# Peter Francis Hickey

# Curriculum Vitae 30 June 2017

# Contact

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- Australian Citizen

# Education and research experience

- 2016-present: **Postdoctoral Fellow** Department of Biostatistics, Johns Hopkins Bloomberg School of Public Health, Baltimore, USA.
  - Advisor: Assistant Professor Kasper Hansen
- 2011-2015: PhD (Statistics) Department of Mathematics and Statistics, The University of Melbourne, Australia.
  - Thesis: The statistical analysis of high-throughput assays for studying DNA methylation
  - Advisors: Professor Terry Speed and Professor Peter Hall
- 2009-2015: Research Assistant Bioinformatics Division, Walter and Eliza Hall Institute, Melbourne, Australia.
  - Advisor: Professor Melanie Bahlo
- 2008-2009: Undergraduate Research Opportunities Program scholar Bioinformatics Division, Walter and Eliza Hall Institute, Melbourne, Australia.
  - Advisor: Professor Melanie Bahlo
- 2006-2009: Bachelor of Science (Mathematics and Statistics) with First Class Honours The University of Melbourne, Australia.
  - Thesis: X chromosome association testing in genome-wide association studies
  - Advisors: Professor Melanie Bahlo and Professor Richard Huggins

#### **Publications**

\*Indicates equal contributions

#### Journal articles

- [1] J. T. Hickey, P. F. Hickey, N. Maniar, R. G. Timmins, M. D. Williams, C. A. Pitcher and D. A. Opar. "A novel apparatus measuring knee flexor strength during various hamstring exercises: A reliability and retrospective study". In: *Journal of Orthopaedic & Sports Physical Therapy* (in press) (2017).
- [2] P. F. Hickey. "Representation and manipulation of genomic tuples in R". In: *The Journal of Open Source Software* 1.1 (May. 2016). DOI: 10.21105/joss.00020. URL: https://doi.org/10.21105/joss.00020.

