Overview The ability of organisms to adapt to new and changing conditions is vital in today's world of climate and other anthropogenic changes. Understanding and predicting this ability to adapt for species is thus key, and is especially relevant in crop species that are needed to sustain food and fuel sources for the world's growing populations. This research project aims to examine the genetic architecture of important traits for adaptation in maize before and after its domestication bottleneck. Two major questions will be addressed: 1) what is the distribution of effect sizes for mutations underlying important phenotypic traits in maize and teosinte, and 2) how do different histories of demography and selection change this genetic architecture over time and impact the species' ability to adapt? Vast resources of genomic and phenotypic data in both teosinte and maize create an ideal system in which to investigate these questions. I propose three objectives to answer these questions using both real and simulated data to create and test theoretical predictions in the system. The first objective is to estimate the distribution of fitness effects of new mutations in teosinte, maize's progenitor. From this result, simulations in objective two will be parameterized and run to simulate the demographic and selective history of maize during its domestication. This creates an expectation for how the genetic architecture of traits in maize, hypothesized to be either under selection during the domestication event or not, should change given its past and can be compared to real data to see if predictions match reality. Objective three then performs further simulations of additional demographic and selective events that varying landraces of maize have experienced to assess if further changes to genetic architecture are detected. This research will be conducted at the University of California, Davis, under the supervision of Dr. Jeffrey Ross-Ibarra and cosupervised by Dr. Kevin Thornton at the University of California, Irvine.

Intellectual Merit The proposed research will greatly expand our understanding of the adaptability of quantitative traits and the relationship this has with the architecture underlying such quantitative traits. Little is known about the genetic architecture of quantitative traits, yet most important traits are quantitative. With the availability of genomic and phenotypic data, this is an area now ripe for further research A standing question in evolutionary biology is whether adaptation generally happens from few loci of large effect or many loci of small effect. This is a difficult question to answer, as the detectability of small effect loci is limited. Traditionally, QTL and GWAS approaches can be used to find important genes underlying traits of interest, but suffer from the aforementioned problem of small effect size loci. This research proposes to resolve this issue and provide deeper insights into the genetic architecture of adaptation. Additionally, conducting this research in the maize/teosinte system informs future research on this important crop and breeding of the species for adapting to future climate change as well as advances our understanding of plant biology and plant genomics. The results could inform crop studies in terms of maintaining genetic diversity for the future in ways that may have much larger long term impacts on the maintenance of diversity as well as on adaptation to changing environments.

Broader Impacts The impacts of this research will span across both the scientific community and the broader public. The research components and results will be made publicly available through online repositories of simulation and analysis code as well as public archiving of the data used. Additionally, I will make an effort to publish results from this work either in open-access journals or on public pre-print servers for other journals. I have a strong record of conference attendance that I will maintain in order to present results to the scientific community, and will also use my presence on Twitter and as a contributor to The Molecular Ecologist online blog to more frequently reach both the scientific and public audiences. Furthermore, I will mentor undergraduate students in the lab and also be able to participate in the educational partnership that has been created between UC Davis and Pioneer High School in Woodland, California. This will allow me to visit high school classrooms to teach about the cutting edge of genomics and evolutionary biology research to young scientists as well as to potentially mentor high school students in the lab or on small-scale projects.

found this here:
http://www.
dailydemocrat.
com/general-news
20140307/

<mark>ic-davis-partnersh</mark>ip-with-pioneer-high-benefits-budding-scientists

Jeff, do you have any collaborators in Mexico or S. America where I could go and teach some sort of workshop on genomics or computational skills for year 3 or something? if so, could be a good thing to propose here, or does that not make sense if they're more doing the field work with maize?

## **Project Description**

#### A. Introduction

A critical goal of evolutionary biology today is to understand organisms' abilities to adapt to new or changing conditions. This is especially relevant in the face of global climate change and other increasingly common anthropogenic changes to the environment (Easterling, 2000). Many important phenotypic traits are quantitative (need a ref?), and in order to understand adaptation of such traits, we must understand the genetic architecture underlying them and how this architecture may be changed as a result of selection and demography. Such knowledge advances our understanding of the process of adaptation and can further benefit crucial goals such as improving crop yields under changing climates.

in addition we want to know whether standing variation is maintained by selection (possibly adaptive) or at mutation/selection balance (load)

in what sense, pe

Genetic architecture can be affected by many factors, including population size, the number of genes, demographic history, and the strength of positive selection and its consistency through time. Demographic history alone plays a major role in the strength of selection on traits and organisms. For example, human demographic history of changing population sizes, migration, and expansion out of Africa have left lasting signatures in the genome (Fu et al., 2014; Gravel et al., 2011; Henn et al., 2015). Domestication bottlenecks have also been shown to impact the genome (e.g. in sunflowers (Renaut and Rieseberg, 2015), torsten - rice?). Such bottlenecks greatly decrease the effective population size  $(N_e)$  which can increase the effect of random genetic drift. The effect size, or strength of selection (s), for a given locus, can also impact the efficiency of selection across the genome (selection effective when  $N_e s > 1$ ). Thus, as demography changes, so does  $N_e s$  and therefore so does the occurrence of different beneficial, deleterious, or neutral loci across the genome.

