CURRICULUM VITAE

Kasper Daniel Hansen

PERSONAL DATA

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Johns Hopkins Bloomberg School of Public Health

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EDUCATION AND TRAINING

Degrees

Ph.D. 2009 University of California, Berkeley

Biostatistics with a designated emphasis in Computational and Genomic Biology

Advisor: Sandrine Dudoit

Cand. Scient. 2002 University of Copenhagen

Statistics

Advisor: Martin Jacobsen

B. Sc. 1998 University of Copenhagen

Statistics and mathematics

Postdoctoral Training

2009–2012 Department of Biostatistics

Johns Hopkins Bloomberg School of Public Health

Advisor: Rafael Irizarry

Visiting

2018 Departments of Statistics and Biology

University of Copenhagen

2004–2005 Department of Biostatistics

University of California, Berkeley

PROFESSIONAL EXPERIENCE

2018–Present Associate Professor

Department of Biostatistics

Nathans-McKusick Institute of Genetic Medicine

Johns Hopkins University

2012–2018 Assistant Professor

Department of Biostatistics

Nathans-McKusick Institute of Genetic Medicine

Johns Hopkins University

2009–2012 Postdoctoral Fellow, Department of Biostatistics

Johns Hopkins University

2002–2004 Research Assistant, Department of Biostatistics

University of Copenhagen, Denmark.

PROFESSIONAL ACTIVITIES

Professional Memberships

American Society of Human Genetics American Statistical Association

Project Development

2012–Present Member of the Bioconductor Technical Advisory Board.

EDITORIAL ACTIVITIES

Editorial Board Membership

Gateway advisor for the Bioconductor Gateway at F1000Research.

Served as referee for

Annals of Applied Statistics

Bioinformatics

Biometrics

Biostatistics

BMC Bioinformatics

F1000Research

Genome Biology

Genome Research

International Journal of Biostatistics

Journal of the American Medical Association (JAMA) Journal of the American Statistical Association (JASA)

Nature Biotechnology

Nature Communications

Nature Ecology and Evolution

Nature Methods

Nature Reviews Genetics

Nucleic Acids Research

PLOS Biology

PLOS Genetics

PLOS ONE

Proceedings of the National Academy of Sciences (PNAS)

RNA

Statistical Applications in Genetics and Molecular Biology

Statistics in Medicine

Review of Proposals

Joint NIH and NSF BIGDATA initiative review panel (2012) Israeli Science Foundation (2019)

HONORS AND AWARDS

2010	Second prize at the MGED poster competition (out of around 50)
2007	Third prize at the Computational and Genomic Biology student retreat
	poster competition
2007	Reshetko Family Scholarship, UC Berkeley
2005	William V. Power Top-off Graduate Award, UC Berkeley
2005	William V. Power Graduate Award, UC Berkeley

Significant awards to trainees:

2014 Jean-Philippe Fortin:

John van Ryzin award for best student paper submitted to ENAR.

PUBLICATIONS

Journal Articles (peer reviewed)

boldface indicates a member of my lab

^{*} indicates equal contributions

[†] indicates corresponding author(s) (if not the senior author)