- [3] A. Keniry, L. J. Gearing, N. Jansz, J. Liu, A. Z. Holik, **P. F. Hickey**, S. A. Kinkel, D. L. Moore, K. Breslin, K. Chen, R. Liu, C. Phillips, M. Pakusch, C. Biben, J. M. Sheridan, B. T. Kile, C. Carmichael, M. E. Ritchie, D. J. Hilton and M. E. Blewitt. "Setdb1-mediated H3K9 methylation is enriched on the inactive X and plays a role in its epigenetic silencing". In: *Epigenetics & Chromatin* 9 (May. 18, 2016), p. 16. ISSN: 1756-8935. DOI: 10.1186/s13072-016-0064-6. URL: http://dx.doi.org/10.1186/s13072-016-0064-6.
- [4] D. G. Phelan, D. J. Anderson, S. E. Howden, R. C. B. Wong, **P. F. Hickey**, K. Pope, G. R. Wilson, A. Pébay, A. M. Davis, S. Petrou, A. G. Elefanty, E. G. Stanley, P. A. James, I. Macciocca, M. Bahlo, M. M. Cheung, D. J. Amor, D. A. Elliott and P. J. Lockhart. "ALPK3-deficient cardiomyocytes generated from patient-derived induced pluripotent stem cells and mutant human embryonic stem cells display abnormal calcium handling and establish that ALPK3 deficiency underlies familial cardiomyopathy". In: *European Heart Journal* 37.33 (Sep. 01, 2016), pp. 2586-2590. ISSN: 0195-668X, 1522-9645. DOI: 10.1093/eurheartj/ehw160. URL: http://dx.doi.org/10.1093/eurheartj/ehw160.
- [5] D. Lacey, **P. F. Hickey**, B. D. Arhatari, L. A. O'Reilly, L. Rohrbeck, H. Kiriazis, X. Du and P. Bouillet. "Spontaneous retrotransposon insertion into TNF 3'UTR causes heart valve disease and chronic polyarthritis". In: *Proceedings of the National Academy of Sciences of the United States of America* 112.31 (Aug. 04, 2015), pp. 9698-9703. ISSN: 0027-8424, 1091-6490. DOI: 10.1073/pnas.1508399112. URL: http://dx.doi.org/10.1073/pnas.1508399112.
- [6] H. Oey, L. Isbel, **P. F. Hickey**, B. Ebaid and E. Whitelaw. "Genetic and epigenetic variation among inbred mouse littermates: identification of inter-individual differentially methylated regions". In: *Epigenetics & Chromatin* 8 (Dec. 12, 2015), p. 54. ISSN: 1756-8935. DOI: 10.1186/s13072-015-0047-z. URL: http://dx.doi.org/10.1186/s13072-015-0047-z.
- [7] P. F. Hickey and M. Bahlo. "X chromosome association testing in genome wide association studies". In: Genetic Epidemiology 35.7 (Nov. 2011), pp. 664-670. ISSN: 0741-0395, 1098-2272. DOI: 10.1002/gepi.20616. URL: http://dx.doi.org/10.1002/gepi.20616.
- [8] M. Bahlo, J. Stankovich, P. Danoy, **P. F. Hickey**, B. V. Taylor, S. R. Browning, Australian, ew Zealand Multiple Sclerosis Genetics Consortium (ANZgene), M. A. Brown and J. P. Rubio. "Saliva-derived DNA performs well in large-scale, high-density single-nucleotide polymorphism microarray studies". In: *Cancer Epidemiology, Biomarkers & Prevention* 19.3 (Mar. 2010), pp. 794-798. ISSN: 1055-9965, 1538-7755. DOI: 10.1158/1055-9965.EPI-09-0812. URL: http://dx.doi.org/10.1158/1055-9965.EPI-09-0812.
- [9] L. G. Riley, S. Cooper, **P. F. Hickey**, J. Rudinger-Thirion, M. McKenzie, A. Compton, S. C. Lim, D. Thorburn, M. T. Ryan, R. Giegé, M. Bahlo and J. Christodoulou. "Mutation of the mitochondrial tyrosyl-tRNA synthetase gene, YARS2, causes myopathy, lactic acidosis, and sideroblastic anemia-MLASA syndrome". In: *American Journal of Human Genetics* 87.1 (Jul. 09, 2010), pp. 52-59. ISSN: 0002-9297, 1537-6605. DOI: 10.1016/j.ajhg.2010.06.001. URL: http://dx.doi.org/10.1016/j.ajhg.2010.06.001.

#### Preprints and under review

- [1] L. F. Rizzardi\*, **P. F. Hickey\***, V. Rodriguez DiBlasi, R. Tryggvadóttir, 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". Mar. 24, 2017. DOI: 10.1101/120386. URL: http://biorxiv.org/content/early/2017/03/24/120386.
- [2] The eGTEx Project. "Enhancing GTEx: Bridging the gaps between genotype, gene expression, and disease". (submitted). 2017.
- [3] N. J. Brown, P. F. Hickey, P. J. Lockhart, F. Miriam, C. Bromhead, T. Desai, T. Vick, G. Gillies, H. Mountford, E. Fitzpatrick, L. Gordon, P. Hewson, V. Anderson, M. Delatycki, M. Bahlo, S. J. Wilson and I. E. Scheffer. "Tracing Autism Traits in Autosomal Dominant Families". (submitted). 2016.

## Theses

- [1] P. F. Hickey. "The statistical analysis of high-throughput assays for studying DNA methylation". PhD thesis. Department of Mathematics and Statistics, The University of Melbourne, 2015. URL: https://minerva-access.unimelb.edu.au/handle/11343/55699.
- [2] P. F. Hickey. "X chromosome association testing in genome-wide association studies". Honours Thesis. Department of Mathematics and Statistics, The University of Melbourne, Nov. 05, 2009.

