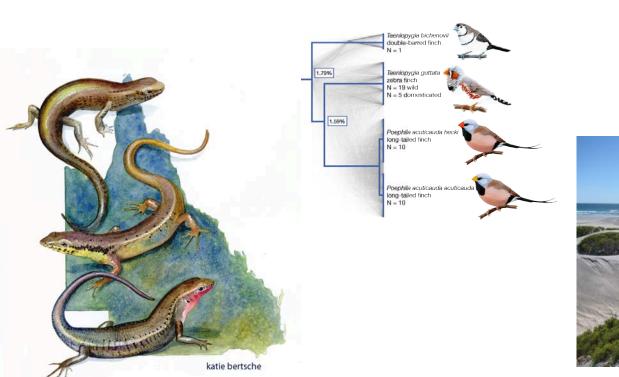


Rabosky Lab University of Michigan

My Background

- Demography, hybrid zones & introgression
- Fine-scale recombination rate estimates

Population genetics & macroevolution





Today's Plan

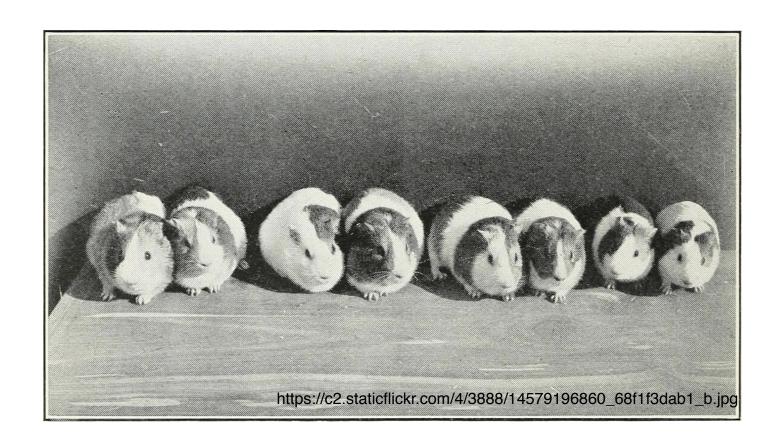
- What is population genomics?
- What are the possibilities and pitfalls?
- Sampling design
- Thought exercises
- Hands-on exercises
- Resource list

What is population genomics?

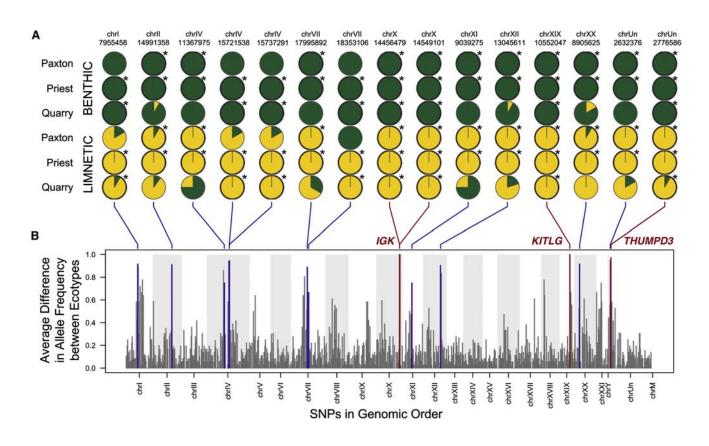
Population genomics has become a bit of an umbrella term.

- Genetic diversity (genotype calling)
- Linkage disequilibrium (phasing)
- Genetic differentiation across landscapes and the genome
- Population structure
- Population demography: parentage, inbreeding, etc.
- Reconstructing demographic history, i.e., expansions
- Molecular evolution
- Identifying targets of selection
- Often, phylogeography gets lumped in here

Don't forget the roots of population genomics: population genetics.



Get genome wide patterns! So fun. So interesting.



The cost margin is negligible compared to traditional approaches.



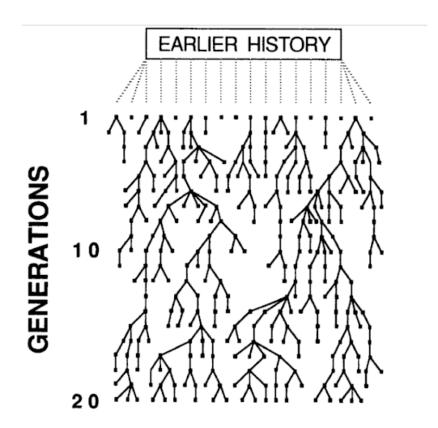
Move from spending time on collecting data to analyzing data.



