



Computational
Biology @
Berkeley

Analysis of Dataset Equity and Genetic Diversity in the 1000 Genomes Project and TCGA

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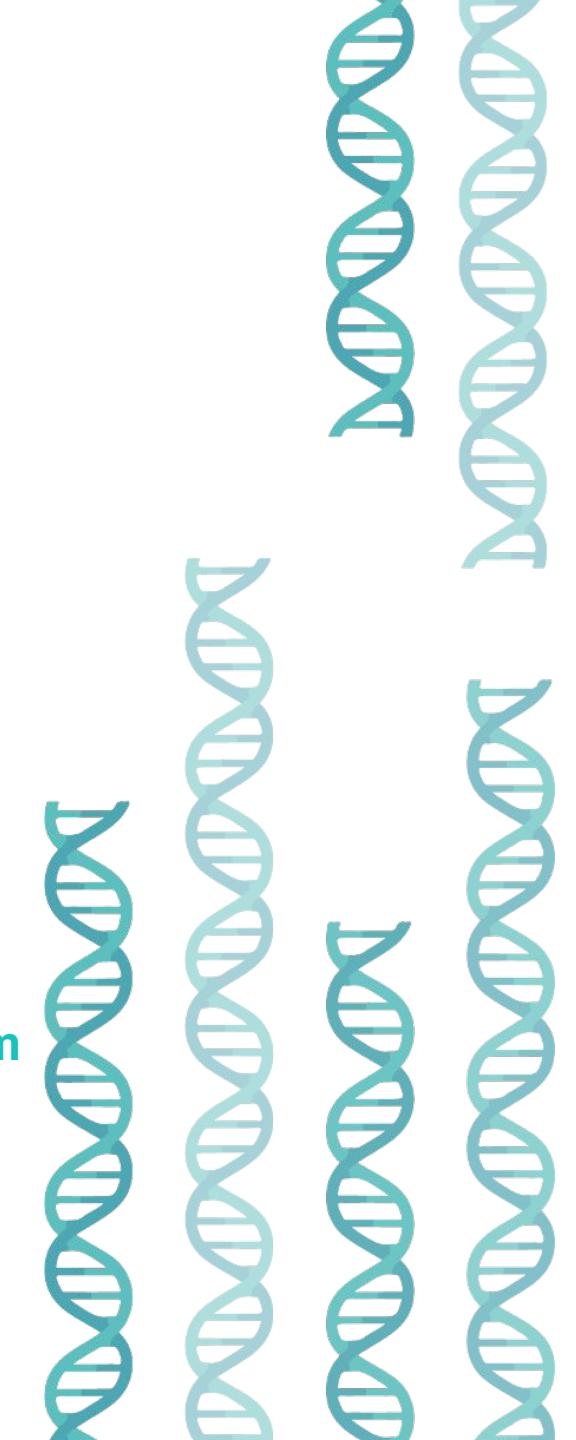


Table of Contents

I Research Background

II Hypothesis & Objectives

III Methodology

IV Results

V Conclusion & Discussion

Introduction

- Many genomic and biomedical datasets that are used in biology research tend to under represent certain populations
- This can lead to biased insights in downstream research
- Students and researchers alike tend to use these datasets without actually questioning how representative it is



Knowledge Gap

The 1000 Genomes Project and TCGA are both large scale genomics consortia that generate publicly available datasets

- The diversity in terms of populations and genetically is not quantified

Chose 1000 Genomes and TCGA as they represent opposite ends of the representation spectrum

Both are widely used benchmark datasets in genomics and precision medicine, so inequities or biases in their composition have real downstream impact on research, tools, and clinical translation

Questions

1. How are the various superpopulations represented within each dataset?
2. How do the various superpopulations vary genetically?
3. How do the datasets compare in regards to these metrics?