Many traits ecologically important for local adaptation have a complex, quantitative genetic basis, and much heterogeneity has been found among these traits both within and among species (Orr, 2001; Slate, 2005). The architectures considered in the proposed research concern the number of underlying loci and their various effect sizes and dominance relationships. Whether this genetic architecture consists of many loci of small effect or few of large effect can play a role in the impact of demography and selection, as well as mutation, on the genome. How the genome may be restructured as a result of various demographic histories and those effects on selection is not well studied, nor how new genes may evolve to underly traits of importance (Long et al., 2003). The genetic architecture underlying traits can affect how easily or quickly local adaptation may occur, or how long and difficult that process may be (Yeaman, 2015; Yeaman and Whitlock, 2011). Such knowledge can therefore greatly contribute to improve breeding and conservation efforts as well as for predicting responses to environmental changes.

better reis:

refs?

My proposed research aims to investigate these important factors (demographic history, selection, and genetic architecture) and their effects and interactions in terms of shaping the architecture of quantitative traits in maize. Maize (*Zea mays*, ssp. *mays*) was domesticated from its wild teosinte ancestor, *Zea mays*, ssp. *parviglumis*, approximately 9,000 years ago in southwestern Mexico (cite Matsuoka et al 2002). Demographic history of the species is supported by historic records of its use and domestication in the Americas (cite). Extensive genomic data is also available, allowing the known demographic history of the species to be examined in terms of genomic signatures of domestication, as well as specifically parameterized to match this demographic and selective past in terms of important values such as effective population sizes, or migration rates. This provides a unique opportunity to compare these closely related subspecies which have undergone various different demographic processes contributing to their present-day genetic make-up.

I propose three objectives which make use of the extensive knowledge and data available in the maize/teosinte system in order to answer the question of **how the genetic architecture of traits changes as a result of demography and selection**. These approaches overcome issues of

similar past studies using GWAS and QTL approaches which suffer from limitations due to inability to detect small effect alleles. Objective one uses existing genomic data in teosinte to characterize the genetic architecture of phenotypically important traits and find the distribution of effect sizes. Objective two then simulates the demographic history of teosinte's domestication into modern-day maize, parameterized from results of objective one. Objective three assesses changes in the distribution of effects sizes that resulted from selection and demography by comparing the results from objective two's simulations to the reality of modern-day maize populations. This research will further our understanding of demography and selection's impacts on the genome and its structure, as well as our ability to detect and predict changes in genetic architecture after certain demographic events.

## B. Research Objectives, Methods & Significance

## Objective I: Investigate the genetic architecture of important traits in teosinte

This first objective aims to estimate the distribution of fitness effects (DFE) in teosinte (*Zea mays* ssp. *parviglumis*), the wild progenitor of domestic maize, and translate this into a distribution of effect sizes. The DFE describes the fitness effects of various mutations that are possible within the genome. Mutations can broadly be classified as beneficial, deleterious, or neutral, but in actuality, mutation effect sizes span a continuum of strongly deleterious to strongly beneficial, with any value in between. The software DoFE (Keightley and Eyre-Walker, 2007) uses of the number of nonsynonymous and synonymous substitutions within genes to calculate the DFE. I will apply this method to sequence data from regions of the genome known to underly important phenotypic traits. Some such quantitative phenotypes include yield, plant height, and flowering time, which are of critical importance to agriculture (cite). From this distribution of fitness effects and fitness measures from these traits, we can then translate these values into effect sizes per allele. I will use genotypic and phenotypic data on teosinte which is available for 5,000 individuals at 16 phenotypic traits (height, kernel traits...) (cite data source). These individuals are the progeny of 70 teosinte individuals sequenced to 25X coverage, providing an ideal resource for this analysis.

I will validate these parameter estimates by using them to parameterize simulations of a population of teosinte to an equilibrium state, and by comparing a genome-wide association study (GWAS) on these simulation results to real GWAS results on teosinte, can establish the accuracy of the parameter estimates. If this comparison retuerns vastly different estimates of effect sizes, I will adjust parameter values and resimulate until a sufficient match is made between real teosinte data and simulated data. The simulation program fwdpy (a Python implementation of fwdpp, Thornton (2014)) allows explicit modeling of the genomic architecture desired: distribution of effect sizes across the desired number of loci, including deleterious mutations and their effects on the phenotype. Values of  $V_G$ ,  $V_A$ , and  $V_D$  can be run to equilibrium, and then the populations sampled in order to test for successful replication of a wild teosinte population.

Characterizing the DFE of teosinte and translating this into a distribution of effect sizes uncovers whether the genetic architecture of important quantitative phenotypic traits in teosinte are underlain by many loci of small effect, few loci of large effect, or any combination therein. Estimating the DFE in teosinte will further inform a larger body of work aimed at understanding how common different genetic architectures are for traits important in adaptation. The DFE is difficult to estimate and not broadly understood in evolutionary biology for any given organism or in terms of how much it may vary across organisms. This difficulty arises particularly for cases where there are many loci of small effects: such small effect sizes are difficult to detect on an individual basis. This has implications for studies attempting to identify loci important in adaptation struggle when each locus out of many may only contribute a small amount to the trait (cite Outi Savolainen's work in Scots pine). Thus, studies that aim to identify loci important in adaptation will benefit from knowledge on

also: Fay, Wycoff and Wu (2001), Smith and Eyre-Walker (2002), Bierne and Eyre-Walker (2004) and Eyre-Walker and Keightley (2009), Stoletzki and Eyre-Walker (2011)

ek, not sure i m escribina this well

should list them

the range of mutation effect sizes they may expect to see when performing GWAS or genotypeenvironment associations.

