# Alex H. Wagner, PhD

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Research Interests Cancer Genomics, Precision Medicine, Medical Informatics

Research Experience

Division of Oncology,

**Instructor in Medicine** 

Washington University School of Medicine

Saint Louis, MO

Advisors:

Obi L. Griffith, PhD Timothy J. Ley, MD

## Postdoctoral Research Scholar

January 2015 to May 2019

June 2019 to Present

McDonnell Genome Institute,

Washington University School of Medicine

Saint Louis, MO

Advisors:

Obi L. Griffith, PhD

Ramaswamy Govindan, MD

#### **Graduate Research Assistant**

August 2010 to December 2014

Coordinated Laboratory for Computational Genomics,

University of Iowa College of Engineering

Iowa City, IA

Advisors:

Terry A. Braun, PhD

Edwin M. Stone, MD, PhD

# **Clinical Laboratory Technologist**

July 2008 to July 2010

Department of Laboratory Medicine and Pathology,

Mayo Clinic, Rochester, MN

Supervisors:

Dianna Bowden

Thomas P. Moyer, PhD

## **Biological Laboratory Aide**

Jan 2007 to Sep 2007

USDA Agricultural Research Service, Iowa State University, Ames, IA Supervisor: David Grant, PhD

## Education

#### University of Iowa, Iowa City, IA

PhD, Computational Genetics, December 2014

- Thesis Topic: Computational Methods for Identification of Disease-Associated Variations in Exome Sequencing
- · Advisors: Terry A. Braun, PhD and Edwin M. Stone, MD, PhD
- GPA: 3.90

Graduate Certificate, Bioinformatics, May 2013

· Advisor: Terry A. Braun, PhD

• GPA: 3.96

# Iowa State University, Ames, IA

BS, Biology, May 2008

- Minor in Mathematics
- Cum Laude
- GPA: 3.51

#### **Extracurricular Education**

High Performance Computing, Gregory Howes, Iowa, Summer 2012 Machine Learning, Andrew Ng, Stanford (Online), Fall 2011 Intro to Databases, Jennifer Widom, Stanford (Online), Fall 2011

#### **Published Works**

- EK Barnell, P Ronning, KM Campbell, K Krysiak, BJ Ainscough, C Ramirez, N Spies, J Kunisaki, ZL Skidmore, F Gomez, L Trani, M Matlock, AH Wagner, SJ Swamidass, M Griffith, OL Griffith (2019) "Standard operating procedure for somatic variant refinement of tumor sequencing data". Genetics in Medicine. doi: 10.1038/s41436-018-0278-z.
- AH Wagner<sup>†</sup>, S Devarakonda<sup>†</sup>, ZL Skidmore, K Krysiak, A Ramu, L Trani, J Kunisaki, ..., M Griffith, OL Griffith, R Govindan (2018) "Recurrent WNT Pathway Alterations are Frequent in Relapsed Small Cell Lung Cancer". Nature Communications. doi: 10.1038/s41467-018-06162-9
- 3. BJ Ainscough<sup>†</sup>, EK Barnell<sup>†</sup>, KM Campbell, **AH Wagner**, TE Rohan, R Govindan, M Griffith, ER Mardis, SJ Swamidass, OL Griffith (2018) "A deep learning approach to automate refinement of somatic variant calling from cancer sequencing data". *Nature Genetics*. doi: 10.1038/s41588-018-0257-y
- 4. AM Danos<sup>†</sup>, DI Ritter<sup>†</sup>, **AH Wagner**, K Krysiak, ..., S Kulkarni, M Griffith, S Madhavan, OL Griffith (2018) "Adapting crowdsourced clinical cancer curation in CIViC to the ClinGen minimum variant level data community-driven standards". *Human Mutation*. doi: 10.1002/humu.23651.
- 5. K Cotto<sup>†</sup>, **AH Wagner**<sup>†</sup>, YY Feng, S Kiwala, AC Coffman, G Spies, A Wollam, NC Spies, OL Griffith, M Griffith (2017) "DGldb 3.0: a redesign and expansion of the drug-gene interaction database". *Nucleic Acids Research*. doi: 10.1093/nar/gkx1143.
- M Griffith, NC Spies, K Krysiak, JF McMichael, AC Coffman, AM Danos, BJ Ainscough, CA Ramirez, DT Rieke, L Kujan, EK Barnell, AH Wagner, ..., OL Griffith (2017) "CIVIC is a community knowledgebase for expert crowdsourcing the clinical interpretation of variants in cancer". Nature Genetics. doi: 10.1038/ng.3774.
- 7. BJ Ainscough, M Griffith, AC Coffman, AH Wagner, J Kunisaki, MNK Choudhary, JF McMichael, RS Fulton, RK Wilson, OL Griffith, ER Mardis

