

Covariance between relatives: A reminder

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1 Covariance between relatives

The covariance between relatives for a purely additive genetic model with two loci in LD is briefly sketched out. The development involves covariance terms between individuals at the same locus, and covariance terms between individuals at different loci. The term for covariances at the same locus is derived first.

1.1 Covariance at a single locus

An example motivates the general case. Imagine a locus denoted A . The genotype of a father is A_1A_2 and of a mother A_3A_4 . Consider two offspring from these parents, and the possible number of alleles shared identical by descent (IBD) between the two. There are 16 possible genotype combinations for the two offspring genotypes (arranged in a 4×4 table, where the columns are the possible genotypes for offspring 1, and the rows the possible genotypes for offspring 2). The number of alleles shared IBD between the two offspring i and j , N_{ij} , can take the following values

- $N_{ij} = 2$ (4 cases out of 16)
- $N_{ij} = 1$ (8 cases out of 16)
- $N_{ij} = 0$ (4 cases out of 16)

Therefore

$$\begin{aligned} E(N_{ij}) &= 0 \Pr(N_{ij} = 0) + 1 \Pr(N_{ij} = 1) + 2 \Pr(N_{ij} = 2) \\ &= 1 \frac{1}{2} + 2 \frac{1}{4} = 1 \end{aligned}$$

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and the expected proportion of alleles shared IBD is

$$\frac{E(N_{ij})}{2} = a_{ij} \quad (1)$$

where a_{ij} is also known as the expected additive genetic relationship between i and j , which is the element in the i th row and j th column of the additive genetic relationship matrix A . In the present example the expected proportion is $a_{ij} = 0.5$, the expected number is 1, but the two full-sibs can share 0, 1 or 2 alleles IBD, with probabilities 1/4, 1/2 and 1/4, respectively.

Denote the additive genetic value, or breeding value of individual j

$$g_j = \alpha z_j$$

where α is the additive genetic effect for a locus (or additive effect of a gene substitution), and z_j is the centred genotypic code for the locus. Due to the centring of z

$$E(g_j|\alpha) = \alpha E(z_j) = 0.$$

The additive genetic variance in the population contributed by the locus is

$$V_g = E(g_j^2|\alpha) = \alpha^2 \text{Var}(z_j).$$

Consider the covariance between offspring i and j , conditional on N_{ij} . There are three possible outcomes

- $N_{ij} = 0$,

$$\begin{aligned} \text{Cov}(g_i, g_j | N_{ij} = 0) &= E(g_i, g_j | N_{ij} = 0) - E(g_i | N_{ij} = 0) E(g_j | N_{ij} = 0) \\ &= E(g_i | N_{ij} = 0) E(g_j | N_{ij} = 0) - E(g_i | N_{ij} = 0) E(g_j | N_{ij} = 0) = 0, \end{aligned}$$

because if individuals do not share alleles IBD, the g 's are independent.

- $N_{ij} = 1$,

$$\text{Cov}(g_i, g_j | N_{ij} = 1) = \frac{1}{2} V_g,$$

the gametic variance.

- $N_{ij} = 2$,

$$\text{Cov}(g_i, g_j | N_{ij} = 2) = V_g,$$

the additive genetic variance at the locus. These three cases can be written compactly as

$$\text{Cov}(g_i, g_j | N_{ij}) = \frac{N_{ij}}{2} V_g, \quad N_{ij} = 0, 1, 2. \quad (2)$$

Then, marginally with respect to N_{ij} ,

$$\begin{aligned}
\text{Cov}(g_i, g_j) &= E[\text{Cov}(g_i, g_j | N_{ij})] + \text{Cov}[E(g_i | N_{ij}) E(g_j | N_{ij})] \\
&= E[\text{Cov}(g_i, g_j | N_{ij})] \\
&= \frac{E(N_{ij})}{2} Vg \\
&= a_{ij} Vg
\end{aligned} \tag{3}$$

where the last line uses (1).