# Objective II: Simulate the demographic history of maize domestication from teosinte

This portion of the proposed project aims to use the DFE results of Objective 1 to parameterize a model that will simulate the evolution of maize and its genetic architecture through time during and since its domestication. Maize originated approximately 9,000 years ago in southern Mexico (cite Matsuoka et al 2002, others?) during a single domestication event of *ssp. parviglumis*. Archaeological records also confirm this dating and single location (cite) as well as the subsequent spread and growth in population size of domestic landraces across the Americas into both low-land and highland environments [cite Wilkes, H. G. (1967) Teosinte: The Closest Relative of Maize (Harvard Univ., Cambridge, MA)]. South American landraces of maize also underwent a second bottleneck event during their expansion (cite). Using genetic data, the precise demographic parameters of this history have been estimated (cite Beissinger et al in prep). This provides information on the ancestral effective population size of maize ( $N_a \approx 120,000$ ), the size to which the population was bottlenecked during domestication (5% of this  $N_a$ ), the subsequent size to which populations rapidly expanded (3 times as large as  $N_a$ ), and lastly on the genome-wide mutation rate ( $3.8 \times 10^{-8}$  cite Clark et al 2005 MBE 22, 2304 – 2312.)

The information described above can be used to parameterize simulations of maize genomes undergoing this same demographic history. The DFE of teosinte established from objective 1 provides the necessary parameters for the number of loci and their various effect sizes to be modeled as contributing to each phenotypic trait of interest. Population sizes can be matched to those known during the domestication process, enforcing the desired demographic history, and mutations can be induced at the inferred rates to create simulations as accurate to reality as possible. The forward simulation program fwdpy (a Python implementation of fwdpp, cite) provides the framework for performing these simulations of complex demographies with natural selection. This approach allows explicit modeling of the genomic architecture desired: distribution of effect sizes across the desired number of loci, including deleterious mutations and their effects on the phenotype. Recombination parameters, as well as setting population sizes, and even sampling from simulated populations is possible.

I will simulate the various landraces of maize which each have a different demographic history during their spread across Central and South America into both lowland and highland environments. I can then compare the different genetic architectures that evolve from these various conditions. From the simulation results, I will evaluate how many loci are found to contribute to phenotypically important traits (on average and how variable this number is), how strong the mutational effects are at each locus and if and how this relates to the number of loci contributing. This will allow assessment of how important the details of demography are in determining the genetic architecture of local adaptation to different (lowland vs. highland) or similar (lowland Mexico vs. lowland S. America) conditions.

Objective 2 will inform whether the DFE changes under certain demographies, and if so whether there is any meaningful direction in this change. We may hypothesize changes to a wider or narrower or more skewed distribution of fitness effects depending on the demography at hand. Population bottlenecks are a common demographic occurrence during the geographic spread of populations (cite) which will be included in our simulations. Such bottlenecks reduce population sizes greatly, and this reduction can have several effects on the genome including purging of deleterious alleles (recessive alleles become homozygous and are more efficiently removed), or alternatively may lead to an increase of some deleterious alleles through increased random genetic drift (allele surfing, cite Klopfstein et al 2006). The effects of these processes varies depending on the degree

TL/GWAS. this good for finding genes, but has limitations for understanding process because power, small fx. we propose to resolve this by 1) using popness effects of new mutations and thus effects on a fitnessrelated phenotype 2) simulation-based mate genetic architecture of teosinte 3) w/ architecture known demography to see if demography alone explains maize, or demography + selection

part we need K-Thor for: turning s from DoFE into effect size for quant trait

but maybe as muc as 1E9, citation

will want to think about how to make the comparison.

have/need info on recombination/linkage?

this is cool idea. we will need to esti mate demography. can use dadi and cite Takuno et al. for basic model. or try MSMC and cite Beissinger? this could be cool 3rd objective, looking at how geography affects. not well done in human lit, but good popgen (as you're aware) describing how this

I think this could be interesting, even if we can't compare them all in the end to real data since we only have it for some - unless it might be in the scope of this project to try and generate

this paragraph great for intellectual merit of population size reduction and the length of time over which populations are bottlenecked (cite bottleneck lit). The second bottleneck of S. American populations may thus result in a difference between landraces in terms of genetic architecture of adaptive traits.

These hypothesized changes in the DFE are not well understood in any system and are controversial in humans (cite lohmueller vs. pritchard etc.). Deleterious alleles likely play a large role in many adaptive phenotypes: crop plants have undergone dramatic demographic shifts, usually involving a domestication bottleneck followed by expansion as cultivation spread, and some authors even argue that selection on domestication traits has inadvertently increased the frequency of alleles deleterious for other phenotypes (cite gunther2010). Consistent with this, it has recently been shown that genes associated with a number of quantitative traits in maize are enriched for deleterious alleles compared to randomly chosen genes (cite mezmouk2014). Such information is crucial for understanding variation in phenotype, designing breeding strategies, utilizing diversity from wild relatives, or even engineering new traits using biotechnology.

# Objective III: Estimate and compare the genetic architecture underlying traits in maize post-domestication from simulations and genomic data

With the DFE of teosinte estimated in Objective 1 and the simulated evolution of this DFE from teosinte into domesticated maize in Objective 2, we now are primed to compare the predictions of Objective 2 with real world data from modern maize. Similar to Objective 1's approach, Objective 3 will estimate the DFE of modern maize (*Zea mays mays*). The expectation is that if the estimated demographic model and DFE are reasonable, the genetic architecture of simulated phenotypes should closely mimic that of real data found here. If these results are not recapitulated in the maize data, then this indicates that there are complexities in the demographic history and/or our genomic model parameters that are not well enough understood currently. In this case, we can explore the sensitivity of genetic architecture to changes to the demography or the DFE. Understanding this sensitivity will then lead to improved estimation of these important parameters for the future.

