

# Module 18 Multivariate Analysis for Genetic data

## Session 05: Log-ratio principal component analysis

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# What is compositional data?

- Compositional data consists of variables that are **parts of some whole**.
- Typical examples are proportions, percentages, concentrations.
- The data vectors are constrained and reside in a simplex.
- Compositions provide information about the **relative** values of the parts.
- Aitchison (1986) proposed the **log-ratio approach** to deal with compositional data.

# Compositional data

Compositional data arise in many contexts:

- Mineral composition of rocks (geology)
- Time or budget expenditure (economy)
- Bacterial composition of the gut (microbiology)
- Allele frequencies and genotype frequencies (genetics)
- ...

# Compositional data and spurious correlations

counts

	A	B	C	D	E
s1	10	30	20	50	40
s2	0	30	20	10	90
s3	20	80	10	30	10

full composition

	A	B	C	D	E
s1	0.07	0.20	0.13	0.33	0.27
s2	0.00	0.20	0.13	0.07	0.60
s3	0.13	0.53	0.07	0.20	0.07

subcomposition

	A	B	C	D
s1	0.09	0.27	0.18	0.45
s2	0.00	0.50	0.33	0.17
s3	0.14	0.57	0.07	0.21

R

	A	B	C	D	E
A	1.00	0.87	-0.87	0.50	-0.99
B	0.87	1.00	-1.00	0.00	-0.79
C	-0.87	-1.00	1.00	-0.00	0.79
D	0.50	0.00	-0.00	1.00	-0.62
E	-0.99	-0.79	0.79	-0.62	1.00

R

	A	B	C	D
A	1.00	0.07	-1.00	0.31
B	0.07	1.00	-0.14	-0.93
C	-1.00	-0.14	1.00	-0.24
D	0.31	-0.93	-0.24	1.00

- Correlations can be **spurious** due to the existence of a linear constraint
- Ordinary PCA of the data will display a spurious correlation structure

# Principles of Compositional Data Analysis (CoDA)

## Principles:

- Scale invariance
- Permutation invariance
- Subcompositional coherence

## Typical CoDA approach:

- In order to satisfy these principles, we use a [log-ratio transformation](#) of the data.
- Analyse the data by applying the classical statistical methods to the log-ratio transformed data.

## Some notation

A composition of  $D$  parts

$$\mathbf{x} = (x_1, x_2, \dots, x_D)$$

The sample space is the **simplex**

$$S^D = \left\{ \mathbf{x} = (x_1, x_2, \dots, x_D) \mid x_i > 0, i = 1, 2, \dots, D; \sum_{i=1}^D x_i = \kappa \right\}$$

The **closure** operation  $\mathcal{C}$  to the constant  $\kappa > 0$  (usually 1)

$$\mathcal{C}(\mathbf{x}) = \left( \frac{\kappa x_1}{\sum_{i=1}^D x_i}, \frac{\kappa x_2}{\sum_{i=1}^D x_i}, \dots, \frac{\kappa x_D}{\sum_{i=1}^D x_i} \right)$$

# Log-ratio transformations

- Additive log-ratio transformation (alr; ratios of two parts)

$$alr(\mathbf{x}) = \left[ \ln \left( \frac{x_1}{x_D} \right), \ln \left( \frac{x_2}{x_D} \right), \dots, \ln \left( \frac{x_{D-1}}{x_D} \right) \right],$$

- Centred log-ratio transformation (clr; ratios of one part against all)

$$clr(\mathbf{x}) = \left[ \ln \left( \frac{x_1}{g_m(\mathbf{x})} \right), \ln \left( \frac{x_2}{g_m(\mathbf{x})} \right), \dots, \ln \left( \frac{x_D}{g_m(\mathbf{x})} \right) \right],$$

- Isometric log-ratio transformation (ilr; ratios of geometric means of subcompositions)

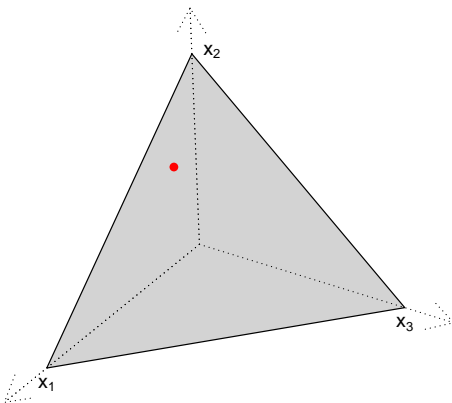
$$ilr(\mathbf{x}) = \left[ \ln \left( \frac{g_m(\mathbf{x}_a)}{g_m(\mathbf{x}_b)} \right), \dots, \ln \left( \frac{g_m(\mathbf{x}_c)}{g_m(\mathbf{x}_d)} \right) \right],$$

