular presentations at group and departmental level. Presenting monthly for stakeholders from industry on project progress as part of OpenTargets private-public collaboration.

Languages

■ Strong reading, writing and speaking competencies for English and Dutch (native).

Internships

During M.Sc.

Jan – Jun 2014

■ Connecting multiple gene expression signatures with candidate drugs for boosting heart regeneration potential.

Supervisor Dr. Francisco Azuaje

Institute CRD-Santeé, Luxembourg, Luxembourg

Grade 9/10

https://github.com/npklein/drugFinder

Aug - Dec 2013

Clustering of MS/MS spectra to improve peptide and protein identification. Supervisor Dr. Tham Pham

Institute Cancer Center Amsterdam, Amsterdam, Netherlands

Grade 9/10

During B.Sc.

Feb – Jun 2012

■ Methods to assess the reproducibility and coverage of tandem mass spectrometry data in phosphoproteomics experiments.

Supervisor Dr. David Martin

Institute Wellcome Trust, Dundee, Scotland

Grade 9/10

https://github.com/npklein/pyMSA

Sep - Dec 2010

■ How widespread are microProteins and how did they evolve?

Supervisor Dr. Sue Rhee

Institute Carnegie Institue for Life Sciences, Stanford, USA

Grade 8/10

https://github.com/npklein/miP3

Teaching

Courses and seminars

16 Jun 2022

Lecture basics of eQTL analysis and colocalisations
Wellcome Sanger departmental retreat

Teaching (continued)

Lecture on gene expression + workshop differential gene expression and pathway analysis

Data Science for Life Sciences Master students of the Hanze University of Applied Sciences Groningen

Course material at npklein.github.io/Teaching-modules/.

Sep – Oct 2017 ■ Introduction to Biomolecular Sciences

Co-coordinated, created, and supervised various genetics workshops. Biomolecular Science Master students of the Rijksuniversiteit Groningen Course material at npklein.github.io/iBMS.

26 Aug 2016 ■ Invited Talk + Workshop Genotyping from RNAseq

Lecture + workshop on how to call genotypes from RNAseq data at the Estonian Genome Center.

22 Jun 2016 Enriching biobanks using genotype data BIKE Summer School seminar

Student (co)-supervision

Mar 2021 − **Z**iying Ke

PhD student

Metabolic pathway perturbation in single cells

May 2022 − Jul 2022 ■ Haerin Jang

PhD rotation project

Drug target identification through single cell eQTL mapping of immune response regulation in health and SLE

Mar 2021 – May 2021 📕 Isaac Garcia-Salinas

PhD rotation project

Determining putative causal genes of neuropsychiatric lupus through genotype and brain gene expression data

Jan 2020 – Jun 2020 Martijn Vochteloo

Master thesis

Investigation of Cell Type Mediated eQTL Regulatory Effects in Psychiatric and Neurodegenerative Diseases

Apr 2019 – Jun 2019 ■ Fabian VogelPohl

Bachelor internship

Unraveling the brain specific molecular network of the SCGE gene, the major gene for myoclonus dystonia.

Sep 2018 – Apr 2019 ■ Omar El Garwany

Master thesis

Brain eQTLs show more enrichment for neuro-psychotic diseases than blood eQTLs

Teaching (continued)

Feb - Jul 2017

Anne Tjallingii

M.Sc. internship
Influence of a diabetes mellitus type II on the susceptibility of getting

tuberculosis

Jan – Jun 2016

Carlos Urzua

M.Sc. internship

Allele-specific signals in gluten sensitive t-cell clones

Misc

Jun 2022

Marking students for M.Sc. course Genomics of complex disease Cambridge University, United Kingdom

Aug 2020, 2021

■ External examiner for various bachelor thesis projects H.A.N., Nijmgen, the Netherlands

Grants

07 Mar 2016

One tool to interpret and assess impact of genetic variants and predict disease-causing genes using the BBMRI - BIOS data (Co-Applicant)
 € 60,000

Miscellaneous Experience

Conference poster presentations

17 October 2019

■ eQTL analysis in brain cortex samples of 3,833 individuals and 31,684 blood samples reveals distinct regulatory effects of disease-associated genetic variants

ASHG (Poster)

17 June 2018

■ Allele specific expression identifies rare variants as cause for extreme allelic imbalance

ESHG (Poster)

25 May 2016

■ Genotyping of all public RNA-sequencing data for large scale trans-QTL and ASE studies

ESHG (Poster)

8 Sep 2014

■ Connecting multiple gene expression signatures with candidate drugs for boosting heart regeneration potential ECCB (Poster)

Other

Okt 2019 – Dec 2019

Research visit at Biogen - including invited talk

Aug 2018

■ Leena Peltonen Summer School Summer school

Miscellaneous Experience (continued)

Jul 2016 Microsoft Research PhD Summer School
Summer school

Research Publications (Reverse chronological order)

