

GHEP kinship exercise 2024: Advanced level

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General instructions

This is a multiple-choice test consisting of 20 questions. For each question exactly one alternative is correct. You are free to use whatever software you like, but keep in mind that some programs have built-in conventions (e.g., rounding) that may affect the output. If your answer does not precisely match any of the options, choose the closest one.

Files needed to complete the test

- *cousins-data.txt* / *cousins-ibd.txt*. Data for Part II.
- *siblings-data.txt* / *siblings-ibd.txt*. Data for Part III.
- *db.txt*. Allele frequencies for 23 STR markers. (If your software requires database size, use N = 1000.)

Assumptions throughout

- No linkage between markers, no linkage disequilibrium, no deviations from HW equilibrium.
- No drop-outs, drop-ins, silent alleles or mutations.
- Pedigree founders are non-inbred and unrelated to each other.
- The total genetic length of the autosome (chromosomes 1–22) is 3391 cM.

Some definitions

Homologous alleles are *identical by descent* (IBD) if they have the same origin within a given pedigree. The *IBD coefficients* ($\kappa_0, \kappa_1, \kappa_2$) of non-inbred individuals A and B, are the probabilities of sharing respectively 0, 1 and 2 alleles IBD at a random autosomal locus. They are related to the kinship coefficient φ by the formula $\varphi = \kappa_1/4 + \kappa_2/2$.

The *IBD triangle* (Figure 1) is a convenient tool for visualising IBD coefficients. Note that, since $\kappa_0 + \kappa_1 + \kappa_2 = 1$, any two of them suffice to deduce the third; the choice of κ_0 and κ_2 is simply my personal preference. The online tool [QuickPed](#) may be useful for calculating IBD coefficients and plotting them in the IBD triangle.

Traditional coefficients like κ and φ measure the **expected** IBD sharing based on the pedigree. In contrast, the *realised* (or *genomic*) relatedness between A and B refers to the **actual** IBD segments they share as a result of recombination (Figure 2). We denote by (k_0, k_1, k_2) the actual proportions of the autosome, in terms of genetic length, where they share 0, 1 and 2 alleles IBD, respectively. The *realised kinship coefficient* is given by $\varphi_R = k_1/4 + k_2/2$.

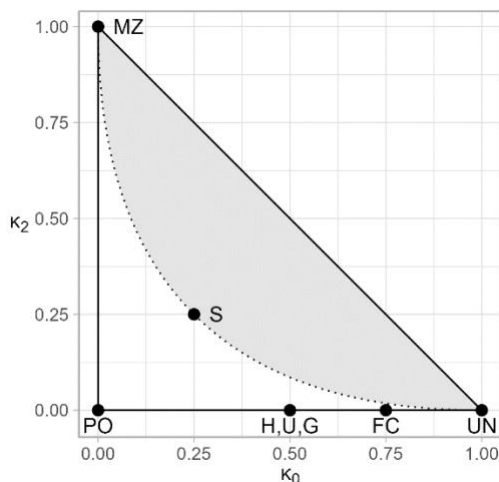


Figure 1. The IBD triangle. FC=first cousins; G=grandparent-grandchild; H=half sibs; MZ=monozygous twins; PO=parent-offspring; S=full sibs; U=uncle-nephew; UN=unrelated

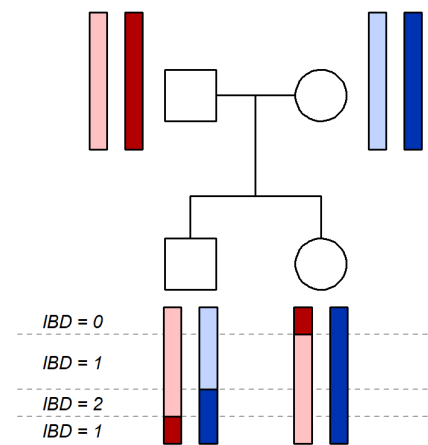


Figure 2. An example of the realised IBD sharing between siblings. The chromosome is divided in segments with IBD status 0, 1 or 2

Questions

Part I: Warm-up

We consider a situation where two individuals, A and B, are typed with a tri-allelic marker. The alleles are labelled 1,2,3, and the allele frequencies are p_1, p_2, p_3 , respectively.