#### Technical reports, conference papers and published abstracts

- [1] I. Scheffer, K. Williams, C. Green, K. Pereira, N. J. Brown, **P. F. Hickey**, V. Lukic, G. Gillies, M. Delatycki, P. J. Lockhart, M. Bahlo and S. J. Wilson. "The Victorian Collaborative Autism Study: A family and community study of the genetics of autism spectrum disorder". In: *Journal of Intellectual Disability Research*. Ed. by J. of Intellectual Disability Research. Vol. 60. 7. 2016, p. 732. DOI: 10.1111/jir.12305.
- [2] D. G. Phelan, G. R. Wilson, K. Pope, G. Gillies, J. Sim, M. Bahlo, **P. F. Hickey**, C. Bromhead, P. A. James, D. du Sart, M. Delatycki, R. Leventer, D. J. Amor and P. J. Lockhart. "Identification and characterisation of a novel hypertrophic cardiomyopathy gene". In: *Pathology: The Journal of the Royal College of Pathologists of Australasia*. Ed. by P. T. J. of the Royal College. Vol. 46, 2014, pp. S91-S92.
- [3] D. G. Phelan, G. R. Wilson, J. Sim, M. Bahlo, **P. F. Hickey**, P. A. James, D. du Sart, M. Delatyki, D. J. Amor and P. J. Lockhart. "Identification and characterisation of a novel hypertrophic cardiomyopathy gene". In: *Global Heart*. Ed. by G. Heart. Vol. 9. 1. Elsevier, 2014, p. e316. DOI: http://dx.doi.org/10.1016/j.gheart.2014.03.2365.

# Commentaries and meeting reports

[1] **P. F. Hickey** and M. D. Robinson. *Genomics by the beach*. Apr. 14, 2014. DOI: 10.1186/gb4171. URL: http://dx.doi.org/10.1186/gb4171.

# Computer skills and software development

Very proficient with statistical computing, particularly R, and strategies for managing and analysing large (multi-gigabyte to terabtye size) datasets, especially genomics data. Familiar with C, C++, and various databases — both as standalone tools and as integrations with and extensions of R — as well as Python, Unix shell tools, and cluster job schedulers.

Developer and contributor to several R packages through the Bioconductor project. Also developed and published Python software. Additional projects are available from https://github.com/PeteHaitch.

#### R packages

Download statistics are reported for years since I have been a substantial author/contributor to the package (data from http://bioconductor.org/packages/stats/).

#### Author

- Genomic Tuples: Representation and Manipulation of Genomic Tuples
  - Number of downloads (2014-2017): 497, 2426, 2022, 1006
- bsseq: Analyse, Manage and Store Bisulfite Sequencing Data

- Number of downloads (2014-2017): 5049, 6631, 8419, 5661

#### Contributor

- minfi: Analysing Illumina Infinium DNA Methylation Arrays
  - Number of downloads (2017): 20165

# Python packages

#### Author

• methtuple: A caller for DNA methylation events that co-occur on the same DNA fragment from high-throughput bisulfite sequencing data, such as whole-genome bisulfite-sequencing

# Funding

#### Grants

- 2015: Bioconductor Travel Grant
  - To attend the Bioconductor meeting in Seattle, USA
- 2013: EMBL Australia Travel Grant
  - To attend the EMBL PhD Symposium in Heidelberg, Germany

# **Scholarships**

- 2013: Victorian Life Sciences Computation Initiative PhD Top-Up Scholarship.
- 2012: Statistical Society of Australia (Victoria Branch) scholarship to attend the Young Statisticians Conference (2013)
- 2011-2015: Australian Postgraduate Award
- 2009: Maurice Belz scholarship, The University of Melbourne
  - A competitive scholarship awarded to complete Honours degree in statistics, stochastic processes or operations research
- 2009: Alan W Harris Honours scholarship, The Walter and Eliza Hall Institute of Medical Research

#### Awards

- 2015: Edith Moffat Travel Award
  - To interview for international for postdoctoral positions and attend the European Bioconductor meeting
- 2013: Prize for best lightning talk, Australian Epigenetics Conference 2013
- 2013: Prize for third best oral presentation, Young Statisticians Conference 2013
- 2010: Best presentation (Statistics), 2010 Victorian Mathematics and Students' Conference