- (2016) "DoCM: a database of curated mutations in cancer". *Nature methods*. doi: 10.1038/nmeth.4000.
- 8. M Griffith, OL Griffith, K Krysiak, ZL Skidmore, MJ Christopher, JM Klco, A Ramu, TL Lamprecht, **AH Wagner**, ..., TJ Ley (2016) "Comprehensive genomic analysis reveals FLT3 activation and a therapeutic strategy for a patient with relapsed adult B-lymphoblastic leukemia". *Experimental hematology*. doi: 10.1016/j.exphem.2016.04.011.
- 9. ZL Skidmore, **AH Wagner**, R Lesurf, KM Campbell, J Kunisaki, OL Griffith, M Griffith (2016) "GenVisR: Genomic Visualizations in R". *Bioinformatics*. doi: 10.1093/bioinformatics/btw325.
- AH Wagner, AC Coffman, BJ Ainscough, NC Spies, ZL Skidmore, KM Campbell, K Krysiak, D Pan, JF McMichael, JM Eldred, JR Walker, RK Wilson, ER Mardis, M Griffith\*, OL Griffith\* (2016) "DGIdb 2.0: mining clinically relevant drug?gene interactions". *Nucleic Acids Research*. doi: 10.1093/nar/gkv1165.
- SS Whitmore, AH Wagner, AP DeLuca, AV Drack, EM Stone, BA Tucker, S Zeng, TA Braun, RF Mullins, TE Scheetz (2014) "Transcriptomic analysis across nasal, temporal, and macular regions of human neural retina and RPE/choroid by RNA-Seq". Experimental Eye Research. doi:10.1016/j.exer.2014.11.001
- TP Sharma, CM McDowell, Y Liu, AH Wagner, D Thole, BP Faga, RJ Workinger, TA Braun, AF Clark (2014) "Optic nerve crush induces spatial and temporal gene expression patterns in retina and optic nerve of BALB/cJ mice". Molecular Neurodegeneration. doi: 10.1186/1750-1326-9-14
- 13. TA Braun, RF Mullins, AH Wagner, J Andorf, R Johnston, B Bakall, AP DeLuca, G Fisherman, R Weleber, A Cideciyan, S Jacobson, V Sheffield, B Tucker, EM Stone (2013) "Non-exomic and synonymous variants in ABCA4 are an important cause of Stargardt disease". Human Molecular Genetics. doi: 10.1093/hmg/ddt367
- 14. **AH Wagner**, KR Taylor, AP DeLuca, TL Casavant, RF Mullins, EM Stone, TE Scheetz, TA Braun (2013), "Prioritization of Retinal Disease Genes: An Integrative Approach." *Human Mutation*. doi: 10.1002/humu.22317
- 15. **AH Wagner**, VN Anand, W Wang, JE Chatterton, D Sun, AR Shepard, N Jacobson, L Pang, AP DeLuca, TL Casavant, TE Scheetz, RF Mullins, TA Braun, AF Clark (2013) "Exon-level expression profiling of ocular tissues". *Experimental Eye Research*. doi: 10.1016/j.exer.2013.03.004
- AP DeLuca, AH Wagner, KR Taylor, B Faga, D Thole, VC Sheffield, ..., TA Braun. (2011). "Sequencing and disease variation detection tools and techniques". IEEE/ACS International Conference on Computer Systems and Applications (AICCSA). doi: 10.1109/AICCSA.2011.6126607