- [1] J. A. Fahrner[†], W.-Y. Lin, R. C. Riddle, L. Boukas, V. B. DeLeon, S. Chopra, S. E. Lad, T. R. Luperchio, K. D. Hansen, and H. T. Bjornsson[†]. "Precocious chondrocyte differentiation disrupts skeletal growth in Kabuki syndrome mice". JCI Insight (2019). DOI: 10.1172/jci.insight.129380.
- [2] G. A. Carosso, L. Boukas, J. J. Augustin, H. N. Nguyen, B. L. Winer, G. H. Cannon, J. D. Robertson, L. Zhang, K. D. Hansen, L. A. Goff, and H. T. Bjornsson. "Precocious neuronal differentiation and disrupted oxygen responses in Kabuki syndrome". JCI Insight (2019). DOI: 10.1172/jci.insight.129375.
- [3] L. Myint, R. Wang, L. Boukas, K. D. Hansen, L. A. Goff, and D. G. Avramopoulos. "A screen of 1,049 schizophrenia and 30 Alzheimer's-associated variants for regulatory potential". *American Journal of Medical Genetics Part B: Neuropsychiatric Genetics* (2019). In Press. DOI: 10.1002/ajmg.b.32761.
- [4] P. Wulfridge, B. Langmead, A. P. Feinberg, and **K. D. Hansen**. "Analyzing whole genome bisulfite sequencing data from highly divergent genotypes". *Nucleic Acids Research* (2019). In Press. DOI: 10.1093/nar/gkz674.
- [5] L. Myint, D. G. Avramopoulos, L. A. Goff, and K. D. Hansen. "Linear models enable powerful differential activity analysis in massively parallel reporter assays". BMC Genomics 20 (2019), p. 209. DOI: 10.1186/s12864-019-5556-x.
- [6] L. Boukas, J. M. Havrilla, P. F. Hickey, A. R. Quinlan, H. T. Bjornsson[†], and K. D. Hansen[†]. "Coexpression patterns define epigenetic regulators associated with neurological dysfunction". Genome Research 29.4 (2019). In press, pp. 532–542. DOI: 10.1101/gr.239442. 118.
- [7] L. F. Rizzardi^{*}, **P. F. Hickey**^{*}, V. R. DiBlasi, R. Tryggvadottir, C. M. Callahan, A. Idrizi, **K. D. Hansen**[†], and A. P. Feinberg[†]. "Neuronal brain region-specific DNA methylation and chromatin accessibility are associated with neuropsychiatric disease heritability". *Nature Neuroscience* 22.2 (2019), pp. 307–316. DOI: 10.1038/s41593-018-0297-8.
- [8] V. Gaysinskaya, B. F. Miller, C. De Luca, G. W. van der Heijden, K. D. Hansen, and A. Bortvin. "Transient reduction of DNA methylation at the onset of meiosis in male mice". Epigenetics & Chromatin 11.1 (2018), p. 15. DOI: 10.1186/s13072-018-0186-0.
- [9] W. T. Barrington, P. Wulfridge, A. E. Wells, C. M. Rojas, S. Y. Howe, A. Perry, K. Hua, M. A. Pellizzon, K. D. Hansen, B. H. Voy, B. J. Bennett, D. Pomp, A. P. Feinberg, and D. W. Threadgill. "Improving metabolic health through precision dietetics in mice". Genetics 208.1 (2018), pp. 399-417. DOI: 10.1534/genetics.117.300536.
- [10] M. Ramos, L. Schiffer, A. Re, R. Azhar, A. Basunia, C. Rodriguez, T. Chan, P. Chapman, S. R. Davis, D. Gomez-Cabrero, A. C. Culhane, B. Haibe-Kains, K. D. Hansen, H. Kodali, M. S. Louis, A. S. Mer, M. Riester, M. Morgan, V. Carey, and L. Waldron. "Software for the Integration of Multiomics Experiments in Bioconductor". Cancer Research 77.21 (2017), e39–e42. DOI: 10.1158/0008-5472.CAN-17-0344.
- [11] L. Collado-Torres*, A. Nellore*, K. Kammers, S. E. Ellis, M. A. Taub, **K. D. Hansen**, A. E. Jaffe[†], B. Langmead[†], and J. T. Leek[†]. "Reproducible RNA-seq analysis using recount2". *Nature Biotechnology* 35.4 (2017), pp. 319–321. DOI: 10.1038/nbt.3838.