II Hypothesis & Objectives

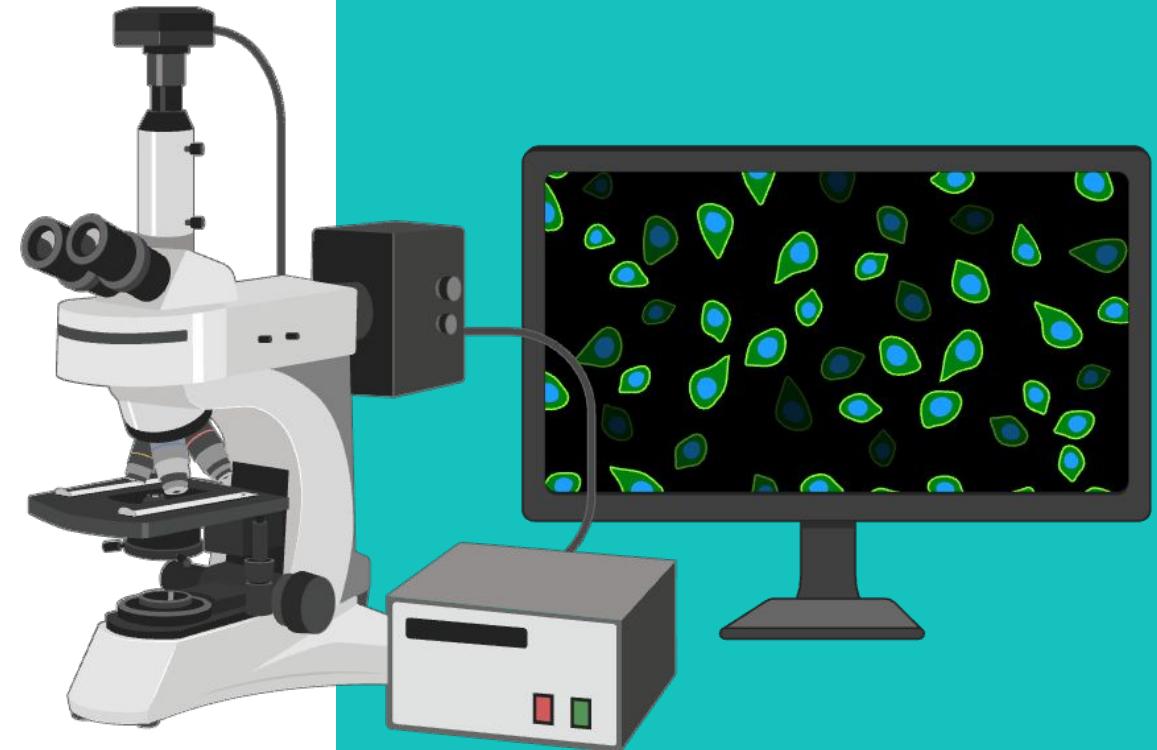
Hypothesis:

We hypothesize that both datasets will underrepresented the AFR superpopulation in their sampling

Research Methodology

We ran elementary statistics on both datasets

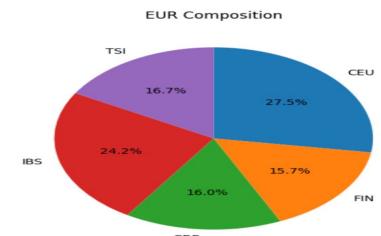
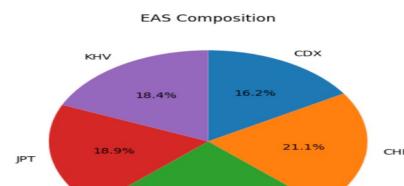
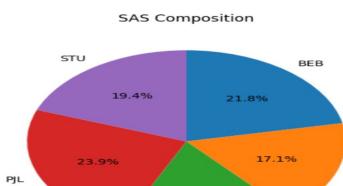
- Calculated population distribution
- Measured genomic diversity for available groups
- Compared representation gaps between datasets
- Interpreted patterns using demographic and evolutionary theory



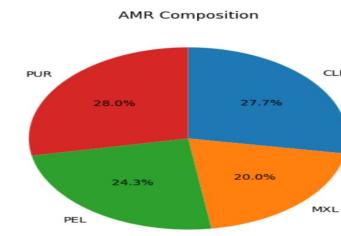
Use the [Resource section](#) to swap this figure