1. Suppose the marker lies in a region where A and B have IBD status 0. If A has genotype 1/1, the genotype of B is
 - a) 2/3
 - b) 2/2 or 3/3
 - c) 2/2, 2/3 or 3/3
 - d) anything except 1/1
 - e) anything
2. Suppose the marker lies in a region with IBD = 1. If A has genotype 1/2, the genotype of B is
 - a) 1/3 or 2/3
 - b) 1/1, 2/2, 1/3 or 2/3
 - c) anything except 1/2
 - d) anything except 3/3
 - e) anything

In the next three questions, we assume that the genotypes of both individuals are unknown.

3. Given that the marker is in a region with IBD = 2, the probability that A and B are homozygous for the same allele, is
 - a) 0
 - b) $p_1^2 + p_2^2 + p_3^2$
 - c) $p_1(1 - p_1) + p_2(1 - p_2) + p_3(1 - p_3)$
 - d) $p_1(1 - p_1)^2 + p_2(1 - p_2)^2 + p_3(1 - p_3)^2$
 - e) 1
 4. Given that the marker is in a region with IBD = 1, the probability of a *full match* (i.e., A and B have the same genotype) is
 - a) 0
 - b) $p_1^2 + p_2^2 + p_3^2$
 - c) $p_1(1 - p_1) + p_2(1 - p_2) + p_3(1 - p_3)$
 - d) $p_1(1 - p_1)^2 + p_2(1 - p_2)^2 + p_3(1 - p_3)^2$
 - e) 1
 5. Given that the marker is in a region with IBD = 1, the probability of a *partial match* (i.e., exactly one shared allele) is
 - a) 0
 - b) $p_1^2 + p_2^2 + p_3^2$
 - c) $p_1(1 - p_1) + p_2(1 - p_2) + p_3(1 - p_3)$
 - d) $p_1(1 - p_1)^2 + p_2(1 - p_2)^2 + p_3(1 - p_3)^2$
 - e) 1
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Part II: A case of cousins

Emma and Carlos are about to get married, but suspect that they are related to each other. They consult a geneticist, who types them with 15 standard STR markers. The resulting genotypes are given in the file *cousins-data.txt*, along with the physical location of each marker. Allele frequencies can be found in *db.txt*.

Note: Recall that linkage is to be ignored in LR calculations. The locations are only used in Exercise 8.

6. Use all 15 markers to compute the LR comparing the hypothesis that Emma and Carlos are first cousins, to the unrelated alternative. The total LR, rounded to two decimals, is
- 0.10
 - 0.75
 - 1.00
 - 3.14
 - 13.14

In a more exhaustive approach the geneticist compares the following 5 hypotheses, illustrated in Figure 3:

- H1: first cousins
- H2: first cousins once removed
- H3: second cousins
- H4: third cousins
- H5: unrelated

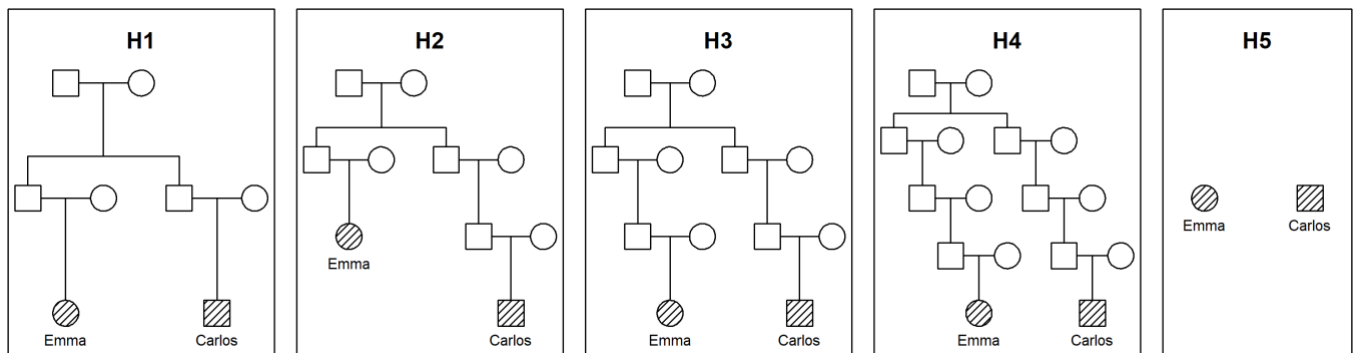


Figure 3. Hypotheses for Question 7

7. The LR comparing the most likely hypothesis with the second most likely, is approximately
- 1.00
 - 1.12
 - 1.45
 - 3.61
 - 10.18

Not completely satisfied with the STR analysis, Emma and Carlos undergo whole-genome sequencing, and the geneticist uses this to identify regions of IBD sharing between them. The result is given in the file *cousins-ibd.txt*, reproduced in its entirety below. In all segments Emma and Carlos share one haplotype, i.e., IBD = 1.