- $g_m(\mathbf{x})$  is the **geometric mean** of the components of the composition  $\mathbf{x}$

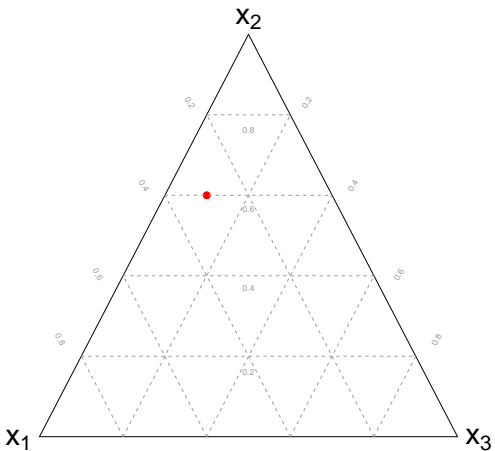
$$g_m(\mathbf{x}) = \left( \prod_{i=1}^D x_i \right)^{1/D} \quad \ln(g_m(\mathbf{x})) = \frac{1}{D} \sum_{i=1}^D \ln(x_i) \quad g_m(\mathbf{x}) = e^{\bar{y}} \quad y_i = \ln(x_i)$$



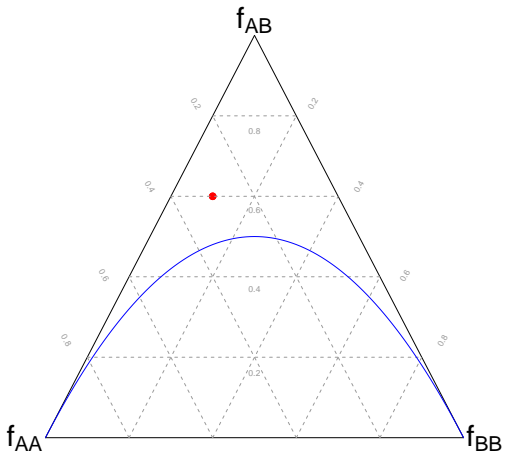
# Visualizing 3 part compositions: ternary diagram



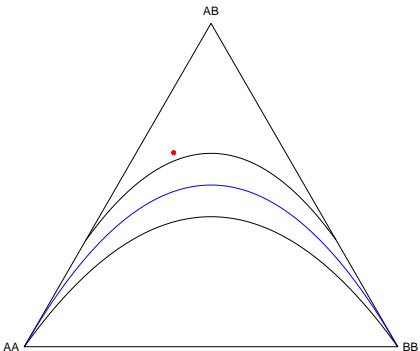
# How to read a ternary diagram?



# The ternary diagram in genetics: De Finetti diagram

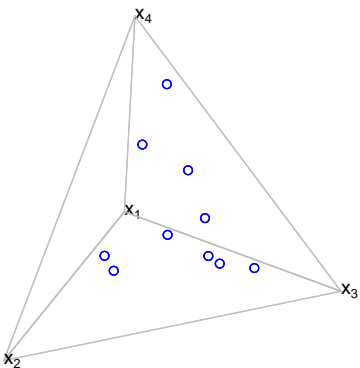


# Is there Hardy-Weinberg equilibrium?



Graffelman, J. & Morales-Camarena, J. (2008) Graphical tests for Hardy-Weinberg equilibrium based on the ternary plot. *Human Heredity* 65(2): 77-84

# Plotting four-part compositions



# Plotting compositional data

- Three part compositions can be visualised in a [ternary diagram](#)
- Four part compositions can be visualised in a [tetrahedron](#)
- Larger compositions can be visualised, in an approximate manner, in a [compositional biplot](#)
- Even three- and four part compositions are often better shown in a compositional biplot, as this represents them in an [unconstrained space](#).
- Compositional biplots are obtained by [log-ratio principal component analysis](#) (LR-PCA).