We will recapitulate the methods described in Objective 1 using genomic data from 55 sequenced individuals from a maize population as well as 2,500 individuals from the GWAS datasets.

to chat to you more about the data Jeff, realizing I didn't take good enough notes on things

broader impacts: simulation pipeline for arbitrary DFE, architecture, selection. claim method to fit to data (do we make that part of the proposal)? info useful for breeders – learn impact of breeding strategies on traits?

# C. Training Objectives

This research fellowship will provide me with an ideal opportunity to learn the skills I need to enhance my research ability in the fields of genomics and computational biology, both areas in which I expect to continue my future research and which are greatly expanding in evolutionary biology. I will gain many skills related to genomic data analysis through Objectives 1 and 3, learning bioinformatics and analysis skills for large datasets. I have minimal exposure and experience directly working with such data from my dissertation, thus making this a vital step in my career. Genomic technology and data are growing at an incredibly fast pace, and working directly with such data will teach me the most up to date, accurate, and efficient approaches. I will also improve my skills of computational biology through the proposed simulations in Objective 2 and be able to learn a new and useful programming language used widely in evolutionary biology, Python.

# D. Career Development & Future Research

My career goal is to become an independent, academic researcher who is able to push the boundaries of population genetics and evolutionary biology. As I have in my dissertation research, I aim

careful, you don't want to make this sound like an extension of your PhD

somewhere here maybe talk about how with increasing data, pop and quant gen are becoming same?

think of bett

i like the open science angle

if you are considering alternative careers, i'm rather proud of fact i have good track record helping pdocs get jobs: 3 assistant profs, 2 in seed industry, 1 in NGO, 1 govmt. research scientist

yes, I definitely like this, in my head am also considering non-academic careers but worry about coming off as undecided in the proposal?

Jeff, feel free to make that sound better

Likewise Kevin, fe free to modify

should probably ac something about Farm and Kevin's to study population genetic and genomic processes both empirically from real world data and theoretically through simulations. I believe one of the strongest ways to advance our knowledge is through such comparisons of situations where every parameter, current and historical, is known, as well as the evolutionary outcome (simulations) to natural situations where processes or effects that are poorly understood or still unknown to us can diverge from theoretical expectations and provide the basis for further study and investigation into these processes.

The skills I will develop during this fellowship, as described in section C, will benefit my career and put me on the cutting edge for analyses of the newest genomic data and the most recent computational approaches for biological simulations. Interacting with Dr. Ross-Ibarra, as well as other researchers at UC Davis, and with Dr. Kevin Thornton at UC Irvine, on a regular basis will be both intellectually stimulating and rewarding experiences that will help me accomplish my career goals. Drs. Ross-Ibarra and Thornton are both at the forefront of a popular movement for open science, making all stages of the research process transparent to any interested parties, and providing products such as data and code immediately and publicly. This is a work ethic I strongly agree with and hope to contribute to as an independent researcher. Our work together will better equip me with the tools and experience that make open science easy, efficient, and profitable for all. I believe that this will equip me as a competitive, knowledgeable, and independent researcher able to conduct interesting and useful research throughout my future research program on topics of local adaptation, demographic history, population structure and genetic architecture of important traits.

# E. Sponsoring Scientists and Host Institution

The University of California Davis (UCD) is the ideal place to conduct the proposed research. UCD has a world-renowned program in evolutionary biology and faculty in population genetics who are at the top of the field. Jeff Ross-Ibarra is an expert on teosinte, maize, its domestication, and the associated population genetics and genomics of the system. Kevin Thornton is an accomplished quantitative geneticist and computational biologist at UC Irvine, who will also contribute greatly to this research. They will both serve as effective and capable mentors for my post-doctoral research. In particular, Jeff has been studying the maize/teosinte system for many years with a great network of collaborators providing vast resources of data. His work has contributed largely to our knowledge of this system, and more generally on domestication and adaptation as evolutionary processes. Kevin is also the developer and maintainer of fwdpy, the python package proposed for completing the simulations in Objective 2. He will thus serve as a great resource in terms of knowing the exact capabilities of the simulation method and any assumptions of its model that must be taken into account. Furthermore, the Department of Ecology and Evolution, the Department of Plant Biology, and the Department of Plant Sciences at UCD have many exceptional faculty doing research relevant to my interests, providing many research groups to interact with on a daily basis for potential collaborations or feedback on this research. For example, I look forward to interacting with scientists interested in population genetics, such as Graham Coop, and in adaptation, such as Johanna Schmitt. UCD has the necessary computing resources for our proposed work, and as described, vast sources of knowledge and experience on the topics I plan to investigate, ensuring the success of this work. I am excited to join and contribute to UCD's active and vibrant scientific community.

## F. Milestones & Timeline

definitely we go for year 3!

want to do the DFE of maize and teosinte at the sam time? no reason we can't right, and then do the simulations. Though if it takes time, teosinte should be prioritize so the sims can be parameterized and started while the maize DFE is estimated

yup, should be easy to do both at same time.

yeah, we might think about what advantages we have over humans and how human research can benefit from plants!