- [12] S. C. Zheng, S. Beck, A. E. Jaffe, D. C. Koestler, K. D. Hansen, A. E. Houseman, R. A. Irizarry, and A. E. Teschendorff. "Correcting for cell-type heterogeneity in epigenome-wide association studies: revisiting previous analyses". *Nature Methods* 14.3 (2017), pp. 216–217. DOI: 10.1038/nmeth.4187.
- [13] L. Myint, A. Kleensang, L. Zhao, T. Hartung, and K. D. Hansen. "Joint bounding of peaks across samples improves differential analysis in mass spectrometry-based metabolomics". Analytical Chemistry 89.6 (2017), pp. 3517–3523. DOI: 10.1021/acs.analchem.6b04719.
- [14] **J.-P. Fortin**, T. Triche Jr, and **K. D. Hansen**. "Preprocessing, normalization and integration of the Illumina HumanMethylationEPIC array with minfi". *Bioinformatics* 33.4 (2017), pp. 558–560. DOI: 10.1093/bioinformatics/btw691.
- [15] J. S. Benjamin, G. Pilarowski, G. Carosso, L. Zhang, J. Farner, D. L. Huso, L. A. Goff, H. Vernon, K. D. Hansen, and H. T. Bjornsson. "A ketogenic diet rescues hippocampal memory defects in a mouse model of Kabuki syndrome". Proceedings of the National Academy of Sciences 114.1 (2017), pp. 125–130. DOI: 10.1073/pnas.1611431114.
- [16] A. Nellore, A. E. Jaffe, J.-P. Fortin, J. Alquicira-Hernández, L. Collado-Torres, S. Wang, R. A. Phillips, N. Karbhari, K. D. Hansen, B. Langmead[†], and J. T. Leek[†]. "Human splicing diversity and the extent of unannotated splice junctions across human RNA-seq samples on the Sequence Read Archive". Genome Biology 17 (2016), p. 266. DOI: 10.1186/s13059-016-1118-6.
- [17] S. V. Andrews*, C. Ladd-Acosta*, A. P. Feinberg, **K. D. Hansen**, and M. D. Fallin. ""Gap hunting" to to characterize clustered probe signals in Illumina methylation array data". *Epigenetics and Chromatin* 9 (2016), p. 56. DOI: 10.1186/s13072-016-0107-z.
- [18] X. Li, Y. Liu, T. Salz, **K. D. Hansen**, and A. P. Feinberg. "Whole genome analysis of the methylome and hydroxymethylome in normal and malignant lung and liver". *Genome Research* 26.12 (2016), pp. 1730–1741. DOI: 10.1101/gr.211854.116.
- [19] A. Nellore[†], C. Wilks, **K. D. Hansen**, J. T. Leek, and B. Langmead[†]. "Rail-dbGaP: analyzing dbGaP-protected data in the cloud with Amazon Elastic MapReduce". *Bioinformatics* 32.16 (2016), pp. 2551–2553. DOI: 10.1093/bioinformatics/btw177.
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- [24] **J.-P. Fortin** and **K. D. Hansen**. "Reconstructing A/B compartments as revealed by Hi-C using long-range correlations in epigenetic data". *Genome Biology* 16 (2015), p. 180. DOI: 10.1186/s13059-015-0741-y.
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- [26] X. Hong*, K. Hao*, C. Ladd-Acosta*, K. D. Hansen, H.-J. Tsai, X. Liu, X. Xu, T. A. Thornton, D. Caruso, C. A. Keet, Y. Sun, G. Wang, W. Luo, R. Kumar, R. Fuleihan, A. M. Singh, J. S. Kim, R. E. Story, R. S. Gupta, P. Gao, Z. Chen, S. O. Walker, T. R. Bartell, T. H. Beaty, M. D. Fallin, R. Schleimer, P. G. Holt, K. C. Nadeau, R. A. Wood, J. A. Pongracic, D. E. Weeks, and X. Wang. "Genome-wide association study identifies peanut allergy-specific loci and evidence of epigenetic mediation in US children." Nature Communications 6 (2015), p. 6304. DOI: 10.1038/ncomms7304.
- [27] W. Huber[†], V. J. Carey, R. Gentleman, S. Anders, M. Carlson, B. S. Carvalho, H. C. Bravo, S. Davis, L. Gatto, T. Girke, R. Gottardo, F. Hahne, K. D. Hansen, R. A. Irizarry, M. Lawrence, M. I. Love, J. MacDonald, V. Obenchain, A. K. Oleś, H. Pagès, A. Reyes, P. Shannon, G. K. Smyth, D. Tenenbaum, L. Waldron, and M. Morgan. "Orchestrating high-throughput genomic analysis with Bioconductor". Nature Methods 12.2 (2015), pp. 115–121. DOI: 10.1038/nmeth.3252.
- [28] J.-P. Fortin, A. Labbe, M. Lemire, B. W. Zanke, T. J. Hudson, E. J. Fertig, C. M. Greenwood, and K. D. Hansen. "Functional normalization of 450k methylation array data improves replication in large cancer studies". Genome Biology 15 (2014), p. 503. DOI: 10.1186/s13059-014-0503-2.
- [29] M. J. Ziller, **K. D. Hansen**, A. Meissner[†], and M. J. Aryee[†]. "Coverage recommendations for methylation analysis by whole-genome bisulfite sequencing". *Nature Methods* 12.3 (2015), pp. 230–232. DOI: 10.1038/nmeth.3152.
- [30] H. T. Bjornsson*†, J. S. Benjamin*, L. Zhang, J. Weissman, E. E. Gerber, Y.-C. Chen, R. G. Vaurio, M. C. Potter, **K. D. Hansen**, and H. C. Dietz. "Histone deacetylase inhibition rescues structural and functional brain deficits in a mouse model of Kabuki syndrome." *Science Translational Medicine* 6.256 (2014), 256ra135. DOI: 10.1126/scitranslmed.3009278.
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- [32] A. C. Frazee, S. Sabunciyan, **K. D. Hansen**, R. A. Irizarry, and J. T. Leek. "Differential expression analysis of RNA-seq data at single-base resolution." *Biostatistics* 15.3 (2014), pp. 413–426. DOI: 10.1093/biostatistics/kxt053.