Chr	startMB	endMB	startCM	endCM
3	55.28	98.00	71.71	108.40
4	187.98	189.44	199.80	202.89
5	170.68	180.75	174.66	197.08
9	101.98	112.07	101.00	112.06
11	37.09	98.83	55.18	99.72
11	122.33	131.01	127.17	143.25
12	8.23	15.26	20.31	32.13

8. Of the 15 STR markers, the number that lie in an IBD region is
- 0
 - 1
 - 2
 - 3
 - 4
9. The observed proportion k_1 of the autosome with IBD status 1, is approximately
- 1.1%
 - 1.8%
 - 2.7%
 - 4.3%
 - 6.3%
10. Of the following relationships, the one whose κ_1 is closest to the observed k_1 for Emma and Carlos, is
- first cousins
 - first cousins once removed
 - second cousins
 - third cousins
 - unrelated
11. According to the DNA painter tool, <https://dnainter.com/tools/sharedcmv2>, the IBD sharing between Emma and Carlos is *not compatible* with (i.e., outside the reported range of)
- first cousins
 - first cousins once removed
 - second cousins
 - third cousins
 - several of the above
12. Figure 4 shows the distribution of IBD segments in 200 simulations of first, second and third cousins. Based on this plot, the observed data is only compatible (in the sense of being inside the 95% data ellipse) with
- first cousins (FC)
 - second cousins (SC)
 - third cousins (TC)
 - first and second cousins
 - second and third cousins

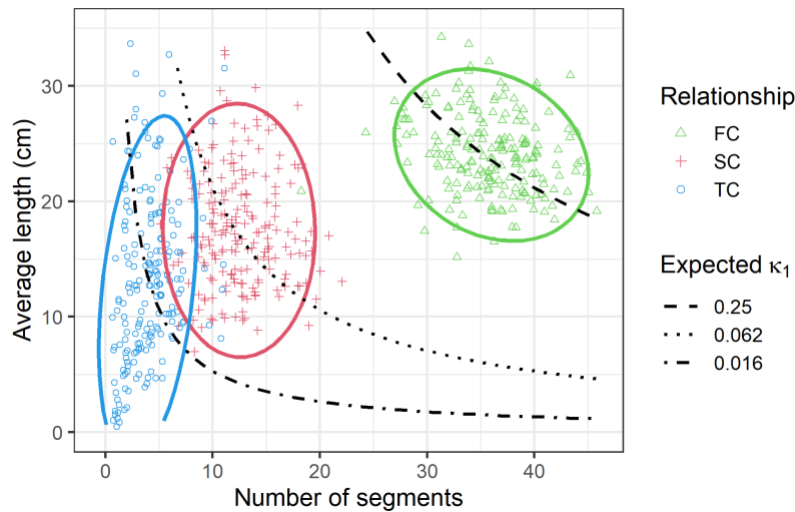


Figure 4. IBD distributions for 1st, 2nd and 3rd cousins

Part III: A case of sibship

This case involves 4 male individuals, labelled A, B, C and D. It is believed that all four have the same mother, but the paternities are unclear. Genotypes for A, B, C, D at 23 forensic markers can be found in *sibship-data.txt*, with allele frequencies in *db.txt* as before.

13. The LR comparing A and B being full siblings versus half siblings, is approximately
 - a) 0.68
 - b) 1.00
 - c) 431.47
 - d) 133506.3
 - e) None of the above

14. The LR comparing C and D being full siblings versus half siblings, is approximately
 - a) 0.68
 - b) 1.00
 - c) 431.47
 - d) 133506.3
 - e) None of the above

We now make the following assumptions:

- i) all four have the same mother
- ii) C and D are full siblings
- iii) each pair among A,B,C,D is either half or full siblings, with no further relationships or inbreeding in the pedigree

15. The number of possible hypotheses (pedigrees) connecting A,B,C,D is
 - a) 5
 - b) 6
 - c) 7
 - d) 8
 - e) 9

16. Let H^* denote the hypothesis that all four are full siblings. Assuming a flat prior on set of hypotheses from the previous question, the posterior probability of H^* given the marker data, is approximately
- 0.00
 - 0.01
 - 0.73
 - 0.99
 - 1.00

