Chaolong Wang, Ph.D.

Curriculum Vitae updated on April 17, 2015

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Singapore 138672, Singapore Website: http://chaolongwang.github.io

Personal Citizenship: People's Republic of China

INFORMATION Languages: English, Chinese Mandarin, Cantonese, Hokkien (Min Nan)

RESEARCH Interest Population genetics, statistical genetics/genomics, disease gene mapping, integrative genomics,

INTEREST next-generation sequencing data analysis, high dimensional data analysis

Professional 01/2015-Now: **Principal Investigator**, Computational & Systems Biology

Appointments Genome Institute of Singapore, A*STAR, Singapore

09/2012-12/2014: **Research Fellow**, Department of Biostatistics

Harvard University, Boston, Massachusetts, USA

• Postdoctoral research on statistical genetics with Dr. Xihong Lin

EDUCATION 08/2012: **Ph.D. in Bioinformatics**

University of Michigan, Ann Arbor, Michigan, USA

• Ph.D. advisor: Dr. Noah Rosenberg (now at Stanford University)

• Thesis: Statistical methods for analyzing human genetic variation in diverse populations

• 01/2012-08/2012: "Post"-doctoral research on statistical genetics with Dr. Gonçalo Abecasis

04/2011: M.A. in Statistics and M.S. in Bioinformatics

University of Michigan, Ann Arbor, Michigan, USA

07/2008: **B.S. in Physics** Peking University, Beijing, China

Selected Awards

- Charles J. Epstein Trainee Award semifinalist, American Society of Human Genetics, 2013
- Stellar Abstract Award in the 6th Annual PQG Conference, Harvard University, 2012
- HHMI International Student Research Fellowship, Howard Hughes Medical Institute, 2011-2012
- Rackham Predoctoral Fellowship, University of Michigan, 2011 (declined)
- DeLill Nasser Award, Genetics Society of America, 2011
- Departmental Fellowship (Biomedical Sciences), University of Michigan, 2008
- May Fourth Scholarship, Peking University, 2006
- Dean's List, Hong Kong University of Science & Technology, 2005

TEACHING EXPERIENCE 06/2014 & 12/2014: Sequence Analysis Workshop Instructor, University of Michigan Lecture and hands-on practical on Estimates of Genetic Ancestry

Doctars and names on presented on Estimates of Science (Interest)

10/2013: PQG Short Course Lecturer, Harvard School of Public Health Statistical methods for ancestry inference with applications to disease gene mapping

01/2011-04/2011: Graduate Student Instructor, University of Michigan BIOSTAT 646 - High throughput molecular genomic and epigenomic data analysis

Manuscripts submitted H Chen*, C Wang*, MP Conomos, AM Stilp, Z Li, T Sofer, AA Szpiro, W Chen, JM Brehm, JC Celedon, SS Redline, GJ Papanicolaou, TA Thornton, CC Laurie, K Rice, X Lin. Control for population structure and relatedness for binary traits in genetic association studies using logistic mixed models. Submitted. (* Co-first author)

X Lin, S Lee, M Wu, C Wang, H Chen, Z Li, X Lin. Test for rare variants by environment interactions in sequencing association studies. In revision.

PUBLICATIONS

- C Wang, X Zhan, L Liang, GR Abecasis, X Lin (2015). Improved ancestry estimation for both genotyping and sequencing data using projection Procrustes analysis and genotype imputation. American Journal of Human Genetics, in press.
- C Wang*, X Zhan*, J Bragg-Gresham, HM Kang, D Stambolian, E Chew, K Branham, J Heckenlively, The FUSION Study, RS Fulton, RK Wilson, ER Mardis, X Lin, A Swaroop, S Zöllner, GR Abecasis (2014). Ancestry estimation and control of population stratification for sequence-based association studies. *Nature Genetics*, 46:409-415. (* Co-first author)
- X Zhan*, DE Larson*, C Wang*, DC Koboldt, Y Sergeev, 52 other coauthors, ER Mardis, A Swaroop, GR Abecasis (2013). Identification of a rare coding variant in Complement 3 associated with age-related macular degeneration. Nature Genetics 45: 1375-1379. (* Co-first author)
- C Wang, KB Schroeder, NA Rosenberg (2012). A maximum-likelihood method to correct for allelic dropout in microsatellite data with no replicate genotypes. Genetics 192: 651-669.
- C Wang, S Zöllner, NA Rosenberg (2012). A quantitative comparison of the similarity between genes and geography in worldwide human populations. PLoS Genetics 8: e1002886. [Featured in Science 337: 1151, 2012]
- TJ Pemberton, C Wang, JZ Li, NA Rosenberg (2010). Inference of unexpected genetic relatedness among individuals in HapMap Phase III. American Journal of Human Genetics 87: 457-464. [Featured in Am. J. Hum. Genet. 87: 447-448, 2010 and Genetics 186(2): NP, 2010]
- C Wang, ZA Szpiech, J Degnan, M Jakobsson, TJ Pemberton, JA Hardy, AB Singleton, NA Rosenberg (2010). Comparing spatial maps of human population-genetic variation using Procrustes analysis. Statistical Applications in Genetics and Molecular Biology 9: 13.
- JT Mosher, TJ Pemberton, K Harter, C Wang, EO Buzbas, P Dvorak, C Simon, SJ Morrison, NA Rosenberg (2010). Lack of population diversity in commonly used human embryonic stem-cell lines. New England Journal of Medicine 362: 183-185. [Featured in Nature 462: 945, 2009]
- NM Kopelman, L Stone, C Wang, D Gefel, MW Feldman, J Hillel, NA Rosenberg (2009). Genomic microsatellites identify shared Jewish ancestry intermediate between Middle Eastern and European populations. BMC Genetics 10: 80.
- L Huang, C Wang, NA Rosenberg (2009). The relationship between imputation error and statistical power in genetic association studies in diverse populations. American Journal of Human Genetics 85: 692-698. [Featured in Am. J. Hum. Genet. 85: 539-540, 2009 and Nat. Rev. Genet. 10: 817, 2009]
- CL Wang, KW Au, CK Chan, HW Lau, KY Szeto (2008). Detecting hierarchical organization in complex networks by nearest neighbor correlation. Studies in Computational Intelligence 129: 487-494 (Conference Proceedings of NICSO 2007).