Year 1	Estimate DFE in teosinte, get simulations working and compare to teosinte
Year 2	finish comparison to teosinte, simulate maize
Year 3	expand to other datasets with demography + local adaptation + expansion?

## G. Broader Impacts

The proposed research will have wide-ranging impacts for both the public and the scientific community. I will ensure that my results are available to the public at all stages of these projects by maintaining code and scripts online at my GitHub account, which will allow other researchers to access analysis methods or data cleaning tools as well as simulation details and parameters which can provide a building block from which further research can be conducted. I will present new findings at international conferences and submit publications to open-access pre-print servers. I will also be able to broadcast my work more widely to the public through a strong online presence I maintain on Twitter, blog posts I can contribute to The Molecular Ecologist, a blog I have contributed to in the past. The wide-ranging impacts of this research on corn as a crop species useful for both food and fuel resources is also of an innate broad impact, as understanding the genetics underlying adaptation will ensure a viable future for such crop species in the future. Lastly, this research will contribute greatly to my own career development, improving my knowledge on genomics and working in an economically important crop species. I will be able to learn these vital tools as well as to teach them to undergraduate students in the lab and into the future as the field of genomics continues to grow.

genetic architecture underlying traits affects how easy/hard and quick/slow local adaptation can occur. important for breeding, conservation, predicting response to climate change, etc. useful info for crops/domesticated species (and also things like disease in humans?)

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## Biographical Sketch — Kimberly Julie Gilbert

#### A. Professional Preparation

Institution, Location	Major	Degree	Year
University of Virginia, USA	Biology	B.Sc.	2010
University of British Columbia, Canada	Zoology	Ph.D.	2016 (expected)

#### **B. Publications**

#### Five Publications Most Closely Related to the Proposed Project

- 1. **Gilbert KJ**, MC Whitlock (2015) Evaluating methods for estimating local effective population size with and without migration. *Evolution*, 68(8), 2154-2166.
- 2. **Gilbert KJ**, MC Whitlock (2015)  $Q_{ST}$ - $F_{ST}$  comparisons with unbalanced half-sib designs. *Molecular Ecology Resources*, 15(2), 262-267.
- 3. Caplins SA, **KJ Gilbert**, C Ciotir, J Roland, SF Matter, N Keyghobadi (2014) Landscape structure and the genetic effects of a population collapse. *Proceedings of the Royal Society B*. 281: 20141798; doi: 10.1098/rspb.2014.1798
- 4. Keller SR, **KJ Gilbert**, PD Fields, DR Taylor (2012) Bayesian inference of a complex invasion history revealed by nuclear and chloroplast genetic diversity in the colonizing plant, *Silene latifolia*. *Molecular Ecology*, 21(19), 4721-4734.
- 5. Whitlock MC, **KJ Gilbert** (2012) *Q*s⊤ in a hierarchically structured population. *Molecular Ecology Resources*, 12(3), 481–483.

#### Four Other Significant Publications

- 1. Santiso X, L Lopez, **KJ Gilbert**, R Barreiro, MC Whitlock, R Retuerto (2015) Patterns of genetic variation within and among populations in *Arbutus unedo* and its relation with selection and evolvability. *Perspectives in Plant Ecology, Evolution and Systematics*, 17(3), 185-192.
- 2. Vines TH, RL Andrew, DG Bock, MT Franklin, **KJ Gilbert**, NC Kane, EJ Kleynhans, J-S Moore, BT Moyers, S Renaut, DJ Rennison, T Veen, S Yeaman (2013) Mandated archiving greatly improves access to research data. *FASEB Journal*, 27(4), 1304-1308.
- 3. **Gilbert KJ**, RL Andrew, DG Bock, MT Franklin, NC Kane, J-S Moore, BT Moyers, S Renaut, DJ Rennison, T Veen, TH Vines (2012) Recommendations for utilizing and reporting population genetic analyses: The reproducibility of genetic clustering using the program STRUCTURE. *Molecular Ecology*, 21(20), 4925-4930.
- 4. Vines TH, AYK Albert, RL Andrew, F Débarre, DG Bock, MT Franklin, **KJ Gilbert**, J-S Moore, S Renaut, DJ Rennison (2014) The availability of research data declines rapidly with age. *Current Biology*, 24, 94-97.

## C. Select Conference Presentations (chosen from 11 presentations)

- 2015 Validating SNP loci underlying local adaptation in lodgepole pine; KJ Gilbert, S Yeaman, KE Lotterhos, KA Hodgins, H Suren, JA Holliday, S Nadeau, SN Aitken, MC Whitlock *Poster,* 15<sup>th</sup> ESEB Congress. Lausanne. Switzerland
- 2014 Evaluating methods for estimating effective population size in the presence of migration; KJ Gilbert & MC Whitlock *Oral presentation, Evolution, Raleigh, USA*
- 2012 Range expansion and adaptation across heterogeneous environments; KJ Gilbert & MC Whitlock Poster, Evo-WIBO Conference (Evolutionary Biology in the Pacific Northwest), Port Townsend, USA
- 2011 Inferred invasion history of *Silene latifolia* into North America utilizing population genetic data and approximate Bayesian computation; KJ Gilbert, SR Keller, PD Fields, DR Taylor *Poster,* 13<sup>th</sup> Congress of the European Society for Evolutionary Biology, Tuebingen, Germany