<sup>&</sup>lt;sup>†</sup>Denotes Co-First Authorship

#### Interim Products

- 1. **AH Wagner**, B Walsh, G Mayfield, ..., M Griffith, OL Griffith, A Margolin (2018) "A harmonized meta-knowledgebase of clinical interpretations of cancer genomic variants". *bioRxiv*. doi: 10.1101/366856
- 2. AM Danos<sup>†</sup>, K Krysiak<sup>†</sup>, EK Barnell<sup>†</sup>, ..., **AH Wagner**, S Madhavan, M Griffith, OL Griffith (2019) "The ClViC knowledge model and standard operating procedures for curation and clinical interpretation of variants in cancer". *bioRxiv*. doi: 10.1101/700179

# Submitted Publications

- 1. **AH Wagner**, B Walsh, G Mayfield, ..., M Griffith, OL Griffith, A Margolin (Under secondary review, *Nature Genetics*) "A harmonized meta-knowledgebase of clinical interpretations of somatic cancer genomic variants".
- 2. AM Danos<sup>†</sup>, K Krysiak<sup>†</sup>, EK Barnell<sup>†</sup>, ..., **AH Wagner**, S Madhavan, M Griffith, OL Griffith (Under primary review, *Genome Medicine*) "The CIVIC knowledge model and standard operating procedures for curation and clinical interpretation of variants in cancer".

# Other Published Works

 AH Wagner, K Krysiak, KM Campbell, EK Barnell (2019) "Cancer Genomics for the Clinician: Tumor Heterogeneity" [Book Chapter]. Springer Publishing Group. ISBN-13: 9780826168672

# Funding Research and Fellowship Awards

•	NHGRI K99/R00	2019-present
•	NCI F32 Postdoctoral Fellowship	2017–2019

## **Training Awards**

•	NCI T32 Postdoctoral Training in Cancer Biology	2016–2017
•	NIGMS T32 Predoctoral Training Grant in Genetics	2013–2014
•	NIGMS T32 Predoctoral Training Grant in Bioinformatics	2011–2013
	<ul> <li>Consecutive annual awards granted for 2011-2012 and 2012</li> </ul>	2-2013.

# Merit Research Awards

ICTS Precision Medicine Abstract Award

January 16, 2018

- Precision Medicine Symposium Institute of Clinical and Translational Science Washington University in Saint Louis
- The most outstanding research was selected from more than 75 applicants to present to Eric Green, the director of the National Human Genome Research Institute (NHGRI).
- D.C. Spriestersbach Dissertation Prize Nominee

2015

- Genetics Program Nomination
- This biennial award recognizes excellence in doctoral research. Each
  of the twenty biological/life sciences programs at the University of Iowa
  nominates one dissertation submitted between July 1, 2013 and June 30,
  2015 to compete for the award.
- Outstanding Student Research Award Comp Bio

2012-2013

 This annual award recognizes a single student in the College of Engineering for exemplary research in the fields of bioinformatics and computational biology.

#### **Travel and Trainee Awards**

Trainee Abstract Award

May 2019

· Curating the Clinical Genome

**Annual Conference** 

Washington D.C.

1st Place Student/Trainee Travel Award

August 2018

Cancer Genomics Consortium

**Annual Conference** 

Nashville, TN

AACR-Bristol Myers Squibb Scholar-in-Training Award

April 2018

American Association for Cancer Research

Annual Conference

Chicago, IL

NSF Travel Grant

July 2013

 International Society for Computational Biology Intelligent Systems for Molecular Biology Annual Conference Berlin, Germany

Graduate Student Senate Travel Grant

May 2013

 Association for Research in Vision and Ophthalmology Annual Conference Seattle, WA

## **Oral Presentation Awards**

Annual Bioinformatics Retreat, University of Iowa

August 16, 2013

· Best Student Talk

Midwest Eye Research Symposium

July 6, 2012

· Outstanding Oral Presentation, 2nd Place

#### **Poster Presentation Awards**

Interdisciplinary Health Research Poster Session

April 23, 2013

· Best Poster Award, Center on Aging

## Presentations

<sup>†</sup>Oral Presentation; <sup>††</sup>Invited Speaker; \*Upcoming

#### **External Presentations**

AMIA 2019 Annual Conference, Washington, D.C.
 \*November 2019
 \*Panelist, computational representation of patient data