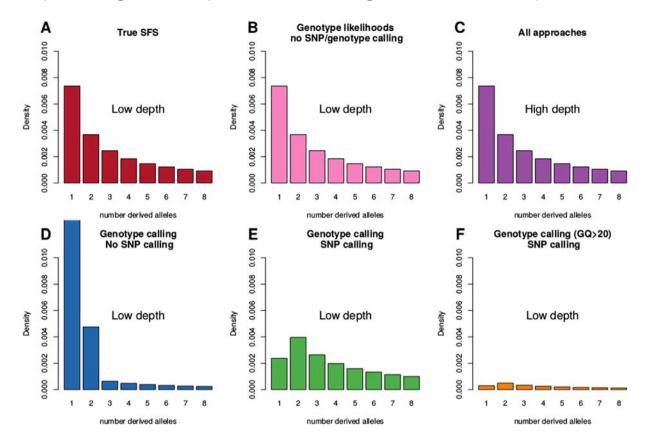




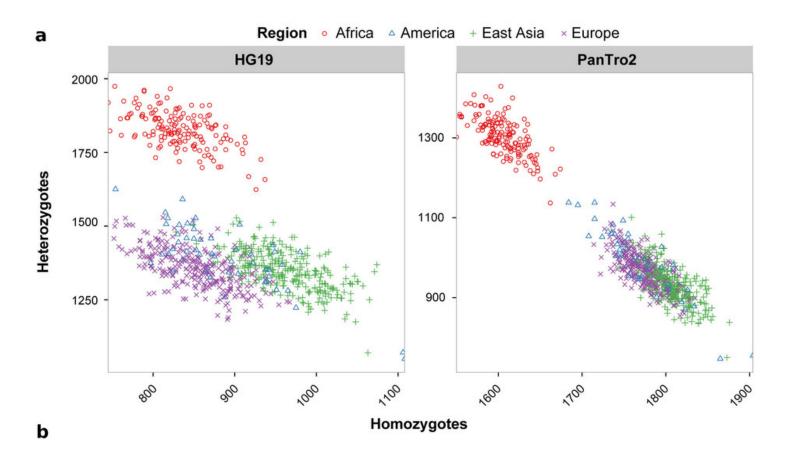
Every individual sampled is equivalent to sampling a population.



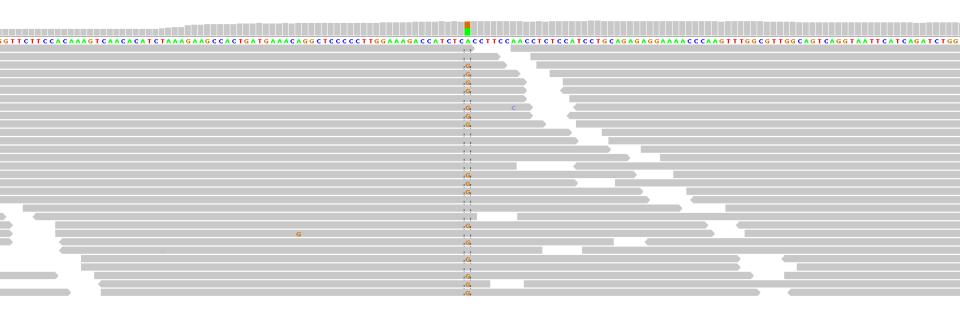
Accuracy of genotype calling: how do you filter?



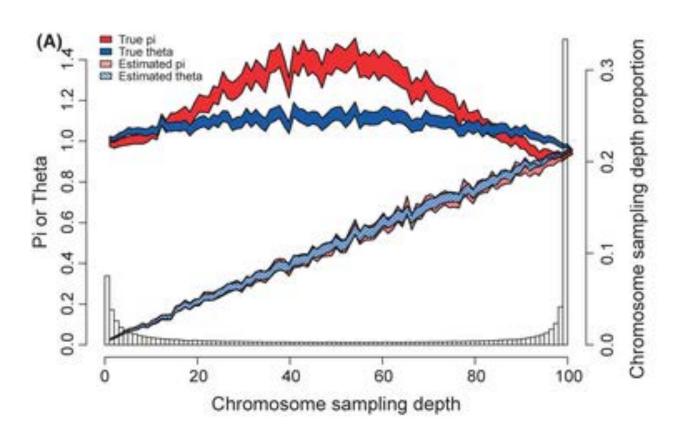
Bias, especially ascertainment bias and reference bias.