## Log-ratio PCA based on the centered log-ratio transformation

$$clr(x_i) = \ln \left( \frac{x_i}{g_m(\mathbf{x})} \right) = \ln \left( \frac{x_i}{(\prod x_j)^{1/D}} \right) = \ln(x_i) - \frac{1}{D} \sum_{i=1}^D \ln(x_i)$$

In matrix form:

$$\mathbf{X}_l = \ln(\mathbf{X})$$

$$\mathbf{X}_{clr} = \mathbf{X}_l \mathbf{H}_r$$

$$\mathbf{X}_{cclr} = \mathbf{H}_c \mathbf{X}_{clr} = \mathbf{H}_c \mathbf{X}_l \mathbf{H}_r$$

where

$$\mathbf{H}_r = \mathbf{I} - \frac{1}{D} \mathbf{1}\mathbf{1}' \quad \mathbf{H}_c = \mathbf{I} - (1/n)\mathbf{1}\mathbf{1}'.$$

The log-transformed data is **double-centered**

Do a standard PCA of the  $\mathbf{X} = \mathbf{X}_{cclr}$

# Singular value decomposition (SVD)

Log-ratio PCA can be performed by the SVD:

$$\mathbf{X}_{\text{clr}} = \mathbf{UDV}' \text{ with } \mathbf{U}'\mathbf{U} = \mathbf{I} \text{ and } \mathbf{V}'\mathbf{V} = \mathbf{I}.$$

Possible biplot coordinates (row markers  $\mathbf{F}$  and column markers  $\mathbf{G}$ )

- $\mathbf{F} = \mathbf{UD}$  and  $\mathbf{G} = \mathbf{V}$  (the [form biplot](#))
- $\mathbf{F} = \mathbf{U}$  and  $\mathbf{G} = \mathbf{VD}$  (the [covariance biplot](#))
- $\mathbf{F} = \mathbf{UD}^{1/2}$  and  $\mathbf{G} = \mathbf{VD}^{1/2}$  (the [symmetric biplot](#))
- The [form biplot](#) will approximate the [Aitchison distances](#) between the compositions.
- The Aitchison distance is the Euclidean distance between the clr transformed compositions.



# The zero problem

The zero issue:

- Compositional data analysis generally considers the [simplex to be open](#)
- Zeros are not admitted

Important questions:

- How many zeros do you have?
- What kind of zeros do you have?
  - [Rounding zeros](#) (below detection limit)
  - [Count zeros](#) (related to sampling effort)
  - [Essential or structural zeros](#) (impossible outcome)

Solutions:

- For rounding or count zeros, [impute](#) a reasonable non-zero amount, for structural zeros, [stratify](#).

# Compositional biplot interpretation

- The origin represents the geometric mean of the compositions.
- Biplot vectors represent **clr transformed parts**.
- **Links** between vectors represent **pairwise log-ratios**:

$$\text{clr}(x_1) - \text{clr}(x_2) = \ln \left( \frac{x_1}{g_m(\mathbf{x})} \right) - \ln \left( \frac{x_2}{g_m(\mathbf{x})} \right) = \ln \left( \frac{x_1}{x_2} \right)$$

- The **link length** represents the **standard deviation of the corresponding log-ratio**.

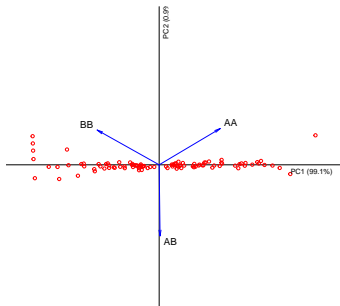
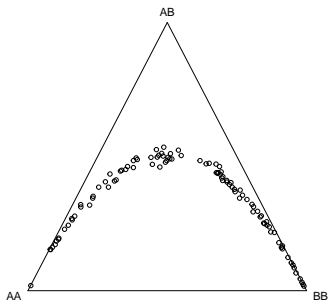
$$\begin{aligned} \|\mathbf{f}_i - \mathbf{f}_j\|^2 &= \mathbf{f}_i' \mathbf{f}_i + \mathbf{f}_j' \mathbf{f}_j - 2\mathbf{f}_i' \mathbf{f}_j \\ &= \text{Var}(\text{clr}(x_i)) + \text{Var}(\text{clr}(x_j)) - 2\text{Cov}(\text{clr}(x_i), \text{clr}(x_j)) \\ &= \text{Var} \left( \ln \left( \frac{x_i}{g_m(\mathbf{x})} \right) - \ln \left( \frac{x_j}{g_m(\mathbf{x})} \right) \right) = \text{Var} \left( \ln \left( \frac{x_i}{x_j} \right) \right). \end{aligned}$$

- Close to coincident biplot vectors suggest **proportionality of parts**.
- Cosines of angles between links represent **correlations between log-ratios**
- Collinear biplot vectors suggest a **one-dimensional pattern for a subcomposition**.