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- [35] M. L. Smith[†], K. A. Baggerly, H. Bengtsson, M. E. Ritchie, and **K. D. Hansen**[†]. "illuminaio: An open source IDAT parsing tool for Illumina microarrays". F1000Research 264.2 (2013). DOI: 10.12688/f1000research.2-264.v1.
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- [37] **K. D. Hansen**[†], B. Langmead[†], and R. A. Irizarry[†]. "BSmooth: from whole genome bisulfite sequencing reads to differentially methylated regions". *Genome Biology* 13 (2012), R83. DOI: 10.1186/gb-2012-13-10-r83.
- [38] B. R. Herb, F. Wolschin, **K. D. Hansen**, M. J. Aryee, B. Langmead, R. Irizarry, G. V. Amdam[†], and A. P. Feinberg[†]. "Reversible switching between epigenetic states in honeybee behavioral subcastes." *Nature Neuroscience* 15.10 (2012), pp. 1371–1373. DOI: 10.1038/nn. 3218.
- [39] J. Tung[†], L. B. Barreiro, Z. P. Johnson, K. D. Hansen, V. Michopoulos, D. Toufexis, K. Michelini, M. E. Wilson, and Y. Gilad[†]. "Social environment is associated with gene regulatory variation in the rhesus macaque immune system." *Proceedings of the National Academy of Sciences* 109.17 (2012), pp. 6490–6495. DOI: 10.1073/pnas.1202734109.
- [40] S. Munshaw, H. S. Hwang, M. Torbenson, J. Quinn, K. D. Hansen, J. Astemborski, S. H. Mehta, S. C. Ray, D. L. Thomas, and A. Balagopal. "Laser captured hepatocytes show association of butyrylcholinesterase gene loss and fibrosis progression in hepatitis C-infected drug users." Hepatology 56.2 (2012), pp. 544–554. DOI: 10.1002/hep.25655.
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- [42] K. D. Hansen*, W. Timp*, H. C. Bravo*, S. Sabunciyan*, B. Langmead*, O. G. McDonald, B. Wen, H. Wu, Y. Liu, D. Diep, E. Briem, K. Zhang, R. A. Irizarry†, and A. P. Feinberg†. "Increased methylation variation in epigenetic domains across cancer types". *Nature Genetics* 43.8 (2011), pp. 768–775. DOI: 10.1038/ng.865.
- [43] **K. D. Hansen**, Z. Wu, R. A. Irizarry[†], and J. T. Leek[†]. "Sequencing technology does not eliminate biological variability". *Nature Biotechnology* 29.7 (2011), pp. 572–573. DOI: 10.1038/nbt.1910.
- [44] A. N. Brooks, L. Yang, M. O. Duff, **K. D. Hansen**, J. W. Park, S. Dudoit, S. E. Brenner[†], and B. R. Graveley[†]. "Conservation of an RNA regulatory map between Drosophila and mammals". *Genome Research* 21.2 (2011), pp. 193–202. DOI: 10.1101/gr.108662.110.