Comparison of Diversity Metrics in 1000s Genomes

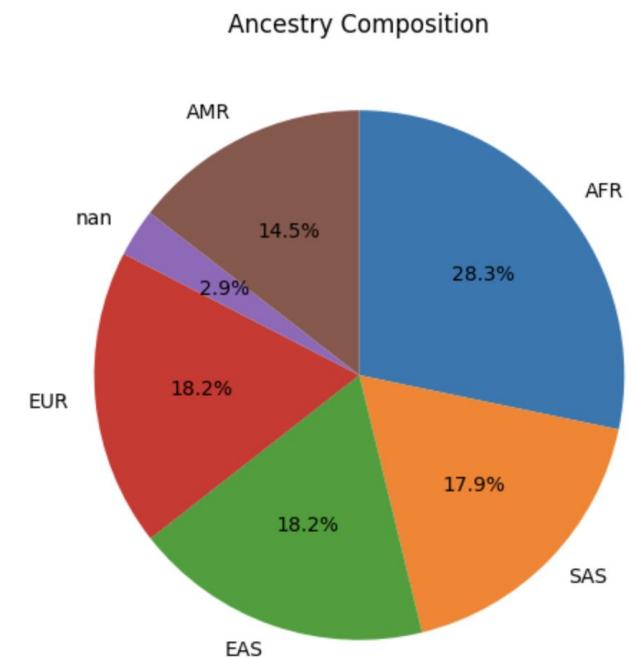
- Pretty equitable
- Overall even distribution
- Signifies more equal sampling and a better representation of data to be used



nan Composition



population	delta_%	shannon_diversity	kl	chi_p_vals	chi_stat	pi	fst	pair
super	NA	1.565213315	0.017038189	0.921226965	0.923076923	NA	NA	NA
sub	NA	3.276566952	NA	NA	NA	NA	NA	NA



Comparison of Diversity Metrics in 1000 Genomes and TCGA

- Computed same statistics for both datasets
- Compared them to find any significant results
- Brainstormed ways to improve possible inequities

```

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Date: 2025-10-28
Organization: Computational Biology @ Berkeley

This file automates/generalizes the computation of:
- Population counts and percentages by ancestry group and superpopulation
- Graphing the population data
- Shannon Diversity Index
...

import pandas as pd
import matplotlib.pyplot as plt
import math

population_col_title = ''
num_indiv_col_title = ''
percent_col_title = ''
superpopulation_col_title = ''
super_num_indiv_col_title = ''
super_percent_col_title = ''

path = ''

def set_path(in_path):
    path = in_path

def set_column_titles(in_population_col_title, in_num_indiv_col_title, in_percent_col_title, in_superpopulation_col_title, in_super_num_indiv_col_title, in_super_percent_col_title):
    ...
    Different data sets can have different names for the same concepts displayed in the columns.
    This function generalizes the columns (that we need) and allows the user to input what its called in their dataset.
    ...
    population_col_title = in_population_col_title
    num_indiv_col_title = in_num_indiv_col_title
    percent_col_title = in_percent_col_title
    superpopulation_col_title = in_superpopulation_col_title
    super_num_indiv_col_title = in_super_num_indiv_col_title
    super_percent_col_title = in_super_percent_col_title
    return population_col_title, num_indiv_col_title, percent_col_title, superpopulation_col_title, super_num_indiv_col_title, super_percent_col_title

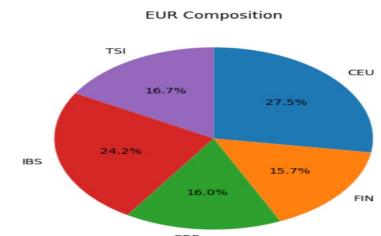
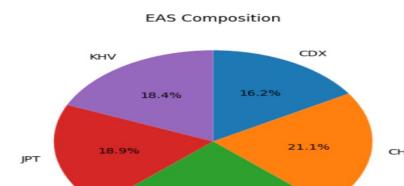
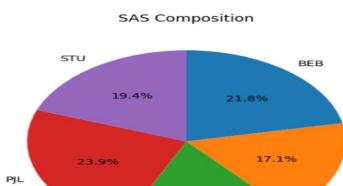
def read_data(file_path):
    """Reads data from a CSV file into a pandas DataFrame."""
    data = pd.read_csv(file_path)

