# **Spit For Science Genetic Data**

# **Quality analysis by sample**

v0.6.0 30 October 2023

Table of genotype array quality statistics per sample. This is a tab-delimited text file.

## **Fields**

- 1. FID = family ID
- 2. IID = individual ID
- 3. ClinSex = self-reported sex from the clinical database
- 4. Xhet = proportion of heterozygous genotypes on the X chromosome
- 5. YcallRate = proportion of non-missing genotypes on the Y chromosome
- GeneticSex = estimated sex based on Xhet and YcallRate; 1=male, 2=female, NA=inconclusive
- 7. SexChr = inferred sex chromosome count
- 8. CallRate = proportion of non-missing genotypes across the genome
- 9. AutoHet = proportion of heterozygous genotypes on the autosomes
- 10. ClosestAncestry = approximate inferred ancestry
- 11. SRancestry = self-reported ancestry
- 12. SexMismatch = Is GeneticSex inconsistent with ClinSex? 1=yes, 0=no
- 13. AncestryMismatch = Is ClosestAncestry inconsistent with SRancestry? 1=yes, 0=no
- 14. QCfail = did the sample fail quality control for reasons other than SexProblems? 1=yes, 0=no
- 15. Reasons = if QCfail=1, reason(s) for failure

# **Details**

#### **FID**

Family ID as listed in the clinical database.

#### TTD

Individual ID ("TAG" ID). All letters have been converted to uppercase as necessary.

#### ClinSex

Self-reported biological sex from the clinical database. 1=male, 2=female, 3=other, 4=prefer not to answer, 999=unknown/don't know.

#### Xhet

Proportion of heterozygous genotypes on the X chromosome. Calculated using bcftools 1.9.

#### **YcallRate**

Proportion of non-missing genotypes on the Y chromosome. Calculated using PLINK 1.9b6.21.

#### GeneticSex

Inferred genetic sex based on Xhet and YcallRate. Inferred to be male (GeneticSex=1) if Xhet < 0.05 and YcallRate > 0.75. Inferred to be female (GeneticSex=2) if 0.1 < Xhet < 0.3 and YcallRate < 0.4. Otherwise, inconclusive (GeneticSex=NA).

#### SexChr

Inferred sex chromosome count:

- XY if Xhet < 0.05 and YcallRate > 0.75
- XX if 0.1 < Xhet < 0.3 and YcallRate < 0.4</li>
- XXY if 0.1 < Xhet < 0.3 and YcallRate > 0.75
- unknown otherwise.

#### CallRate

Proportion of non-missing genotypes across the genome, after excluding

- SNPs with call rate <97%
- X chromosome SNPs with call rate <97% in males after setting heterozygous calls to missing.

Calculated using PLINK 1.9b6.21. Individuals were considered to have failed the CallRate step if CallRate < 97%.

#### AutoHet

Proportion of heterozygous genotypes on the autosomes. Calculated using bcftools 1.9. Individuals were considered to have failed the AutoHet step if AutoHet > 6 times the interquartile range (IQR) based on all samples.

### ClosestAncestry

Approximate ancestry based on principal components analysis with 1000Genomes data as a reference. Performed using PLINK 1.9b6.21. Individuals were assigned to have the same continental ancestry as the 1000 Genomes sample "closest" to them, where "closest" was defined as Euclidean distance based on the first 3 principal components. Distances were calculated using R version 3.6.1. AFR = African, EAS = East Asian, EUR = European, SAS = South Asian.

#### **SRancestry**

Self-reported ancestry from the clinical database. Individuals were selected if all 4 grandparents were reported to come from the same continental group. EAS = East Asian, EUR = European, SAS = South Asian, NA = not available.

#### SexMismatch

Was ClinSex inconsistent with GeneticSex? 1=yes, 0=no. Inconsistent if

- GeneticSex is inconclusive
- ClinSex not missing and ClinSex not the same as GeneticSex.

#### **AncestryMismatch**

Was ClosestAncestry inconsistent with SRancestry? 1=yes, 0=no. Inconsistent if SRancestry is not missing and ClosestAncestry is not the same as SRancestry.

# **QCfail**

Did the individual fail quality control for non-sex reasons? 1=yes, 0=no. Failed if

- CallRate <97% (IMISS)
- AutoHet >6 IQR (HET).

### Reasons

If QCfail = 1, reason(s) for failure. NA if QCfail = 0.

# Resources

1000 Genomes data: https://tcag.ca/tools/1000genomes.html (independent samples bundle)

bcftools: https://samtools.github.io/bcftools/bcftools.html

PLINK: <a href="https://www.cog-genomics.org/plink/">https://www.cog-genomics.org/plink/</a>

R: <a href="https://cran.r-project.org/">https://cran.r-project.org/</a>