Journal Articles

- Boulogne, F., Claus, L., Wiersma, H., Oelen, R., Schukking, F., de <u>Klein</u>, N., Li, S., Westra, H.-J., van der Zwaag, B., van Reekum, F., et al. (2023). Kidneynetwork: Using kidney-derived gene expression data to predict and prioritize novel genes involved in kidney disease. *medRxiv*, in press European Journal of Human Genetics 2023.
- de <u>Klein*</u>, N., Tsai*, E. A., Vochteloo*, M., Baird, D., Huang, Y., Chen, C.-Y., van Dam, S., Deelen, P., Bakker, O. B., El Garwany, O., et al. (2023). Brain expression quantitative trait locus and network analysis reveals downstream effects and putative drivers for brain-related diseases. *bioRxiv in press Nature Genetics 2023*.
- Bakker, O. B., Ramirez-Sanchez, A. D., Borek, Z. A., de <u>Klein</u>, N., Li, Y., Modderman, R., Kooy-Winkelaar, Y., Johannesen, M. K., Matarese, F., Martens, J. H., et al. (2021). Potential impact of celiac disease genetic risk factors on t cell receptor signaling in gluten-specific cd4+ t cells. *Scientific reports*, 11(1), 1–15.
- 4 Karabegovic, I., Portilla-Fernandez, E., Li, Y., Ma, J., Maas, S. C., Sun, D., Hu, E. A., Kuhnel, B., Zhang, Y., Ambatipudi, S., et al. (2021). Epigenome-wide association meta-analysis of dna methylation with coffee and tea consumption. *Nature communications*, 12(1), 1–13.
- Schlosser, P., Tin, A., Matias-Garcia, P. R., Thio, C. H., Joehanes, R., Liu, H., Weihs, A., Yu, Z., Hoppmann, A., Grundner-Culemann, F., et al. (2021). Meta-analyses identify dna methylation associated with kidney function and damage. *Nature communications*, 12(1), 1–16.
- Van Rheenen, W., Van Der Spek, R. A., Bakker, M. K., Van Vugt, J. J., Hop, P. J., Zwamborn, R. A., de <u>Klein</u>, N., Westra, H.-J., Bakker, O. B., Deelen, P., et al. (2021). Common and rare variant association analyses in amyotrophic lateral sclerosis identify 15 risk loci with distinct genetic architectures and neuron-specific biology. *Nature genetics*, 53(12), 1636–1648.
- Zhang, Z., van Dijk, F., de <u>Klein</u>, N., van Gijn, M. E., Franke, L. H., Sinke, R. J., Swertz, M. A., & van der Velde, K. J. (2021). Feasibility of predicting allele specific expression from dna sequencing using machine learning. *Scientific reports*, 11(1), 1–11.
- Aguirre-Gamboa*, R., de <u>Klein*</u>, N., di Tommaso*, J., Claringbould, A., van der Wijst, M. G., de Vries, D., Brugge, H., Oelen, R., Vosa, U., Zorro, M. M., et al. (2020). Deconvolution of bulk blood eqtl effects into immune cell subpopulations. *BMC bioinformatics*, 21(1), 1–23.
- de <u>Klein*</u>, N., van Dijk*, F., Deelen, P., Urzua, C. G., Claringbould, A., Vosa, U., Verlouw, J. A., Monajemi, R., AC't Hoen, P., Sinke, R. J., et al. (2020). Imbalanced expression for predicted high-impact, autosomal-dominant variants in a cohort of 3,818 healthy samples. *bioRxiv*.
- Rodius, S., de <u>Klein</u>, N., Jeanty, C., Sanchez-Iranzo, H., Crespo, I., Ibberson, M., Xenarios, I., Dittmar, G., Mercader, N., Niclou, S. P., et al. (2020). Fisetin protects against cardiac cell death through reduction of ros production and caspases activity. *Scientific reports*, 10(1), 1–12.
- Parmar, P., Lowry, E., Cugliari, G., Suderman, M., Wilson, R., Karhunen, V., Andrew, T., Wiklund, P., Wielscher, M., Guarrera, S., et al. (2018). Association of maternal prenatal smoking gfi1-locus and cardio-metabolic phenotypes in 18,212 adults. *EBioMedicine*, 38, 206–216.

- Claringbould*, A., de <u>Klein*</u>, N., & Franke, L. (2017). The genetic architecture of molecular traits. *Current Opinion in Systems Biology*, 1, 25–31.
- Zhernakova, D. V., Deelen, P., Vermaat, M., Van Iterson, M., Van Galen, M., Arindrarto, W., Van't Hof, P., Mei, H., Van Dijk, F., Westra, H.-J., et al. (2017). Identification of context-dependent expression quantitative trait loci in whole blood. *Nature genetics*, 49(1), 139–145.
- Graham, D. B., Lefkovith, A., Deelen, P., de <u>Klein</u>, N., Varma, M., Boroughs, A., Desch, A. N., Ng, A. C., Guzman, G., Schenone, M., et al. (2016). Tmem258 is a component of the oligosaccharyltransferase complex controlling er stress and intestinal inflammation. *Cell reports*, 17(11), 2955–2965.
- Rodius, S., Androsova, G., Gotz, L., Liechti, R., Crespo, I., Merz, S., Nazarov, P. V., de <u>Klein</u>, N., Jeanty, C., Gonzalez-Rosa, J. M., et al. (2016). Analysis of the dynamic co-expression network of heart regeneration in the zebrafish. *Scientific reports*, 6(1), 1–12.
- de <u>Klein</u>, N., Ibberson, M., Crespo, I., Rodius, S., & Azuaje, F. (2015). A gene mapping bottleneck in the translational route from zebrafish to human. *Frontiers in Genetics*, 5, 470.
- de <u>Klein</u>, N., Magnani, E., Banf, M., & Rhee, S. Y. (2015). Microprotein prediction program (mip3): A software for predicting microproteins and their target transcription factors. *International journal of genomics*, 2015.
- Magnani, E., de <u>Klein</u>, N., Nam, H.-I., Kim, J.-G., Pham, K., Fiume, E., Mudgett, M. B., & Rhee, S. Y. (2014). A comprehensive analysis of microproteins reveals their potentially widespread mechanism of transcriptional regulation. *Plant physiology*, 165(1), 149–159.

Books and Chapters

Fiume, E., de <u>Klein</u>, N., Rhee, S. Y., & Magnani, E. (2016). A framework for discovering, designing, and testing microproteins to regulate synthetic transcriptional modules (Vol. 1482). Humana Press, New York, NY.

References

Available on request

^{* =} shared first author