```

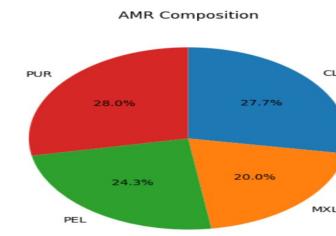
Example code

Diversity Metrics in 1000s Genomes

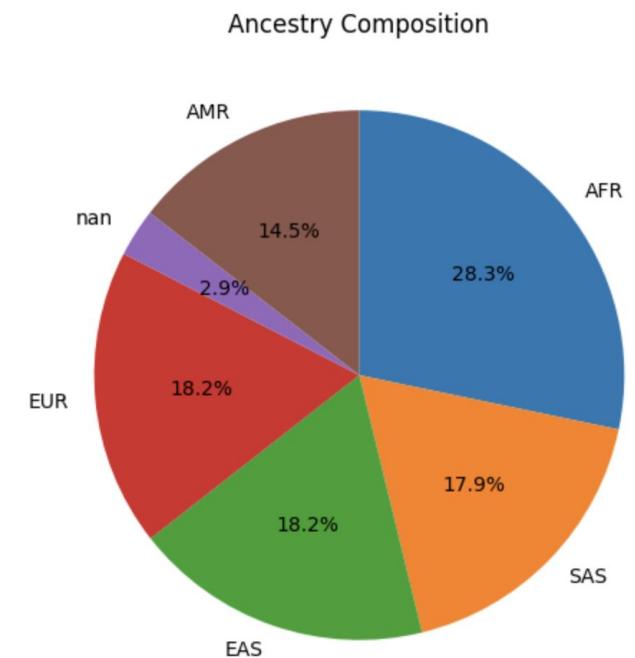
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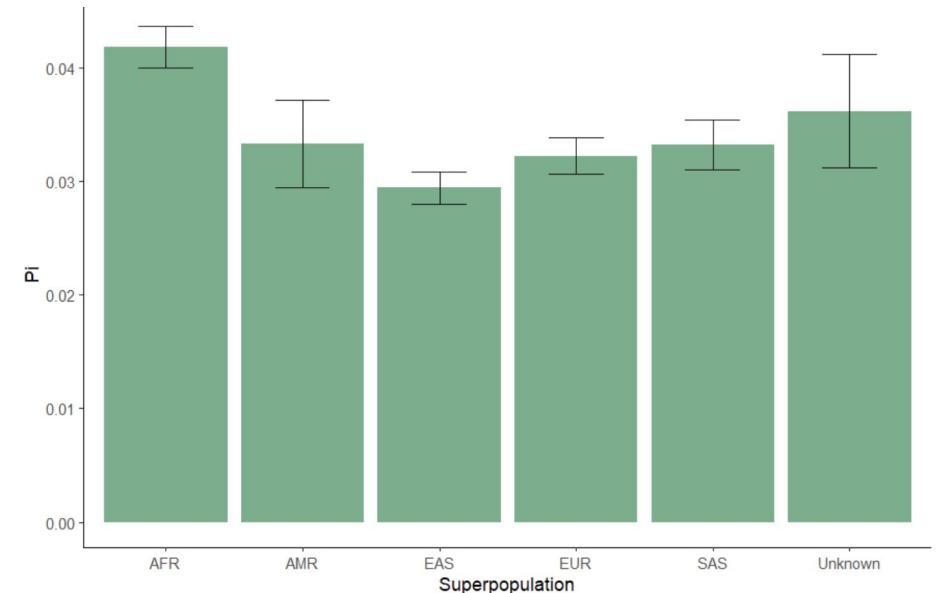


