DeannaMChurch

Genomics and Bioinformatics Leader

contact

focus

3538 22nd St. Boulder, CO 80304

Leading interdisciplinary teams using genomics and computation to accelerate biological dis-

USA covery.

+0 (301) 233 2991

experience

deanna.church@gmail.com

2019-2021

languages

english, mother tongue

Driving customer voice and commercialization of our mammalian editing platform. Ensuring our software

Genomics

Genome editing, Genome assembly, Genome annotation, variant analysis, & Single-cell 'omics

2014-2016

Bioinformatics

Python, R, HTML, XML/XSLT, Javascript, CSS3, SQL & Perl 2001-2014

1990-1992

education

1999-2001	National Center for Biotechnology Information, NIH Postdoctoral fellow Coordinated mouse genome resources.	Bethesda, MD
1997-1999	Laboratory of Janet Rossant, Mt. Sinai Hospital Postdoctoral fellow Applying genomics methods to understand velopmental biology	Toronto, Ontario ding mouse de-
1992-1997	PhD of Biological Sciences The University Phenotype-genotype correlation in the Cri-du-Chat region of ch	y of California, Irvine hromosome 5
1986-1990	Bachelor of Liberal Arts Research project focused on cell cycle regulation using yeast	University of Virginia

additional training

2010	Client-Side Web Development	University of Illinois (O'Reilly)
2009	XSLT/XPATH: Introduction and Syntax	Mulberry Technologies
2007	Introduction to XML	University of Illinois (O'Reilly)

advisory boards and professional service

2018-present	Genome Quebec	Scientific Advisory Board
2012-present	Ensembl	Scientific Advisory Board
2013-2016	The Genome Analysis Center/Earlham Institute	Scientific Advisory Board
2010-2014	Database of Genomic Variants	Scientific Advisory Board
2010-present	American Society of Human Genetics	Member
2008-2013	Advances in Genome Biology and Technology	Organizing Committee Member
2008-2013	International Standards Cytogenomic Arrays	Committee Member
2008-2012	International Mammalian Genome Society	Nomenclature Committee Member
2004-2012	International Mammalian Genome Society	Member
2006-2011	European Conditional Mouse Mutagenesis Program	Scientific Advisory Board
2006-2009	Genome Canada-MORGEN project	Scientific Advisory Board
2005	Mouse Genome Finishing Meeting	Organizer

awards

2012	For extraordinary effort in providing access to the 10	National Library of Medicine 000 genomes project data.
1997-1999	Human Frontiers Postdoctoral Fellowship Awarded to early career scientist to enable them to be country.	Mt. Sinai Hospital proaden their skills in a new
1986-1990	Echols Scholar Awarded to select undergraduates who exhibit acade	University of Virginia demic independence.

presentations

2018-Feb	Beckman Symposium, Technology Innovation and Human Genomics Invited Speaker, "The impact of technology on our view of biology"
2018-Jan	Polyploid Genome Workshop San Diego, CA Invited Speaker, "Linked-Reads for haplotype reconstruction and improved de novo assembly"
2017-Nov	Australasian Genomic Technologies Association Annual Meeting Keynote, "High resolution biology with 10x" Hobart, AUS
2017-Nov	Australasian Genomic Technologies Association Annual Meeting Keynote, "The impact of technology on biology" Hobart, AUS
2017-Apr	NGS Barcelona Barcelona, SP Keynote, "Advancing genomics drop by drop"
2017-Feb	Advance in Genomics and Biotechnology Abstract Selected, "More complete genome analysis" Hollywood, FL
2017-Jan	Personalized World Medicine Conference Speaker, "Expanding the reach of genomic analysis" Palo Alto, CA
2017-Jan	Plant and Animal Genome Conference San Diego, CA Invited Speaker, "Quantitative profiling of large and complex populations by single cell RNA-sequencing"
2017-Jan	Plant and Animal Genome Conference Invited Speaker, "Using Linked-Reads to enable efficient de novo, diploid assembly"
2016-Oct	American Society of Human Genetics Vancouver, CA Abstract Selected, "De novo assembly of individual human haplotypes from diploid samples"
2016-Oct	Genome Reference Consortium Workshop Invited Speaker, "Everyday de novo assembly" Vancouver, CA
2016-Sep	Genome Informatics Hinxton, UK Abstract Selected, "Improving genome analysis using Linked-Reads"
2016-Aug	Cancer Genomics Consortium Denver, CO Invited Speaker, "Complete genome analysis"
2016-May	Sequencing and Finishing in the Future Santa Fe, NM Abstract Selected, "Improving genome analysis using Linked-Reads"
2016-May	Rady Symposium San Diego, CA Invited Speaker, "Advancing Precision Medicine with Linked-Reads"
2016-Feb	Stanford Big Data in Genomics Invited Speaker, "Truly personalized genomics" Palo Alto, CA
2015-Oct	Genome informatics Cold Spring Harbor, NY Invited Speaker, "The Rumsfeldian challenge of developing clinical sequencing tests"
2015-May	The 11th International Workshop on Advanced Genomics Speaker, "Dissecting the diagnostic yield in exome sequencing" Tokyo, JP
2015-Mar	American College of Medical Genetics Annual Meeting Salt Lake City, UT Invited Speaker, "Technical limitations of variant identification and annotation"
2015-Feb	Advance in Genomics and Biotechnology, Pacific Biosciences Workshop Marco Island, FL Invited Speaker, "Finishing genomes, why does it matter"
2014-Nov	Biological Data Sciences Cold Spring Harbor, NY Keynote Speaker, "Analog reporting in a digital age"
2014-Oct	American Society of Human Genetics San Diego, CA Abstract Selected, "The impact of GRCh38 on clinical sequencing"
2014-May	Sequencing and Finishing in the Future Santa Fe, NM Abstract Selected "Dissecting the missing diagnostic yield in exome sequenc-

ing"

publications

articlearticle in peer-reviewed journal bookbooks miscother publications reportresearch reports