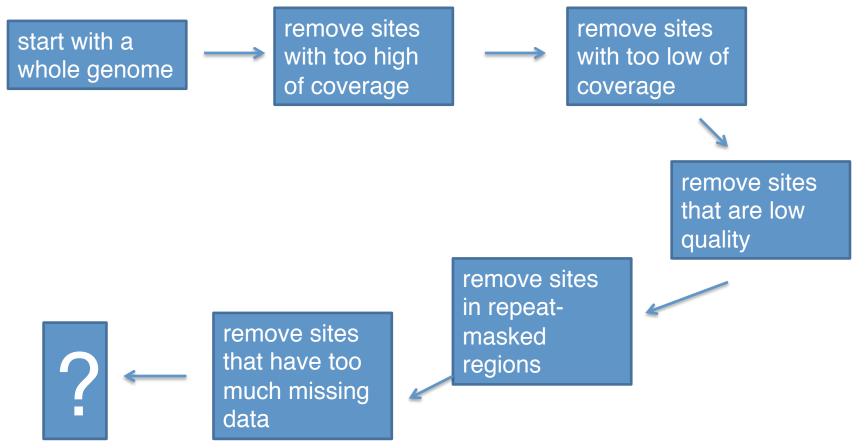
What should your reference be?



How do you handle missing data?

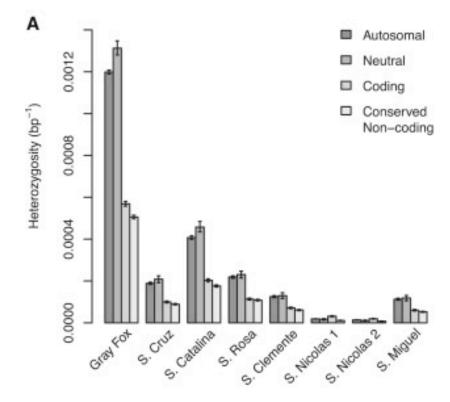


What's your denominator?

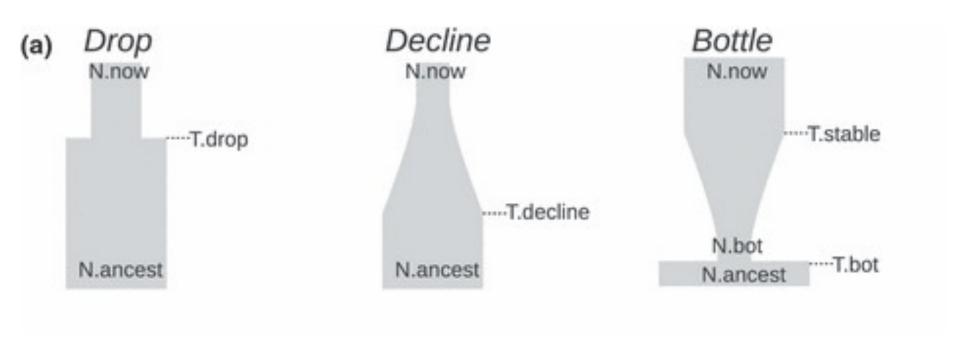


How do you annotate your variants?

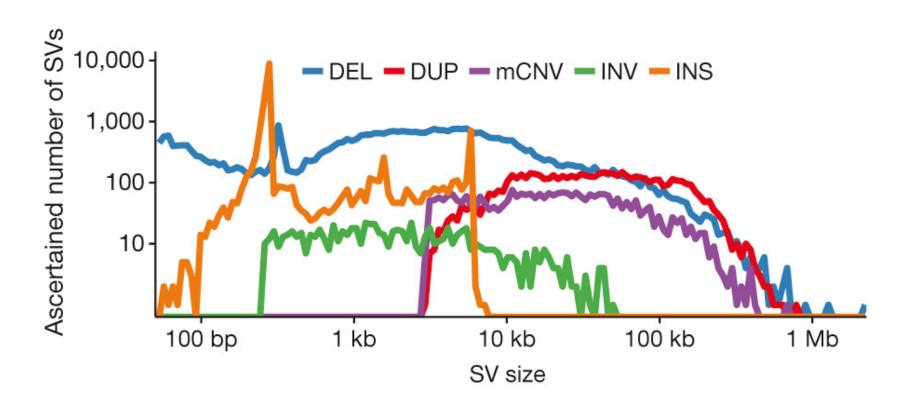




How do you define your neutral expectation?