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Journal Articles, Consortia member (peer reviewed)

- [54] eGTEx Project. "Enhancing GTEx by bridging the gaps between genotype, gene expression, and disease". *Nature Genetics* 12 (2017), pp. 1664–1670. DOI: 10.1038/ng.3969.
- [55] A. Saha, Y. Kim, A. D. H. Gewirtz, B. Jo, C. Gao, I. C. McDowell, GTEx Consortium, B. E. Engelhardt, and A. Battle. "Co-expression networks reveal the tissue-specific regulation of transcription and splicing". Genome Research 27.11 (2017), pp. 1843–1858. DOI: 10.1101/gr. 216721.116.
- [56] F. Yang, J. Wang, GTEx Consortium, B. L. Pierce, and L. S. Chen. "Identifying cis-mediators for trans-eQTLs across many human tissues using genomic mediation analysis". *Genome Research* 27.11 (2017), pp. 1859–1871. DOI: 10.1101/gr.216754.116.

- [57] X. Li, Y. Kim, E. K. Tsang, J. R. Davis, F. N. Damani, C. Chiang, G. T. Hess, Z. Zappala, B. J. Strober, A. J. Scott, A. Li, A. Ganna, M. C. Bassik, J. D. Merker, GTEx Consortium, Laboratory, Data Analysis & Coordinating Center (LDACC)—Analysis Working Group, Statistical Methods groups—Analysis Working Group, Enhancing GTEx (eGTEx) groups, NIH Common Fund, NIH/NCI, NIH/NHGRI, NIH/NIMH, NIH/NIDA, Biospecimen Collection Source Site—NDRI, Biospecimen Collection Source Site—RPCI, Biospecimen Core Resource—VARI, Brain Bank Repository—University of Miami Brain Endowment Bank, Leidos Biomedical—Project Management, ELSI Study, Genome Browser Data Integration & Visualization—EBI, Genome Browser Data Integration & Visualization—UCSC Genomics Institute, University of California Santa Cruz, I. M. Hall, A. Battle, and S. B. Montgomery. "The impact of rare variation on gene expression across tissues". Nature 550.7675 (2017), pp. 239–243. DOI: 10.1038/nature24267.
- [58] M. H. Tan, Q. Li, R. Shanmugam, R. Piskol, J. Kohler, A. N. Young, K. I. Liu, R. Zhang, G. Ramaswami, K. Ariyoshi, A. Gupte, L. P. Keegan, C. X. George, A. Ramu, N. Huang, E. A. Pollina, D. S. Leeman, A. Rustighi, Y. P. S. Goh, GTEx Consortium, Laboratory, Data Analysis & Coordinating Center (LDACC)—Analysis Working Group, Statistical Methods groups—Analysis Working Group, Enhancing GTEx (eGTEx) groups, NIH Common Fund, NIH/NCI, NIH/NHGRI, NIH/NIMH, NIH/NIDA, Biospecimen Collection Source Site—NDRI, Biospecimen Collection Source Site—RPCI, Biospecimen Core Resource—VARI, Brain Bank Repository—University of Miami Brain Endowment Bank, Leidos Biomedical—Project Management, ELSI Study, Genome Browser Data Integration & Visualization—EBI, Genome Browser Data Integration & Visualization—UCSC Genomics Institute, University of California Santa Cruz, A. Chawla, G. Del Sal, G. Peltz, A. Brunet, D. F. Conrad, C. E. Samuel, M. A. O'Connell, C. R. Walkley, K. Nishikura, and J. B. Li. "Dynamic landscape and regulation of RNA editing in mammals". Nature 550.7675 (2017), pp. 249–254. DOI: 10.1038/nature24041.
- [59] GTEx Consortium, Laboratory, Data Analysis & Coordinating Center (LDACC)—Analysis Working Group, Statistical Methods groups—Analysis Working Group, Enhancing GTEx (eGTEx) groups, NIH Common Fund, NIH/NCI, NIH/NHGRI, NIH/NIMH, NIH/NIDA, Biospecimen Collection Source Site—NDRI, Biospecimen Collection Source Site—RPCI, Biospecimen Core Resource—VARI, Brain Bank Repository—University of Miami Brain Endowment Bank, Leidos Biomedical—Project Management, ELSI Study, Genome Browser Data Integration & Visualization—EBI, Genome Browser Data Integration & Visualization—UCSC Genomics Institute, University of California Santa Cruz, Lead analysts: Laboratory, Data Analysis & Coordinating Center (LDACC): NIH program management: Biospecimen collection: Pathology: eQTL manuscript working group: A. Battle, C. D. Brown, B. E. Engelhardt, and S. B. Montgomery. "Genetic effects on gene expression across human tissues". Nature 550.7675 (2017), pp. 204–213. DOI: 10.1038/nature24277.
- [60] T. Tukiainen, A.-C. Villani, A. Yen, M. A. Rivas, J. L. Marshall, R. Satija, M. Aguirre, L. Gauthier, M. Fleharty, A. Kirby, B. B. Cummings, S. E. Castel, K. J. Karczewski, F. Aguet, A. Byrnes, GTEx Consortium, Laboratory, Data Analysis & Coordinating Center (LDACC)—Analysis Working Group, Statistical Methods groups—Analysis Working Group, Enhancing GTEx (eGTEx) groups, NIH Common Fund, NIH/NCI, NIH/NHGRI, NIH/NIMH, NIH/NIDA, Biospecimen Collection Source Site—NDRI, Biospecimen Collection Source Site—RPCI, Biospecimen Core Resource—VARI, Brain Bank Repository—University of Miami Brain Endowment Bank, Leidos Biomedical—Project Management, ELSI Study, Genome Browser Data Inte-