• CGC 2019 Annual Conference, Nashville, TN August 2019

\*\*Community resources for clinical variant classification in cancers\*\*

CGC 2019 Annual Conference, Nashville, TN
 †A new somatic variation model enables precise search strategies for clinical interpretations of patient tumors

Curating the Clinical Genome, Washington, D.C.
 May 2019
 Somatic variant curation standards enable improved identification of relevant clinical interpretations for tumor variants

• GAAGH Implementation Connect Hington LIK	April 2019	
<ul> <li>GA4GH Implementation Connect, Hinxton, UK</li> <li><sup>††</sup> Variant Interpretation for Cancer Consortium: Workstream</li> </ul>	•	
AACR Annual Conference, Atlanta, GA	March 2019	
Cancer genome interpretation with CIViCpy	Maron 2010	
GA4GH-AMED Symposium, Sendai, Japan	March 2019	
<sup>††</sup> The Variant Interpretation for Cancer Consortium, a Gen	omic Knowledge	
Workstream Driver Project of the GA4GH	ŭ	
GA4GH 6th Annual Plenary, Basel, Switzerland	October 2018	
<sup>††</sup> The Variant Interpretation for Cancer Consortium		
<ul> <li>Cancer Genomics Consortium, Nashville, TN</li> </ul>	August 2018	
<sup>†</sup> Coordinating variant interpretation knowledgebases		
improves clinical interpretation of genomic variants in canc		
<ul> <li>American Association for Cancer Research, Chicago, IL</li> </ul>	April 2018	
Standardization and coordination of variant interpretation		
knowledgebases improves clinical genome actionability		
Curating the Clinical Genome, Hinxton, UK	June 2016	
The Drug Gene Interaction Database	E   0040	
AGBT Annual Conference, Orlando, FL  The Drive Come Internation Patch as a	February 2016	
The Drug Gene Interaction Database	July 2012	
<ul> <li>ISMB Annual Conference, Berlin, Germany         Positive and Unlabeled Learning for Prioritization (PULP)     </li> </ul>	July 2013	
ARVO Annual Conference, Seattle, WA	May 2013	
Positive and Unlabeled Learning for Prioritizing Candidate	Way 2010	
Variants in Retinal Degenerative Diseases		
BICB Industry Symposium, Minneapolis, MN	May 2013	
Positive and Unlabeled Learning for Prioritizing Candidate		
Variants in Retinal Degenerative Diseases		
ARVO Annual Conference, Ft. Lauderdale, FL	May 2012	
RNA Sequencing for Identification of Genetic Factors in Retinal Disease		
<ul> <li>Joint Bioinformatics Retreat, Ames, IA</li> </ul>	Aug 2011	
Using RNA Sequencing To Identify And Isolate Causative		
Genetic Factors In Retinal Disease		
Washington University in Saint Louis		
ICTS Precision Medicine Symposium	January 2018	
Coordinating variant interpretation knowledgebases improves		
clinical interpretation of genomic variants in cancers		
Postdoctoral Research Symposium	March 2017	
The Drug Gene Interaction Database		
University of Iowa		
Genetics Retreat 2014	October 2014	
<sup>†</sup> Active Phenotype Acquisition for the Genetic Characterization		
of Heritable Retinal Diseases		
<ul> <li>Engineering Research Open House 2014</li> </ul>	April 2014	
Positive and Unlabeled Learning for Prioritization (PULP)		
Genetics Retreat 2013	October 2013	
Prioritizing Disease Genes in Exome Studies		
Joint Bioinformatics Retreat	August 2013	