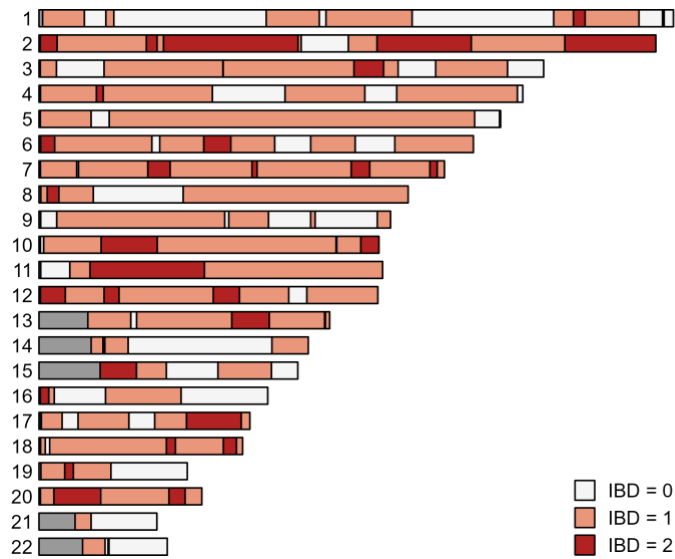


Figure 5. Segments of identity by descent between alleged siblings A and B.

To further investigate the relationship between A and B, whole-genome sequencing is performed, and their IBD segments are determined as shown in Figure 5. The coordinates can be found in the file *sibs-ibd.txt*.

17. In terms of genetic length, the proportion k_1 of the autosome with IBD status 1, is
- 0.25
 - 0.30
 - 0.45
 - 0.50
 - 0.60
18. In the IBD triangle in Figure 6, the realised relationship between A and B corresponds to the point
- P1
 - P2
 - P3
 - P4
 - P5

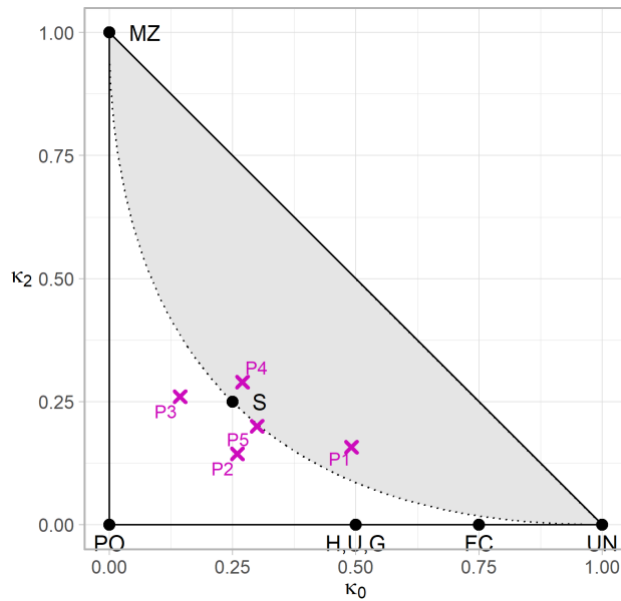


Figure 6. IBD triangle with alternatives for the realised relationship between A and B.

19. Based on the IBD segments, the realised kinship coefficient φ_R between A and B is approximately
- 0.14
 - 0.19
 - 0.22
 - 0.25
 - 0.26

The histogram in Figure 7 shows the realised kinship coefficient in 5000 simulated pairs of full siblings, closely approximated by a normal distribution with mean $\mu = 0.25$ and standard deviation $\sigma = 0.018$ (dashed red curve).

20. Compared with the normal approximation for full siblings, the observed φ_R falls at the
- 0th percentile
 - 5th percentile
 - 10th percentile
 - 15th percentile
 - 20th percentile

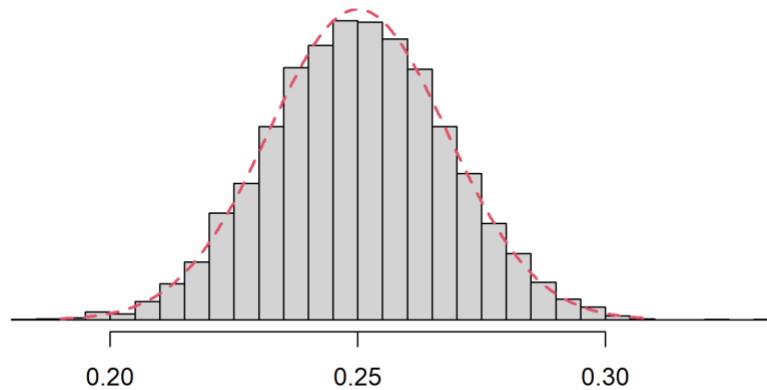


Figure 7. Simulations of the realised kinship between full siblings

GOOD LUCK!