#### D. Grants and Awards

Cordula and Gunter Paetzold Fellowship UBC \$18,000CAD

2015 - 2016





Declined; Zoology Graduate Fellowship UBC \$11,000CAD	2015 – 2016
Ann and William Messenger Graduate Fellowship UBC \$700CAD	2015
Zoology Graduate Fellowship UBC \$11,000CAD	2014 – 2015
Frieda Granot Graduate Scholarship in Interdisciplinary Research UBC \$200CAD	2013 - 2014
Theodore E Arnold Fellowship UBC \$7,750CAD	2013 - 2014
Patrick David Campbell Graduate Fellowship UBC \$8,050CAD	2013 - 2014
Declined; Zoology Graduate Fellowship UBC \$10,000CAD	2013 - 2014
BRITE Fellowship UBC \$10,500CAD per annum	2011 – 2013

#### E. Synergistic Activities

- 1. Working Groups: Participated in the NESCent Reproducible Science Hackathon (2014), a 21-member working group aimed at developing a curriculum and workflow for teaching reproducible science to researchers of any background. Participated in the SimBank NESCent Catalysis Meeting (2014) which was a 25-member working group to create a collection of openly available simulation results to facilitate testing of statistical population genetic and phylogeographic methods.
- 2. Teaching: Teaching assistant for Fundamentals of Evolutionary Biology (Fall 2012-Spring 2013) where I taught three sections per term of 45 students each and lead discussion-based tutorials. Teaching assistant for Fundamentals of Biostatistics (Fall 2013, 2014, 2015). Taught two sections of 70 students total in 2013, and in 2014 and 2015 served as the lab coordinator for 254 and 276 students enrolled in the course, respectively, while teaching one section of 36 and 35 students respectively.
- 3. Service: Served as a Graduate Student Council Member for the American Society of Naturalists (2013-2016, chair 2015-2016). Served as the graduate student representative on the 2014 evolutionary biology CRC2 job search for the Department of Zoology, University of British Columbia. Organize the Biodiversity Research Centre's weekly evolution discussion group (2014-2016), for students, post-docs, and faculty from the departments of Zoology, Botany, Forestry, and Fisheries to discuss current papers in evolutionary biology. Reviewer for Molecular Ecology Resources, Ecology and Evolution, Tree Genetics & Genomes.
- 4. Outreach: Volunteer mist-netting and bird banding with local Vancouver non-profit organization Wild Research (2013-2015) where I participated in winter, spring migration, and fall migration bird monitoring at Iona Island Bird Observatory, taught volunteers proper bird handling, aging, data collection, and mist net extraction techniques, and assisted in educating public visitors to the station about the species conservation and monitoring, and the general tasks of running a banding station.

#### F. Collaborators (Total: 25)

U Alberta Jens Roland; U Basel Peter Fields; U British Columbia Dan Bock, Diana Rennison; U Calgary Sam Yeaman; UC Davis Serena Caplins; U Cincinnati Stephen Matter; CIRB Paris Florence Débarre; Colorado State Brook Moyers; U Colorado Nolan Kane; U Coruna Rodolfo Barreiro, Lúa López; Kwantlen Polytechnic U Michelle Franklin; U Laval Jean-Sébastien Moore; Mol. Ecol. Managing Editor Timothy Vines; U Montreal Sébastien Renaut; U New England (Australia) Rose Andrew; UT Austin Thor Veen; Trent U Claudia Ciotir; U Santiago de Compostela Rubén Retuerto Franco, Xabier Santiso; U Vermont Stephen Keller; U Virginia Douglas Taylor; Western U Nusha Keyghobadi; Women's Health Research Institute Arianne Albert

Graduate Advisor (Total: 1) University of British Columbia Michael C. Whitlock

## Biographical Sketch — Jeffrey Ross-Ibarra

# 1 Professional Preparation

Institution	Area	Degree / Training	Dates
University of California Riverside	Botany	BA, MS	1998, 2000
University of Georgia	Genetics	PhD	2006
University of California Irvine	Genetics	Postdoctoral Research	2008

# 2 Professional Appointments

Position	Institution	Dates
Associate Professor	University of California Davis	2012-present
Assistant Professor	University of California Davis	2009-2012
Profesor de Asignatura	Universidad Nacional Autónoma de México	2001

## 3 Products

## Most Relevant to the Proposed Research

- Mezmouk S, Ross-Ibarra J (2014) The pattern and distribution of deleterious mutations in maize. (2014) G3 4:163-171
- Hufford MB, Xun X, van Heerwaarden J, Pyhäjärvi T, Chia J-M, Cartwright RA, Elshire RJ, Glaubitz JC, Guill KE, Kaeppler S, Lai J, Morrell PL, Shannon LM, Song C, Spinger NM, Swanson-Wagner RA, Tiffin P, Wang J, Zhang G, Doebley J, McMullen MD, Ware D, Buckler ES, Yang S, Ross-Ibarra J (2012) Comparative population genomics of maize domestication and improvement. NATURE GENETICS 44:808-811
- Cook JP, McMullen MD, Holland JB, Tian F, Bradbury P, Ross-Ibarra J, Buckler ES, Flint-Garcia SA (2012) Genetic architecture of maize kernel composition in the Nested Association Mapping and Inbred Association panels. PLANT PHYSIOLOGY 158: 824-834
- van Heerwaarden J, Doebley J, Briggs WH, Glaubitz JC, Goodman MM, Sánchez González JJ, **Ross-Ibarra J** (2011) Genetic signals of origin, spread and introgression in a large sample of maize landraces. PNAS 108: 1088-1092
- Ross-Ibarra J, Tenaillon M, Gaut BS (2009) Historical divergence and gene flow in the genus Zea. GENETICS 181: 1399-1413.