Variation outside of SNPs still challenging to characterize.

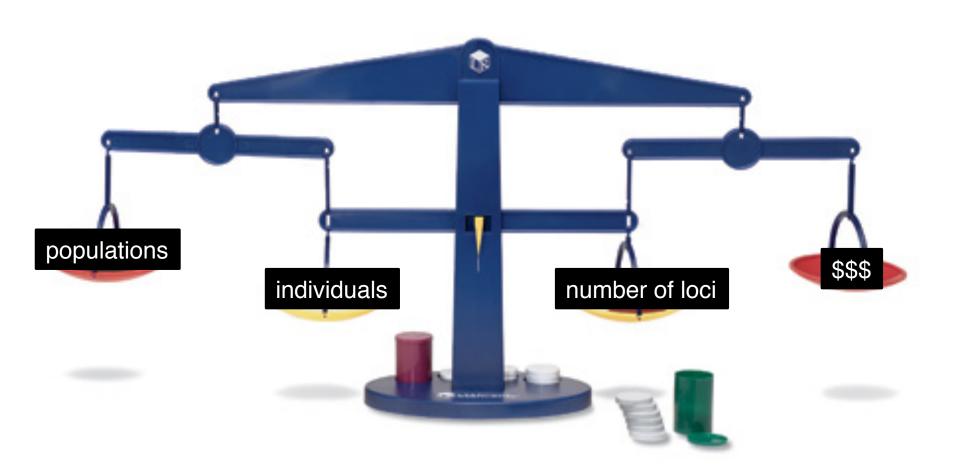


Sampling Design

- How many individuals?
- How many markers?
- What kind of markers are needed?
- What coverage is needed?
- Can anonymous pooling work?
- What is your phylogenetic scale?
- How much money do you have?

... because how you design your study depends on the answers to these questions ...

Sampling Design



Types of Loci

- whole-genome sequencing
- transcriptomes
- restriction-digest based approaches (RADtag, ddRADtag)
- targeted capture
 - exomes,
 - conserved elements
 - RAD loci
- genotyping SNPs using genotypers

You discovered a new endangered mammal species and want to characterize its genetic diversity to manage breeding programs.

How do you do it?



You are working on a widespread lizard species that you think likely consists of several cryptic lineages. You want to understand how many lineages there are and their demographic & divergence history. How do you do it?



You are working on an invasive plant species and want to know what the genetic structure is in the native vs. introduced range. How do you do it?



You are working on a fish that has a wide latitudinal gradient. You suspect that there has been local adaptation across the range. How do you test for this and identify the possible loci under selection?

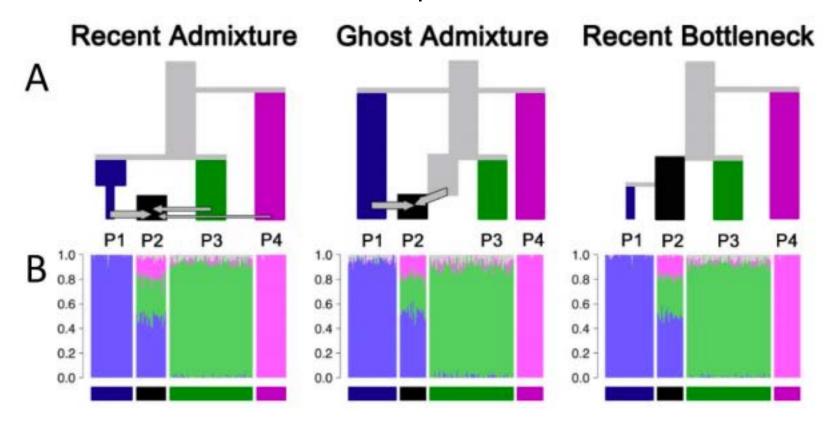


You are working on an insect that has experienced recent massive population declines due to pesticide residues in the environment. Some of its native range remains healthy, and a few susceptible populations have rebounded. How do both test for the genetic effects of these population declines and look for evidence of selective sweeps?