- gration & Visualization—EBI, Genome Browser Data Integration & Visualization—UCSC Genomics Institute, University of California Santa Cruz, T. Lappalainen, A. Regev, K. G. Ardlie, N. Hacohen, and D. G. MacArthur. "Landscape of X chromosome inactivation across human tissues". *Nature* 550.7675 (2017), pp. 244–248. DOI: 10.1038/nature24265.
- [61] modENCODE Consortium. "Identification of functional elements and regulatory circuits by Drosophila modENCODE". Science 330.6012 (2010), pp. 1787–97. DOI: 10.1126/science. 1198374.

Preprints (not peer reviewed)

- * indicates equal contributions
- † indicates corresponding author(s) (if not the senior author) **boldface** indicates a member of my lab

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PRACTICE ACTIVITIES

Software - Bioconductor Project

affxparser A package for parsing output files from Affymetrix microarrays using the Affymetrix Fusion SDK.

bnbc A package to normalize and remove unwanted variation in Hi-C data.

bsseq A package for analyzing whole-genome bisulfite sequencing data.

bumphunter A package implementing a general backend for the bumphunter approach.

cqn A package for normalizing RNA-seq data using the CQN algorithm.

Genominator A package implementing a SQLite based backend for genomic data, including sequencing and microarrays.

illuminaio A package for parsing output for Illumina microarrays. This package is not yet in Bioconductor release.

minfi A package for analysing Illumina's 450k DNA methylation microarray.

mpra A package to analysis massively parallel reporter assays (MPRA).

Rgraphviz A package for visualizing graphs using the Graphviz toolkit.

yamss A package for analyzing MS-based metabolomics experiments.

Software - Other

Myrna is a cloud computing tool for calculating differential gene expression in large RNA-seq datasets. Myrna uses Bowtie for short read alignment and R/Bioconductor for interval calculations, normalization, and statistical testing. These tools are combined in an automatic, parallel pipeline that runs in the cloud (Elastic MapReduce in this case) on a local Hadoop cluster, or on a single computer, exploiting multiple computers and CPUs wherever possible.