Software DEVELOPED

- **LASER**: a package written in C++ for estimating individual ancestry using either sequencing reads or gentyping data. (Wang et al. 2014, Nat. Genet.; Wang et al. 2015, AJHG)
- MicroDrop: a C++ program for estimating and correcting for allelic dropout in microsatellite data without replicated genotyping. (Wang et al. 2012, Genetics).

Conference

- C Wang, Han Chen, X Lin. Control of population stratification in family data using pedigree Presentations information and ancestry principal components. The American Society of Human Genetics 64th Annual Meeting, San Diego, USA, October 2014. (Poster)
 - C Wang, L Liang, GR Abecasis, X Lin. Tracing individual ancestry in a principal components space. The American Society of Human Genetics 63rd Annual Meeting, Boston, USA, October

2013. (Platform talk, Trainee Award Semifinalist)

C Wang, X Zhan, S Zöllner, GR Abecasis. Estimating individual ancestry using next generation sequencing. CSHL Meeting "The Biology of Genomes", Cold Spring Harbor, New York, USA, May 2013. (**Poster**)

C Wang, X Zhan, S Zöllner, GR Abecasis. Estimating individual ancestry using next generation sequencing. HHMI Janelia Conference "Biological Sequence Analysis and Probabilistic Models", Ashburn, Virginia, USA, March 2013. (Poster and blitz talk)

C Wang, X Zhan, S Zöllner, GR Abecasis. Estimating individual ancestry from extremely lowcoverage sequencing data. The 6th Annual PQG Conference "Sequencing and Complex Traits: beyond 1000 Genomes", Boston, Massachusetts, USA, November 2012. (Platform talk, Stellar Abstract Award)

C Wang, NA Rosenberg. A quantitative comparison of the similarity between genes and geography in worldwide human populations. The 12th International Congress of Human Genetics & The American Society of Human Genetics 61st Annual Meeting, Montreal, Canada, October 2011. (Poster)

C Wang, KB Schroeder, NA Rosenberg. A maximum likelihood genotype imputation method to correct for allelic dropout in microsatellite data. ENAR 2011 Spring Meeting, Miami, Florida, USA, March 2011. (Platform talk)

C Wang, KB Schroeder, NA Rosenberg. A genotype imputation method for allelic dropout in microsatellite data. The American Society of Human Genetics 60th Annual Meeting, Washington DC, USA, November 2010. (Poster)

C Wang, NA Rosenberg. Comparing spatial maps of genetic variation and geographic locations using Procrustes analysis. EMBO/EMBL Symposium "Human Variation: Cause and Consequence", Heidelberg, Germany, June 2010. (Poster)

C Wang, ZA Szpiech, J Degnan, M Jakobsson, TJ Pemberton, JA Hardy, AB Singleton, NA Rosenberg. Comparing spatial maps of population-genetic variation using Procrustes analysis. The American Society of Human Genetics 59th Annual Meeting, Honolulu, Hawaii, USA, October 2009. (Poster)

Membership

2009-Now: American Society of Human Genetics American Statistical Association (2010-2011) International Biometric Society, ENAR (2010-2011)

Professional ACTIVITIES

2013-2014: Member, Organizing Committee of the HHMI Alumni Network in Boston Region

2012-2013: Organizer, PQG Short Courses, Harvard School of Public Health

2009-Now: Referee for peer-reviewed journals, including Annals of Human Geneitcs, BMC Bioinformatics, BMC Genetics, Clinical Chemistry, Genetics, Human Biology, Journal of Human Genetics, Molecular Biology and Evolution, PLoS ONE, PNAS, Scientific Reports

Computing SKILLS

Programming: C/C++, Perl, R, MATLAB, Shell Scripting Operating Systems: Linux/Unix, Mac OS X, Microsoft Windows Applications: LATEX, Microsoft Office, Adobe Illustrator, HTML

References Available upon request.