- 1. You discovered a new endangered mammal species and want to explore how much variation is contained in it. How do you do it?
- You are working on a widespread lizard species that you think likely consists of several cryptic lineages. You want to understand how many lineages there are and their demographic & divergence history. How do you do it?
- 3. You are working on an invasive plant species and want to know what the genetic structure is in the native vs. introduced range. How do you do it?
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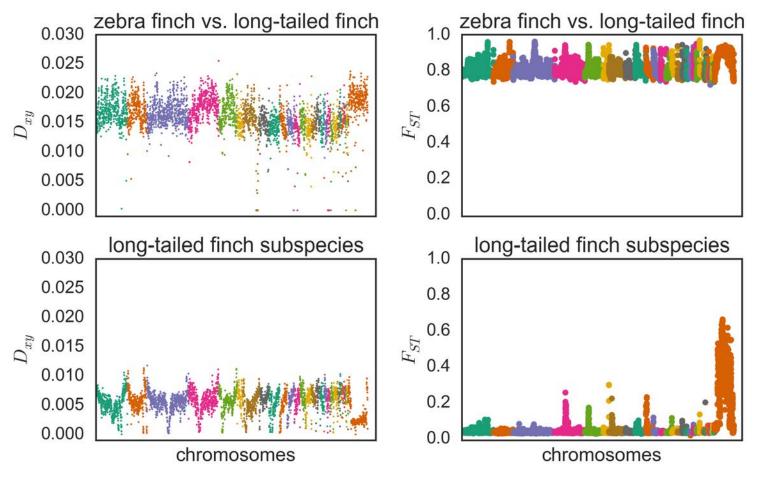
Distinguish between programs that are process-based vs. descriptive.



You will spend most of your time moving between file formats.



Get familiar with your data! Plot, plot, plot.



Almost all methodological innovations start in the human population genetic literature. Read it.

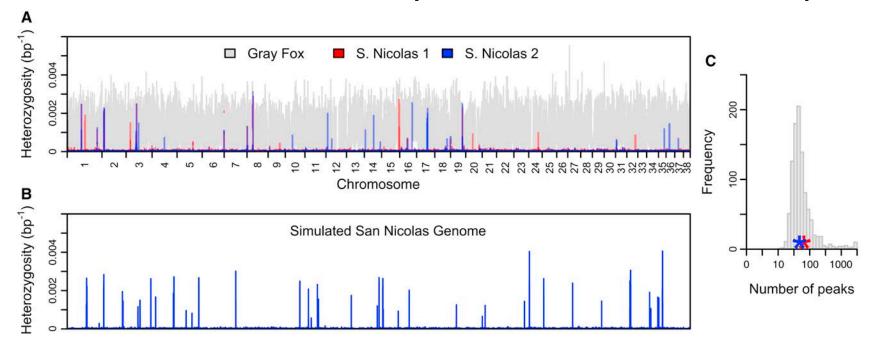


Methods and models for unravelling human evolutionary history

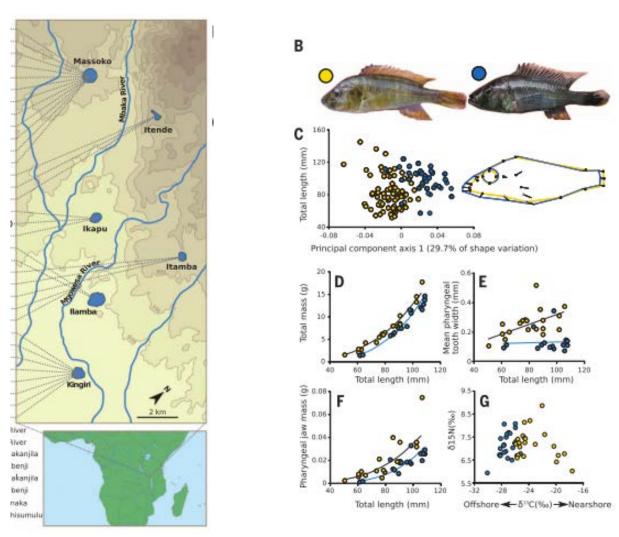
Joshua G. Schraiber and Joshua M. Akey

Abstract | The genomes of contemporary humans contain considerable information about the history of our species. Although the general contours of human evolutionary history have been defined with increasing resolution throughout the past several decades, the continuing deluge of massively large sequencing data sets presents new opportunities and challenges for understanding human evolutionary history. Here, we review the signatures that demographic history imparts on patterns of DNA sequence variation, statistical methods that have been developed to leverage information contained in genome-scale data sets and insights gleaned from these studies. We also discuss the importance of using exploratory analyses to assess data quality, the strengths and limitations of commonly used population genomics methods, and factors that confound population genomics inferences.

How do you bootstrap results? Simulations and non-parametric statistics help.



Tutorial: Cichlid fish



But first, let's look at a VCF!