# Experiment

- Take repeated samples from bi-allelic genetic variants that are in Hardy-Weinberg equilibrium
- Record the genotype composition
- Do a log-ratio PCA of the compositions obtained

# Results



- PC1: log odds of the allele frequency
- PC2: deviation from Hardy-Weinberg equilibrium

	PC1	PC2	PC3	[,1]	[,2]	[,3]
1a	4.837	0.04373	7.28e-32	AA 0.7019	0.417	0.577
1af	0.991	0.00896	1.49e-32	BB -0.7121	0.399	0.577
1ac	0.991	1.00000	1.00e+00	AB 0.0102	-0.816	0.577

$$\ln(f_{AA}) + \ln(f_{BB}) - 2 \ln(f_{AB}) = \ln\left(\frac{f_{AA}f_{BB}}{f_{AB}^2}\right) = \ln\left(\frac{p^2q^2}{(2pq)^2}\right) = c$$

# Worldwide Y-STR dataset

- Purps, J. et al. (2014) A global analysis of Y-chromosomal haplotype diversity for 23 STR loci. *Forensic Science International: Genetics* 12: 12–23.
- Data consists of 23 Y-STRs typed for 19,630 males in 129 populations sampled world-wide,
- Samples stemming from Africa, Asia, Europe, Latin and North America.

## Example: Allele frequencies of Y-STR DYS448 over 129 populations worldwide

Sample	19	20	21	other
Arg-B	37	31	10	14
Arg-F	30	22	13	6
Arg-M	45	34	12	10
Arg-N	23	14	3	10
Arg-S	25	10	6	9
Aus	108	100	24	27
Bel-A	115	61	12	18
Bel-V	63	32	3	7
Ben	4	7	32	8
Bol-M	19	18	4	3
Bol-N	16	33	5	2
Bos	44	46	6	4
Bra-R	53	28	24	18
Bra-SG	35	12	10	4
Bra-SP	54	33	24	9
Chi-B	91	80	27	48
Chi-C	44	19	6	31
Chi-Sh	35	49	6	19
Chi-So	17	4	1	8
Chi-Xi	6	53	4	29
Chi-Xu	53	37	4	51
Chi-Y	31	39	3	28
CoR	75	42	27	22
Cro-C	54	57	8	6
Cro-Z	40	63	10	1
Cze-B	25	45	1	1
Cze-M	13	22	5	2
Den	76	92	10	7
ENG-C	46	24	3	8
ENG-S	66	30	6	12
.	.	.	.	.
.	.	.	.	.
.	.	.	.	.
Wal	85	17	3	13
Zim	3	6	42	4

Sample	19	20	21	other
Arg-B	0.40	0.34	0.11	0.15
Arg-F	0.42	0.31	0.18	0.08
Arg-M	0.45	0.34	0.12	0.10
Arg-N	0.46	0.28	0.06	0.20
Arg-S	0.50	0.20	0.12	0.18
Aus	0.42	0.39	0.09	0.10
Bel-A	0.56	0.30	0.06	0.09
Bel-V	0.60	0.30	0.03	0.07
Ben	0.08	0.14	0.63	0.16
Bol-M	0.43	0.41	0.09	0.07
Bol-N	0.29	0.59	0.09	0.04
Bos	0.44	0.46	0.06	0.04
Bra-R	0.43	0.23	0.20	0.15
Bra-SG	0.57	0.20	0.16	0.07
Bra-SP	0.45	0.28	0.20	0.07
Chi-B	0.37	0.33	0.11	0.20
Chi-C	0.44	0.19	0.06	0.31
Chi-Sh	0.32	0.45	0.06	0.17
Chi-So	0.57	0.13	0.03	0.27
Chi-Xi	0.07	0.58	0.04	0.32
Chi-Xu	0.37	0.26	0.03	0.35
Chi-Y	0.31	0.39	0.03	0.28
CoR	0.45	0.25	0.16	0.13
Cro-C	0.43	0.46	0.06	0.05
Cro-Z	0.35	0.55	0.09	0.01
Cze-B	0.35	0.62	0.01	0.01
Cze-M	0.31	0.52	0.12	0.05
Den	0.41	0.50	0.05	0.04
ENG-C	0.57	0.30	0.04	0.10
ENG-S	0.58	0.26	0.05	0.11
.	.	.	.	.
.	.	.	.	.
.	.	.	.	.
Wal	0.72	0.14	0.03	0.11
Zim	0.05	0.11	0.76	0.07

Purps, J. et al. (2014) A global analysis of Y-chromosomal haplotype diversity for 23 STR loci. Forensic Science International: Genetics 12: 12–23.



## References:

- Aitchison, J. (1986) *The statistical analysis of compositional data*. Chapman & Hall.
- Aitchison, J. (1983) Principal component analysis of compositional data. *Biometrika* 70(1) pp. 57-65.
- Pawlowsky-Glahn, V., Egozcue, J.J. & Tolosana-Delgado, R. (2015) *Modeling and Analysis of Compositional Data*, Chichester, United Kingdom, John Wiley & Sons.