<sup>†</sup>Positive and Unlabeled Learning for Prioritization Interdisciplinary Health Research Poster Session April 2013 Positive and Unlabeled Learning for Prioritizing Candidate Variants in Retinal Degenerative Diseases Genetics Retreat 2012 November 2012 Machine Learning Based Prioritization of Retinal Disease Genes Joint Bioinformatics Retreat October 2012 <sup>†</sup>Prioritization of Retinal Disease Genes: An Integrative Approach · Midwest Eye Research Symposium July 2012 <sup>†</sup>Machine Learning Based Prioritization of Eye Disease Genes Genetics Retreat 2011 February 2012 Exon-level Expression Profiling of Ocular Tissues **Guest Lecturer** December 2019 Variant Interpretation Introduction to Bioinformatics Instructor: Michael Baudis University of Zurich Zurich, Switzerland **Workshop Instructor** August 2019 Bioinformatics: Understanding what's Underneath the Hood Cancer Genomics Consortium Annual Conference Nashville, TN March 2019 **Workshop Leader** Genomic Knowledge Standards AMED-GA4GH Workshop Sendai, Japan Nov 2018 **Teaching Assistant / Lecturer** Advanced Sequencing Technologies and Applications Cold Spring Harbor Laboratories Cold Spring Harbor, NY **Workshop Instructor** August 2018 Introduction to bioinformatics learning resources Cancer Genomics Consortium Annual Conference Nashville, TN Fall 2017-Summer 2019 Instructor Escape from Perlgatory: Developing in Python and Ruby McDonnell Genome Institute Saint Louis, MO **Teaching Assistant / Lecturer** Nov 2017 Advanced Sequencing Technologies and Applications Cold Spring Harbor Laboratories

Teaching

Experience

Cold Spring Harbor, NY

Nov 2016 **Workshop Instructor** CIViC Hackathon Netherlands Cancer Institute (NKI) Amsterdam, Netherlands **Teaching Assistant / Lecturer** Nov 2016 Advanced Sequencing Technologies and Applications Cold Spring Harbor Laboratories Cold Spring Harbor, NY **Teaching Assistant** Fall 2014 051:123 - Bioinformatics Techniques Instructor: Thomas L. Casavant Department of Biomedical Engineering University of Iowa **Guest Lecturer** Spring 2014 051:080 - Bioimaging and Bioinformatics Instructor: Todd E. Scheetz Department of Biomedical Engineering University of Iowa **Teaching Assistant** Spring 2014 051:122 - Computational Genomics Instructor: Thomas L. Casavant Department of Biomedical Engineering University of Iowa **Teaching Assistant** Fall 2013 051:123 - Bioinformatics Techniques Instructor: Terry A. Braun Department of Biomedical Engineering University of Iowa Instructor Fall 2013 Introduction to Bioinformatics Computing with Python Supplement to 051:123 - Bioinformatics Techniques Department of Biomedical Engineering University of Iowa **Teaching Assistant** Fall 2006 **BIOL 313 - Principles of Genetics** Instructor: Jack Girton Department of Biology Iowa State University

Service

## Variant Representation (VR) Group Lead, GA4GH

2018-Present

- Co-Leader of the GA4GH Genomic Knowledge Standards VR subgroup
- Technical co-lead and maintainer of the GA4GH VR Specification

## Co-Director, Variant Interpretation for Cancer Consortium

- Led the design and execution of the VICC knowledge harmonization effort
- Developed the VICC research objectives (summary image) and organized eight expert-led Working Groups to achieve them

## Editor, CIViC Knowledgebase

2015-Present

2018-Present

- Moderated curation of clinical interpretations of genomic variants from biomedical literature
- Editor, with over 450 moderations (#7 all-time leader in moderations)

# Workshop Co-chair, HemOnc / VICC Hackathon

January 2020

 Pre-conference Workshop to AACR Advancing Precision Medicine Drug Development

#### **Peer Review**

<ul> <li>Nucleic Acids Research</li> </ul>	October 2019
Cancer Genetics	August 2018
BMC Cancer	May 2018
Genome Medicine	April 2017
Nature Communications	August 2016
Science Translational Medicine	August 2016

# Software Familiarity

# **Programming languages and environments:**

 Amazon Web Services (AWS), C, C++, Django, Docker, LSF, SQL, Perl, Python (PyCharm, Jupyter Notebooks), R (RStudio), Ruby on Rails (RubyMine), SGE, UNIX

## Common software (bioinformatics):

 Bedtools, Ballgown, BWA, CNVkit, Cufflinks, GATK, GenVisR, GISTIC, GMS, HiSat2, Kallisto, IGV, Picard, Pindel, Pizzly, RNA-SeQC, Samtools, SomaticSniper, Strelka, Stringtie, Tophat, VarScan, VCFTools

## Common software (other):

• Anaconda, Git, GitHub, Homebrew, JIRA, LATEX