population	delta_%	shannon_diversity	kl	chi_p_vals	chi_stat	pi	fst	pair
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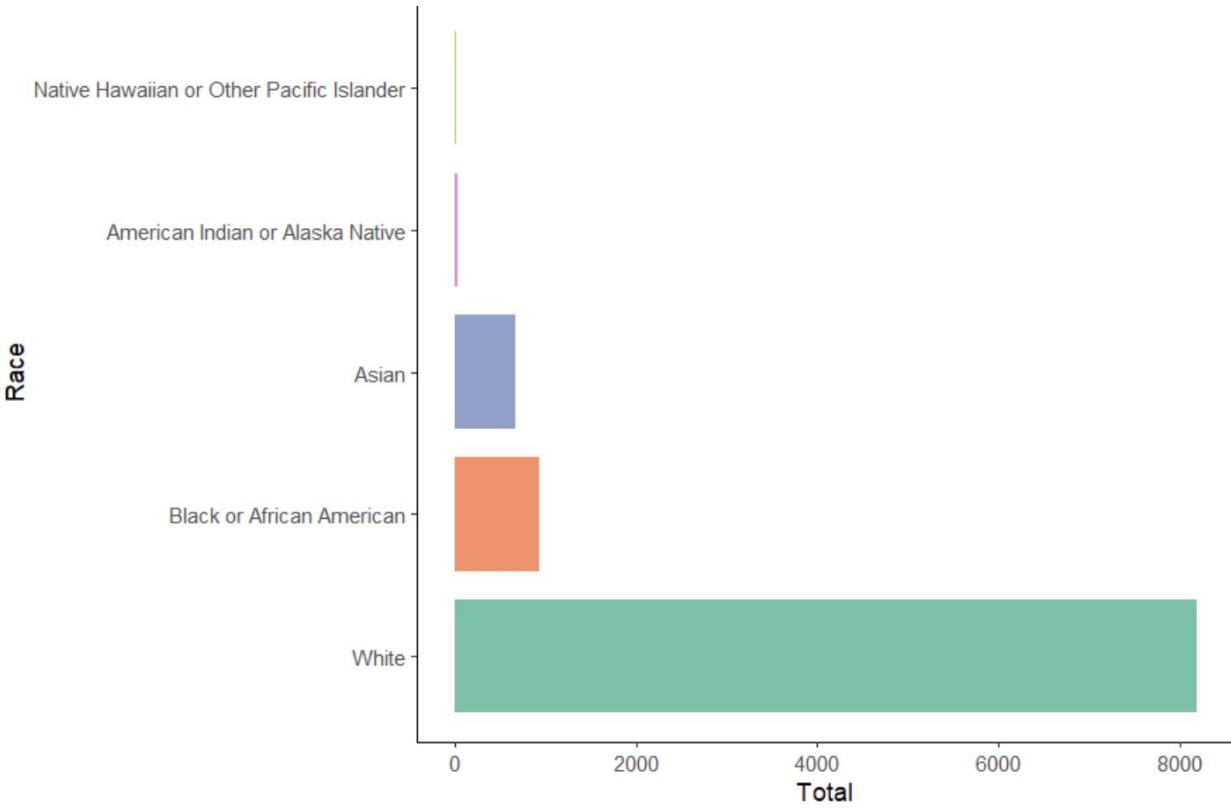
Diversity Metrics in 1000s Genomes

- Genomic diversity shows similar pattern
- Shows a realistic distribution, with AFR the highest
- AFR shows the highest π and $H\Delta$ because of larger long-term effective population size and weaker drift, allowing more mutations to arise and persist
 - Diversity declines in non-African groups due to bottlenecks during the Out-of-Africa expansion
- SAS > EUR \approx EAS because South Asians experienced less severe bottlenecks and more admixture, while Europeans and East Asians underwent additional founder events that reduced diversity to similar levels
- π (nucleotide diversity): the average number of pairwise nucleotide differences per site between two randomly chosen individuals in a population
- $H\Delta$ (expected heterozygosity): the probability that two alleles sampled from the same population are different (i.e., the fraction of sites expected to be heterozygous)



Diversity Metrics in TCGA

- Much more skewed
- More inequitable
- Signifies more biased sampling and a worse representation of data, similar to how the healthcare industry looks like
- No verified and published statistics for genomic diversity, but we assume it to be low due to TCGA's well documented sampling biases



Summary of Results and Drawbacks

- Two very different sides of the spectrum, showing clear disparities in which groups are represented in major genomic datasets
- Indicates that health inequity still exists, because underrepresented populations receive fewer research insights, weaker diagnostic tools, and less effective precision-medicine advances
- Highlights the need for more inclusive data collection and equitable distribution of genomic resources

1000 Genomes Project – Drawbacks

- Underrepresentation of many global populations, which limits downstream analyses of diversity, allele frequencies, and ancestry-specific variant interpretation
- Low sequencing depth (~4–6× WGS for many samples) reduces sensitivity for detecting rare variants and structural variants

TCGA – Drawbacks

- Limited demographic diversity and strong sampling bias, with overrepresentation of White patients and certain cancer types
- Highly heterogeneous data generation protocols (different platforms, labs, and time periods), which can introduce batch effects and complicate cross cohort comparisons

Remaining Questions

- How consistent are these representation gaps across other major biomedical and genomic datasets?
- How much do missing ancestry annotations distort our estimates of diversity and equity?
- Can we quantify how these disparities translate into downstream clinical or research biases?

Next Steps

- Run the same analyses on additional datasets to test whether the disparities we found generalize
- Build a GAN-based model that can automatically detect inequities, generate insights, and provide recommendations for improving dataset representation

THANKS FOR LISTENING