#### Additional Products

- Gerke JP, Edwards JW, Guill KE, Ross-Ibarra J, McMullen MD (2015) The genomic impacts
  of drift and selection for hybrid performance in maize. GENETICS In Press
- Takuno S, Ralph P, Swarts K, Elshire RJ, Glaubitz JC, Buckler ES, Hufford MB, and Ross-Ibarra J (2015) Independent molecular basis of convergent highland adaptation in maize. GENETICS 200:1297-1312
- Wills DM, Whipple C, Takuno S, Kursel LE, Shannon LM, **Ross-Ibarra J**, Doebley JF (2013) From many, one: genetic control of prolificacy during maize domestication. PLoS GENETICS 9(6): e1003604.
- Studer A, Zhao Q, **Ross-Ibarra J**, Doebley J (2011) Identification of a functional transposon insertion in the maize domestication gene *tb1*. NATURE GENETICS 43:1160-1163.

 Gore MA, Chia JM, Elshire RJ, Sun Q, Ersoz ES, Hurwitz BL, Peiffer JA, McMullen MD, Grills GS, Ross-Ibarra J, Ware DH, Buckler ES (2009) A first-generation haplotype map of maize. SCIENCE 326: 1115-1117.

## 4 Synergistic Activities

- Faculty Development Award in recognition of university service, 2015
- Editor, G3, PeerJ, Axios Reviews
- DuPont Young Professor 2012-2014 and faculty advisor DuPont Pioneer graduate student symposium in plant breeding 2012-present
- Functional Genetics of Maize Centromeres US-Mexico exchange program, 2011-present
- Presidential Early Career Award for Scientists and Engineers 2009

## 5 Collaborators and Other Affiliations

#### Collaborators and Co-editors (Total: 56)

Cornell U Peter Bradbury, Jeffrey Glaubitz, Susan McCouch, Qi Sun, Feng Tian, Sharon Mitchell; USDA-ARS Edward Buckler, Sarah Hake, James Holland, Sherry Flint-Garcia, Mike McMullen, Doreen Ware, Jode Edwards; U Southern California Peter Ralph; UC Davis Alan Bennet, Daniel Runcie, Ed Taylor, Graham Coop, Keith Bradnam, Ian Korf, David Neale, Amélie Gaudin; UC Irvine Kevin Thornton; Carnegie Institute Davide Sosso; Stanford Wolf Frommer; LANGEBIO Ruairidh Sawers; U Georgia Kelly Dawe; Arizona State Reed Cartwright; U Missourri James Birchler, Katherine Guill, David Wills; Beijing Genomics Institute Song Chi, Xun Xu; U Wisconsin John Doebley, Jiming Jiang, Shawn Kaeppler; Syngenta William Briggs; Monsanto Lisa Kanizay; Dupont Pioneer Andy Baumgarten, Justin Gerke, Oscar Smith, Tabare Abadie; U Minnesota Roman Briskine, Peter Morrell, Chad Myers, Nathan Springer, Peter Tiffin; MIT Mary Gehring; NC State Major Goodman; INRA Clementine Vitte, Maud Tenaillon; Brigham Young Clinton Whipple; Danforth Center Anthony Studer; Universidad de Guadalajara Jesus Sànchez Gonzàlez; Iowa State Carolyn Lawrence; U Hawaii Gernot Presting; UC Riverside Mitchell Provance

#### **Graduate Advisors and Postdoctoral Sponsors (Total: 3)**

UC Riverside Norman Ellstrand; U Georgia James Hamrick; UC Irvine Brandon Gaut

## Thesis Advisor and Postgraduate Sponsor (Total: 14)

**Postdoctoral:** *Iowa State* Matthew Hufford; *Graduate U Advanced Studies* Shohei Takuno; *U Oulu* Tanja Pyhäjärvi, *KWS* Sofiane Mezmouk; *Wageningen* Joost van Heerwaarden; *USDA* Tim Beissinger; *UC Davis* Kate Crosby, Sayuri Tsukahara, Simon Renny-Byfield, Jinliang Yang **Graduate:** Dianne Velasco, Paul Bilinski, Anna O'Brien, Michelle Stitzer

# Biographical Sketch — Kevin Richard Thornton

# 6 Professional Preparation

Institution	Area	Degree / Training	Dates
University of Puget Sound	Botany	BA	1997, 2000
University of Chicago	Genetics	PhD	2003
Cornell University	Genetics	Postdoctoral Research	2007

## 7 Professional Appointments

Position	Institution	Dates
Associate Professor	University of California Irvine	2012-present
Assistant Professor	University of California Irvine	2007-2012

## 8 Products

## Most Relevant to the Proposed Research

- **Thornton**, K. R. (2014) A C++ template library for efficient forward-time population genetic simulation of large populations. Genetics 98:157-166 PMID: 24950894
- Thornton, K. (2003) libsequence, a C++ class library for evolutionary genetic analysis. Bioinformatics 19(17): 2325-2327 PMID 14630667
- Thornton, K. R., A. J. Foran, and A. D. Long (2013) Properties and modeling of GWAS when complex disease risk is due to non-complementing, deleterious mutations in genes of large effect. PLoS Genetics 9: e1003258. PMID 23437004