## Presentations

#### **Talks**

#### Conferences

- 2017: Neuronal brain region-specific DNA methylation and chromatin accessibility are associated with neuropsychiatric disease heritability. Contributed talk, GTEx Project Community Meeting, Rockville, USA (28/06)
- 2017: Developing statistical methods for large epigenomic studies in the human brain. Contributed talk, ENAR 2017 Spring Meeting, Washington D.C., USA (13/03)
- 2016: New features in bsseq for analysing large whole genome bisulfite-sequencing datasets. Lightning talk, BioC 2016, San Francisco, USA (24/06)
- 2016: The GenomicTuples package. Lightning talk, BioC 2015 Developers' Day, Seattle, USA (20/7)
- 2015: Genomic tuples and DNA methylation patterns. Contributed talk, European Bioconductor Developers' Meeting, Heidelberg, Germany (12/01)
- 2014: Making sense of DNA methylation data. PhD completion seminar, Melbourne, Australia (15/09)
- 2014: Simulating whole-genome DNA methylation data). Contributed talk, Australian Statistical Conference/International Mathematical Statistics Annual Meeting, Sydney, Australia (10/07)
- 2013: Exploiting local dependencies in genome-wide studies of DNA methylation. Contributed talk, Young Statisticians Conference, Melbourne, Australia (07/02)
- 2012: Spatial dependence of CpG-methylation from whole genome bisulfite sequencing. Contributed talk, Epigenomics of Common Diseases Meeting, Baltimore, USA (15/10)
- 2012: Spatial dependence of DNA methylation. Contributed talk, Australian Statistical Conference, Adelaide, Australia (15/07)
- 2010: Bioinformatics Applied statistics in modern molecular biology. Contributed talk (with Davis McCarthy), 2010 Victorian Mathematics and Statistics Students' Conference, Melbourne, Australia (02/07)

#### Other

- 2013: Bioinformatics for bisulfite sequencing. Invited talk, La Trobe University Sequencing Users Group, Melbourne, Australia (28/08)
- 2010: X chromosome association testing in genome wide association studies. Invited talk, Statistical Society of Australia Victorian branch meeting, Melbourne, Australia (24/08)

#### Posters

- 2014: Simulating whole-genome bisulfite-sequencing data. Poster, Lorne Genome, Lorne, Australia (17/01)
- 2013: Simulating whole-genome bisulfite-sequencing data. Poster, Epigenetics 2013, Shoal Bay, Australia (03/12)
- 2011: Analysis of mouse exome sequencing: filtering institute specific single nucleotide variants (SNVs). Poster, GeneMappers 8th Australian Human Gene Mapping Conference, Hobart, Australia (04/04)
- 2010: X chromosome association testing in genome wide association studies. Poster, The International Genetic Epidemiology Society Conference, Boston, USA (12/10)
- 2010: X chromosome association testing in genome wide association studies. Poster, The Australasian Microarray and Associated Technologies Association Conference, Hobart, Australia (16/09)
- 2009: Homozygosity by state analysis in highly inbred pedigrees. Poster, GeneMappers 7th Australian Human Gene Mapping Conference, Sydney, Australia (15/06)

# Teaching experience

## Workshops

• 2016: Analysing DNA methylation data with Bioconductor. BioC 2016, Palo Alto, USA (26/06)

# **Tutoring**

- 2006-2014: Private tutoring
  - One-on-one and small-group tutoring for students studying Masters-level biostatistics, third-year
    university level statistics and first-year university level engineering mathematics. Tutoring senior
    high school students in elementary, intermediate and advanced mathematics subjects and physics.
- 2011: Classroom teaching assistant
  - Assisting in mathematics classes for final year high school students

# Professional activities and service

# Reviewing

Referee for Nature Methods, PLoS Genetics, Genome Biology, Bioinformatics, PLoS Computational Biology, Genetic Epidemiology, Heredity, and F1000.

# Professional society memberships

- Member, Statistical Society of Australia
- Member, International Biometric Society (ENAR)

# References

# **Professor Terry Speed**

**Bioinformatics Division** 

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# Assistant Professor Kasper Hansen

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# Professor Melanie Bahlo

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