

Target genotyping using Capture-Seq technology to characterize the population structure and phylogenetic relationships of peaches from Australia and other parts of the world.

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Background

Germplasm Diversity

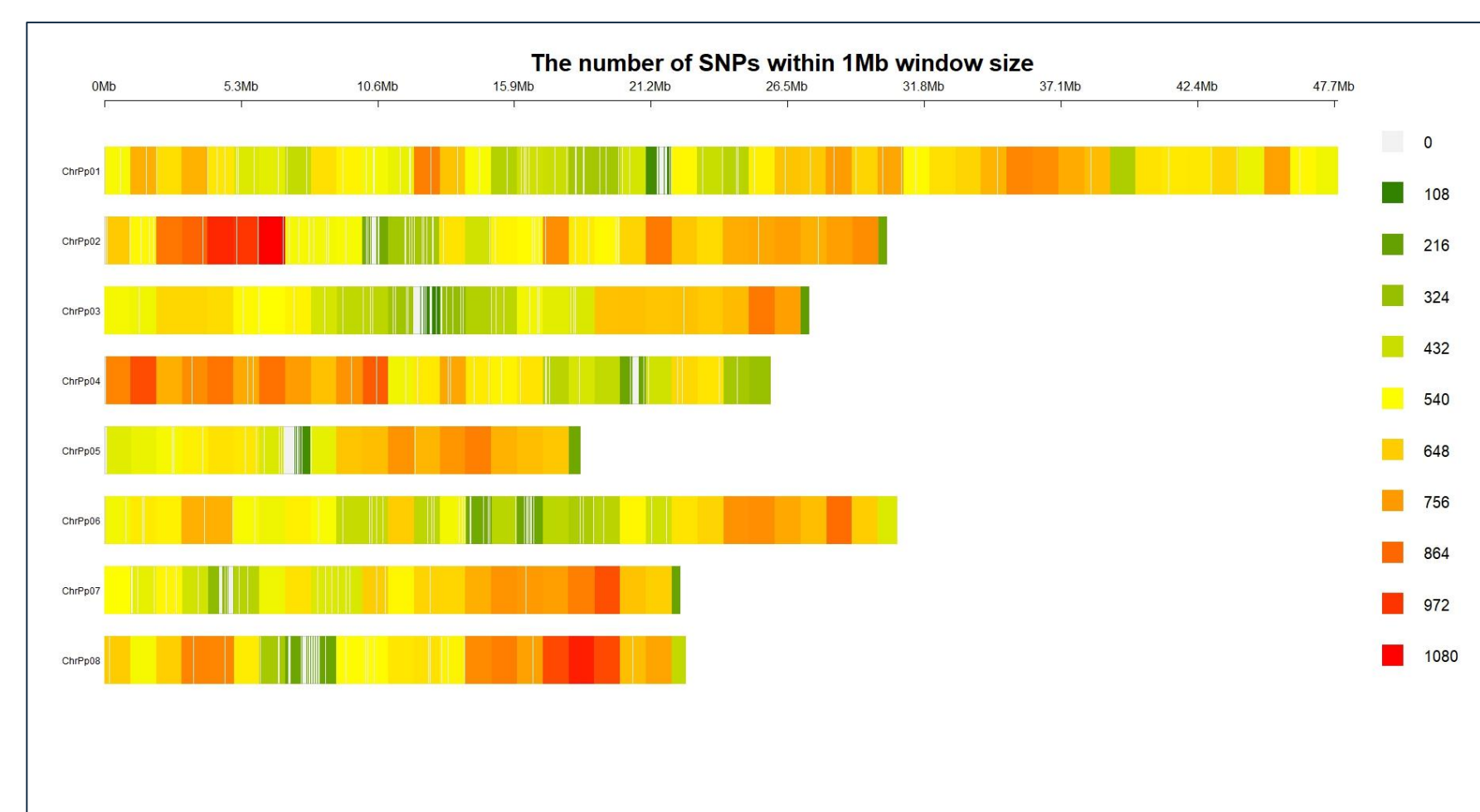
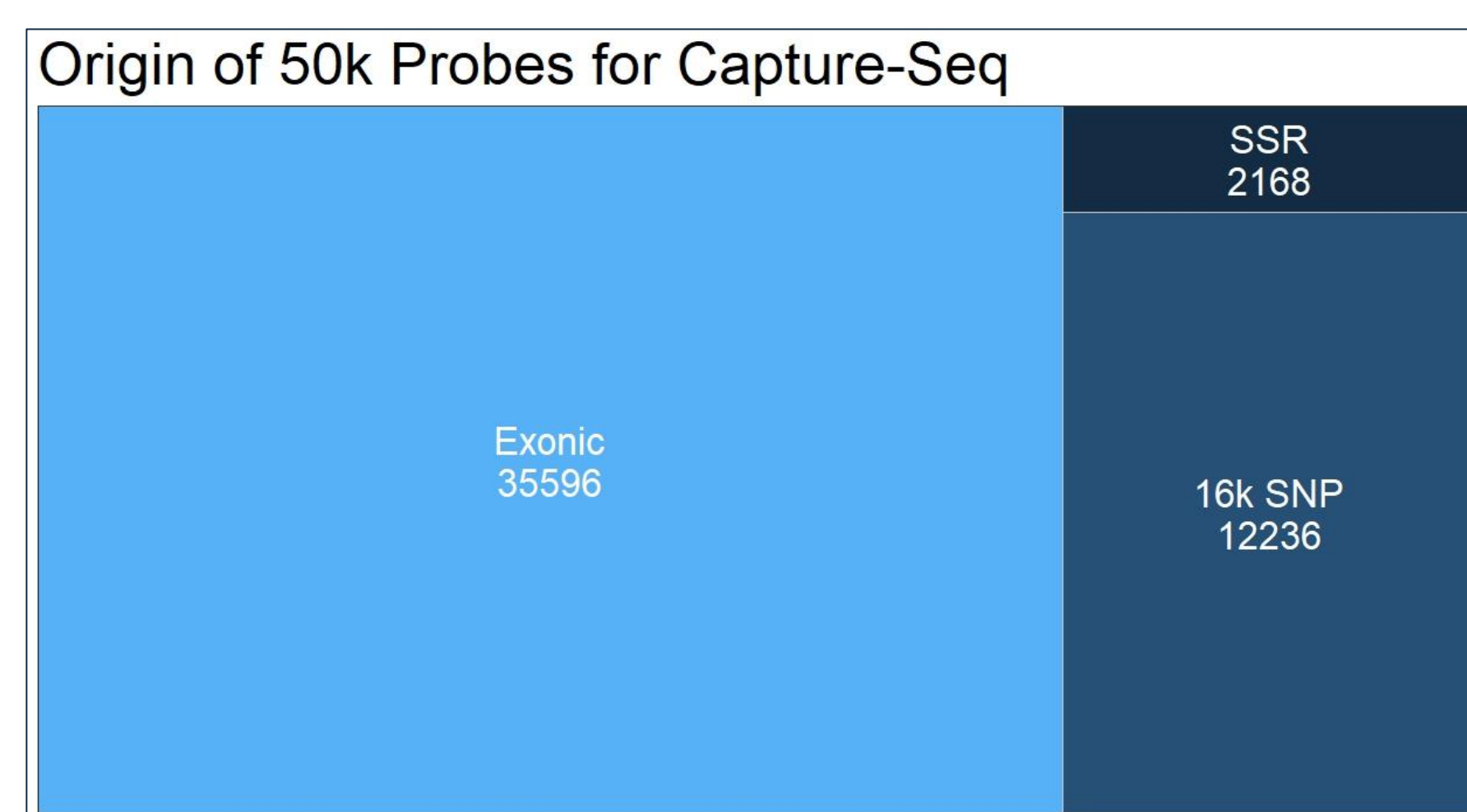
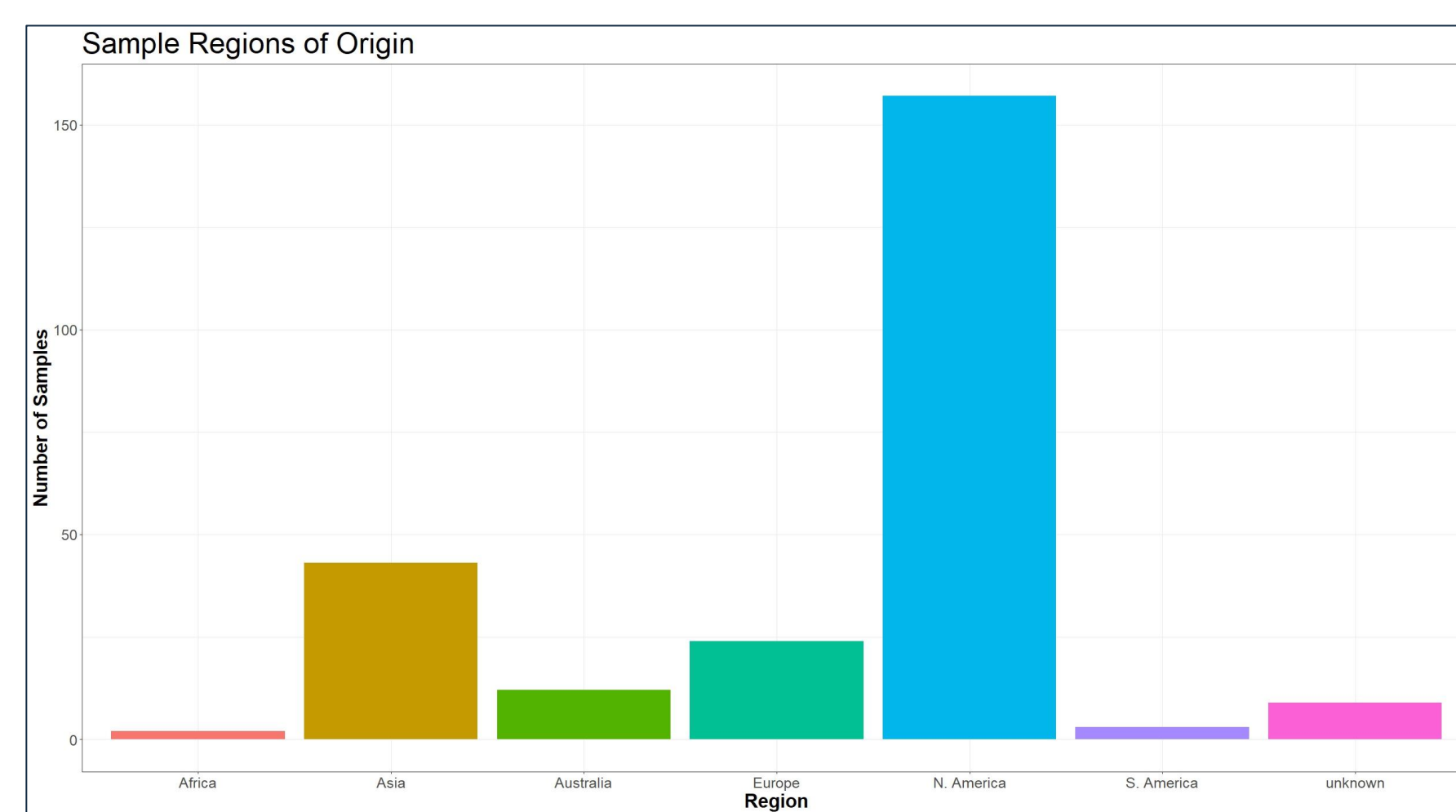
- Peaches in the US are highly homozygous
- A genetic bottleneck in the 1870s resulted in low diversity in US peaches
- Acquiring new diversity would bolster breeding efforts and help combat disease and climate related challenges
- Therefore, identifying potential sources of germplasm diversity is a priority