#### **Additional Products**

- Cridland, J. M., K. R. Thornton and A. D. Long (2015) Gene expression variation in Drosophila melanogaster due to rare transposable element insertion alleles of large effect. Genetics 199: 85-93.
- Baldwin-Brown, J., A. D. Long, and K. R. Thornton (2014) The Power to Detect Quantitative Trait Loci Using Resequenced, Experimentally Evolved Populations of Diploid, Sexual Organisms. Molecular Biology and Evolution 31: 1040-1055. PMID 24441104
- Open-source software: http://molpopgen.github.io/fwdpp/
- Open-source software: http://molpopgen.github.io/libsequence/

# 9 Synergistic Activities

- Open-source software: http://molpopgen.github.io/fwdpy/ This software is unpublished, and will be a key resource for this proposal.
- Editor, G3

## 10 Collaborators and Other Affiliations

## Collaborators and Co-editors (Total: 7)

- Cornell University Andrew G. Clark
- North Carolina State University Trudy Mackay
- Princeton University Peter Andolfatto
- Rochester University Daniel Garrigan, Daven C. Presgraves
- UC Irvine Anthony (Tony) Long
- University of Kansas Stuart MacDonald

#### **Graduate Advisors and Postdoctoral Sponsors (Total: 2)**

University of Chicago Manyuan Long Cornell University Andrew G. Clark

Thesis Advisor and Postgraduate Sponsor (Total: 3)

Postdoctoral: Rebekah R. Rogers PhD Thesis Advisor: Julie M. Cridland, Jaleal S. Sanjak

# **Data Management Plan**

## **Data Types**

This proposal will generate genotype and full-genome sequence data, phenotype data, analytical code, germplasm, and publications.

Data Archiving, Plan for Sharing, Public Access Policy

Genotype and Sequence Data EDIT THIS WHOLE DOCUMENT

## **Dissertation Summary - Kimberly J. Gilbert**

A major obstacle in evolutionary biology is the difficulty of population genetic inference in the face of confounding factors, such as demographic history. My dissertation work has focused on several topics related to this broad area of research:

- 1. Evaluating the ability of statistical genetic methods to estimate effective population sizes in the face of migration (Gilbert and Whitlock, 2015)
- 2. Assessing the factors related to local adaptation at range edges during species expansion
- 3. Validating SNP loci under selection for adaptation to climate in lodgepole pine (*Pinus contorta*) Effective population size,  $N_e$ , is a fundamental parameter in population genetics, evolutionary biology, and conservation biology, yet its estimation can be fraught with difficulties. Several methods to estimate Ne from genetic data have been developed that take advantage of various approaches for inferring  $N_e$ . The ability of these methods to accurately estimate  $N_e$ , however, has not been comprehensively examined. This part of my dissertation work employed seven of the most cited methods for estimating  $N_e$  from genetic data (Colony2, CoNe, Estim, MLNe, ONeSAMP, TMVP, and NeEstimator including LDNe) across simulated datasets with populations experiencing migration or no migration. The simulated population demographies were an isolated population with no immigration, an island model metapopulation with a sink population receiving immigrants, and an isolation by distance stepping stone model of populations. We found considerable variance in performance of these methods, both within and across demographic scenarios, with some methods performing very poorly. The most accurate estimates of  $N_e$  can be obtained by using LDNe, MLNe, or TMVP; however each of these approaches is outperformed by another in a differing demographic scenario. Knowledge of the approximate demography of population as well as the availability of temporal data largely improves  $N_e$  estimates.

Species range edges have boundaries that cannot always be explained ecologically or geographically, which leaves the question of what evolutionary forces may prevent populations at range edges from adapting and expanding the species range further. A large body of theoretical work has investigated many evolutionary parameters' effects on local adaptation in edge populations, but one area lacking in research is that of the interaction of the landscape with the ability to locally adapt. This study investigates how more realistic, heterogeneous environmental gradients (compared to the linear gradients that previous studies investigate) may interact with dispersal distance and the effect size of mutations. I have simulated a range of parameter combinations that show a strong relation of mutation effect size on the ability to spread across the landscape. As environmental heterogeneity increases, migration load (reduction in fitness due to dispersal away from an area previously adapted to) increases, and local adaptation becomes more difficult, especially in smaller populations at the range edge, slowing the speed of expansion across the landscape.

A history of range expansion can confound many inferences that population genetics aims to understand. Identifying the loci that underlie traits contributing to local adaptation is one such inference that is a major goal in evolutionary biology today. The lodgepole pine ( $Pinus\ contorta$ ) is a major timber tree in the Pacific Northwest which has a history of expansion post-glaciation, and either one or putatively a second glacial refugia from which this expansion occurre1.9d. Climate change is spurring foresters to plant trees for future harvest that will be best adapted to future climates for optimal yield, hence identifying loci underlying adaptation to climate change is a key goal. I am conducting a validation study of SNP loci identified through GWAS, genotype-environment association, and  $F_{ST}$  outlier tests to assess how often these methods may produce false positives as a result of population structure and spatial autocorrelation of genetic clines due to range expansion with gradients in environmental variables (i.e. temperature and precipitation). I have sampled a provenance trial (common garden study) in British Columbia to compare performance of populations from a range of native temperatures (MAT -3.7°C - 11°C) planted across test sites of varying temperature (MAT -1.4°C - 5°C) from which I will be able to test if predicted alleles do indeed show increased performance in mature, natural-grown trees.