SNP Markers

- The current SNP chip array from Illumina is inflexible and expensive
- One must buy the whole array, even if you only need a few SNPs
- A probe panel would allow researchers to choose their targets and would expand on the SNP markers currently available
- Genotyping SNP-based methods such as KASP can be done in-house

Our Goals:

- Goal 1: Determine whether Australian peach populations are related to Chinese populations
- Goal 2: Develop regionally-informative SNP markers based on 50k probe panel for genotyping



Methodology

Sample Acquisition

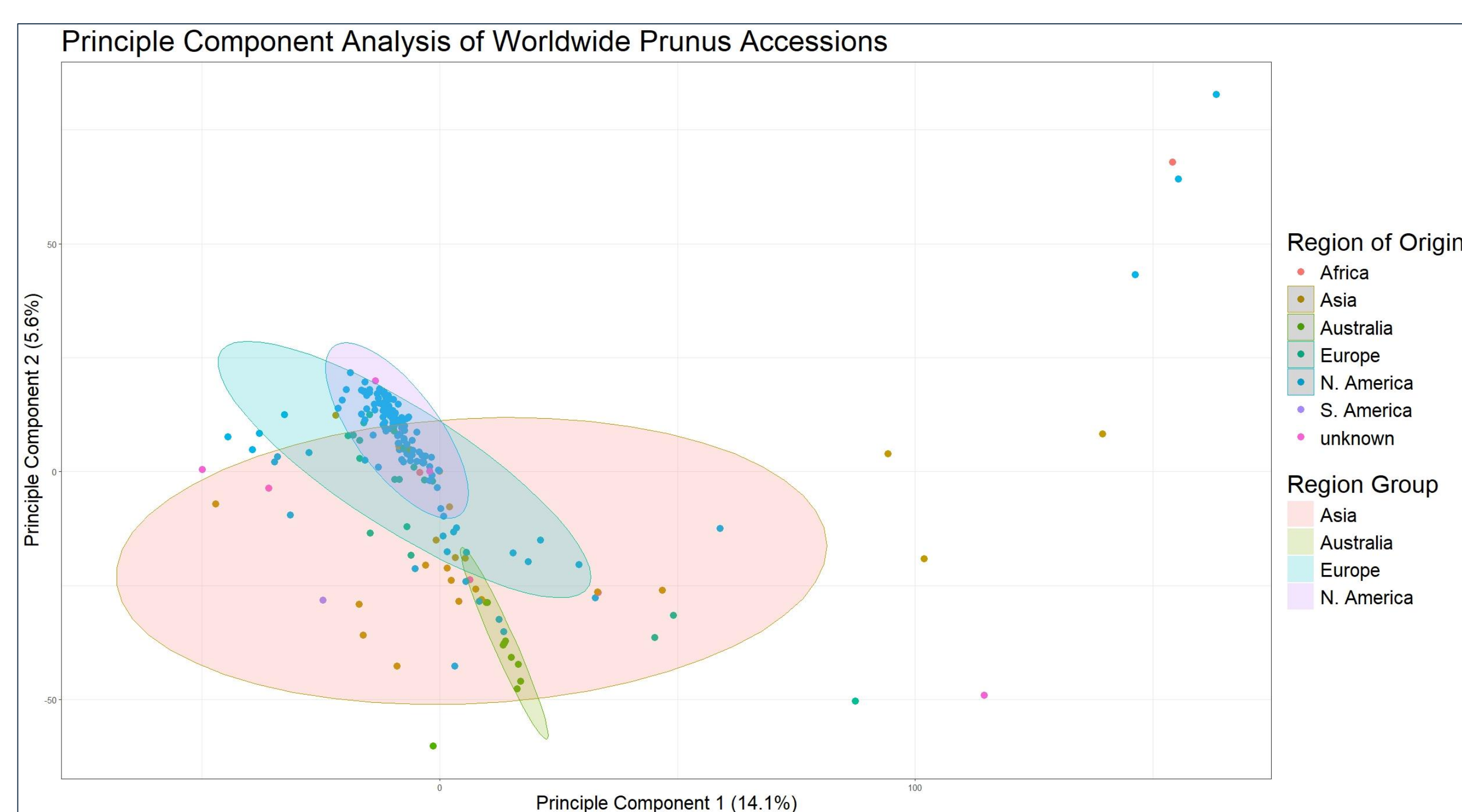
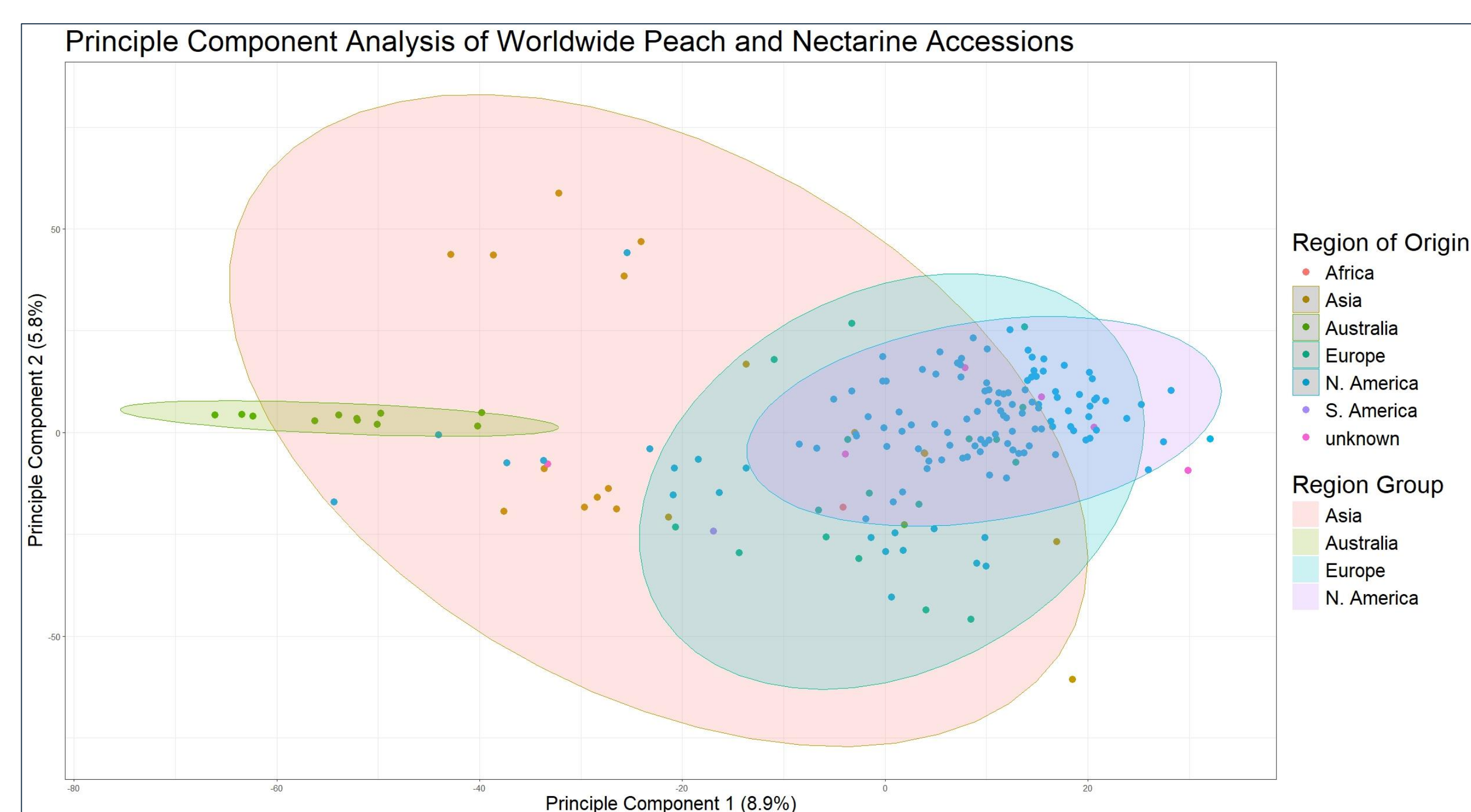
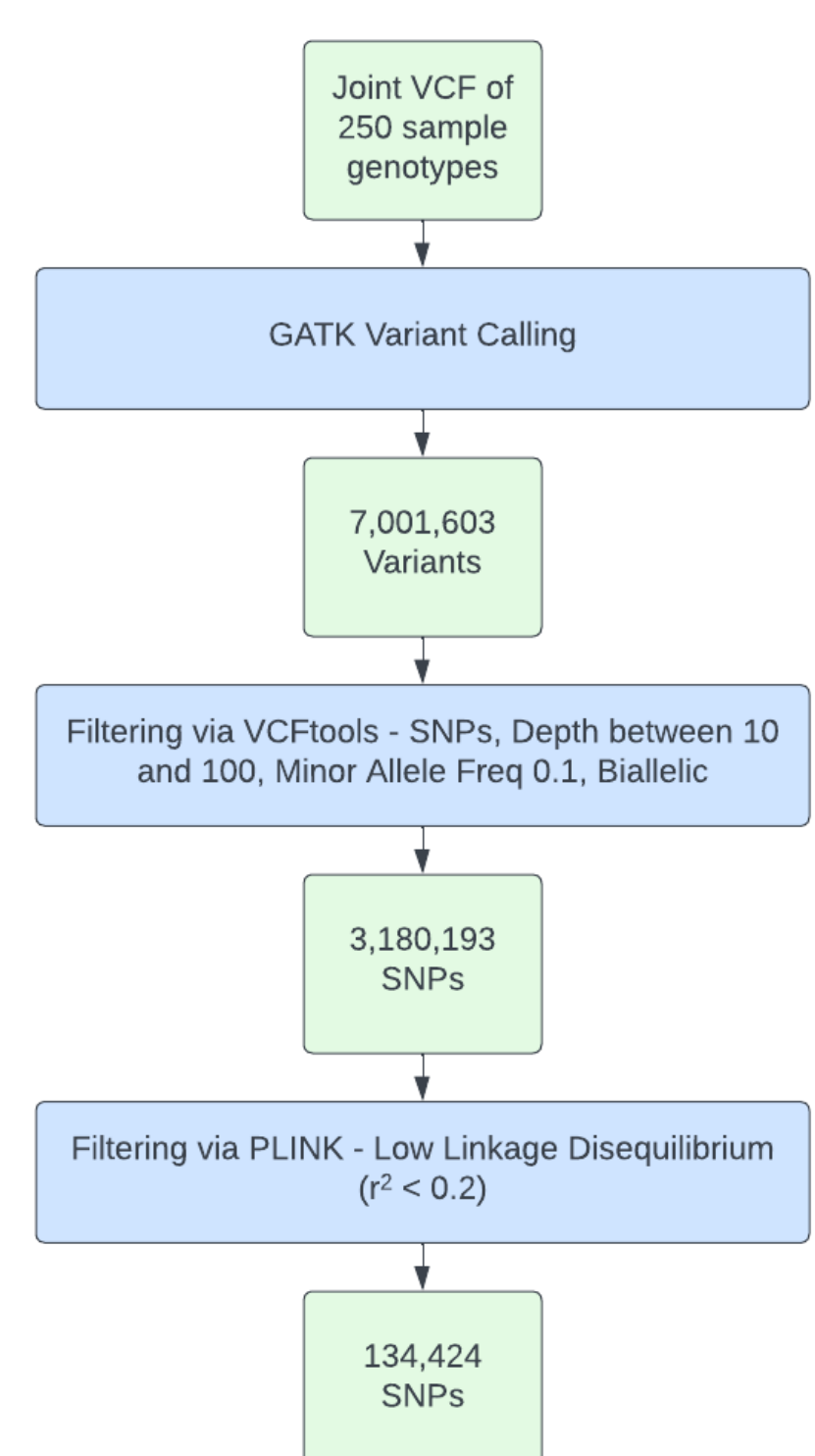
- Dempsey Farm, Griffin, GA
- USDA-ARS GRIN repository
- Australian Collaborators

Genotyping

- Capture-Seq through LGC BioSearch Technologies
- 50k probes developed based on the existing 16k SNP array, SSR markers, and exonic regions

Data Analysis

- 1) Evaluate data quality
 - FastQC and MultiQC
- 2) Align reads to genome and join all into one .vcf
 - VCFtools
- 3) Call variants (indels and SNPs)
 - GATK
- 4) Filter variants
 - SNPs, read depth between 10 and 100, MAF 0.1, low LD, biallelic
 - VCFtools, PLINK
- 5) Analyze results
 - PCA, Structure, heterozygosity, missing data

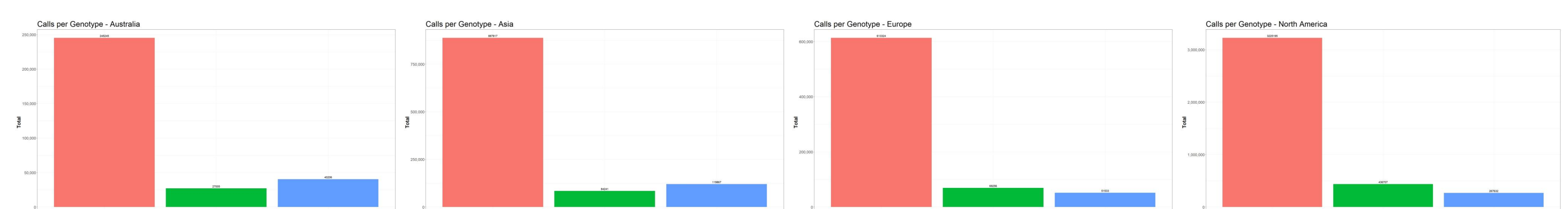
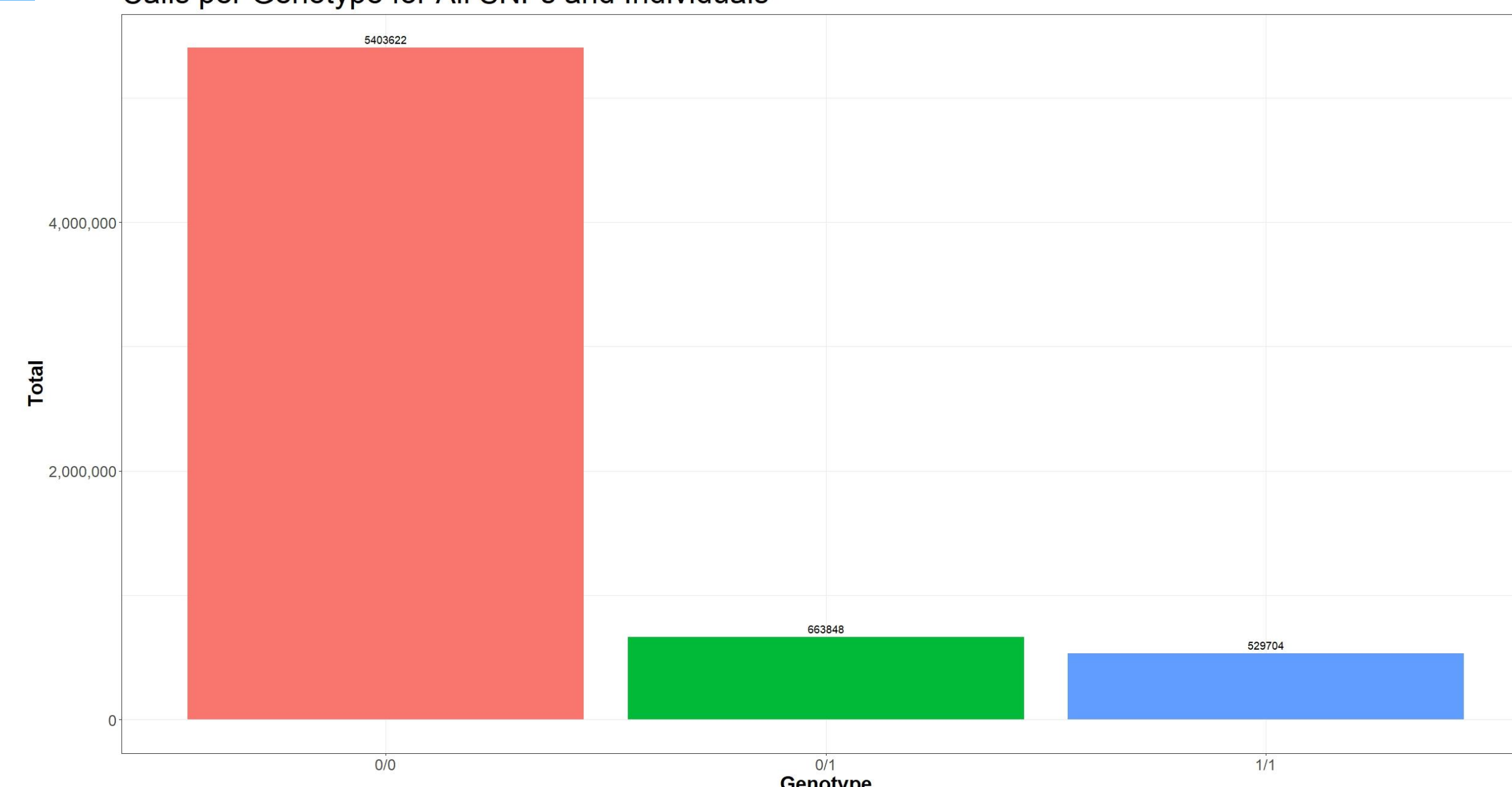


- 221 - *P. persica*
- 196 Peaches
- 25 Nectarines
- 29 - other species
- *P. augustifolia*
- *P. avium*
- *P. davidiana*
- *P. dulcis*
- *P. mira*
- *P. salicina*
- *P. domestica*
- *P. kansuensis*
- *P. mume*
- hybrids

Results

- 134,424 SNPs after filtering
- ~11k SNPs overlap with 16k SNP array
- All SSR-based probes yielded SNPs
- Clear separation of regional accessions
 - Next Step: isolate regionally informative SNP markers
 - Next Step: use these markers to further evaluate Australian germplasm
- 48 individuals with high percent missing data (>60%)
 - To be re-sequenced

Calls per Genotype for All SNPs and Individuals



Acknowledgements

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Chavez Lab



Personal Website