1.2 Covariance involving different loci

Let Θ_{ij} denote the coefficient of coancestry, equal to the probability that alleles from two loci k and l in the randomly drawn gametes from individuals i and j , are IBD. The computation of Θ_{ij} involves the rate of recombination between loci k and l and the number of generations back to the common ancestor of individuals i and j .

At locus k , individual i has genotype z_{ik} , coded as $(0, 1, 2)$ and at locus l , individual j has genotype z_{jl} , also originally coded as $(0, 1, 2)$. Let $z_{ikm} = 0, 1$ and $z_{ikp} = 0, 1$, be the binary random variables representing the maternal and paternal gametic contributions to z_{ik} with similar coding for z_{jl} . Then

$$\begin{aligned}
z_{ik} &= z_{ikm} + z_{ikp}, \\
z_{jl} &= z_{jlm} + z_{jlp}.
\end{aligned}$$

Assume that these binary random variables are centred so that $E(z_{ikm})$, say, is equal to zero, which renders z_{ik} and z_{jl} also centred.

The covariance between z_{ik} and z_{jl} is

$$\text{Cov}(z_{ik}, z_{jl}) = \text{Cov}(z_{ikm} + z_{ikp}, z_{jlm} + z_{jlp}). \tag{4}$$

Let the binary random variable W take the value 1, if a randomly drawn gamete from i is IBD with a randomly drawn gamete from j , and zero otherwise. Then $\Pr(W = 1) = \Theta_{ij}$. There are 4 terms contributing to (4) and all have the following form:

$$\begin{aligned}
\text{Cov}(z_{ikm}, z_{jlm}) &= E(z_{ikm}, z_{jlm}) - E(z_{ikm})E(z_{jlm}) \\
&= E_W[E(z_{ikm}, z_{jlm} | W)] \\
&= E(z_{ikm}, z_{jlm} | W = 1) \Pr(W = 1) + E(z_{ikm}, z_{jlm} | W = 0) \Pr(W = 0) \\
&= D_{kl} \Theta_{ij},
\end{aligned} \tag{5}$$

where D_{kl} , the linkage disequilibrium parameter between loci k and l , is here the covariance between the maternal allele at locus k , and the maternal allele at locus l , and Θ_{ij} is the probability that the gametes drawn are IBD. The second term in the third line vanishes when $W = 0$, because if the gametes are not IBD, they are independent, $E(z_{ikm} | W = 0) =$

$E(z_{ikm}) = 0$ and $E(z_{ikm}, z_{jlm}|W = 0) = E(z_{ikm})E(z_{jlm}) = 0$. Summing over all 4 terms yields

$$\begin{aligned} \text{Cov}(z_{ik}, z_{jl}) &= 4D_{kl}\Theta_{ij} \\ &= 2\tilde{a}_{ij}D_{kl}, \end{aligned} \tag{6}$$

where \tilde{a}_{ij} is the expected additive genetic relationship between i and j , since $\tilde{a}_{ij} = 2\Theta_{ij}$. The covariance between additive genetic values of individuals i and j is

$$\text{Cov}(\alpha_k z_{ik}, \alpha_l z_{jl} | \alpha_k, \alpha_l) = 2\tilde{a}_{ij} \alpha_k \alpha_l D_{kl}. \tag{7}$$

In (6) and (7) \tilde{a}_{ij} is used to distinguish it from a_{ij} in (3). The latter involves the probability of IBD of alleles at single loci, whereas the former considers the probability of IBD at pairs of alleles in gametes from two loci.

1.3 Remarks

The covariance between relatives in multiloci systems is a subject of difficult entry. An exact general treatment involving only pairs of loci constitutes a formidable challenge leading to unwieldy expressions, as shown by Weir and Cockerham (1977). The curious reader may wish to glance with awe at formula (6) for the genetic variance in their article, that is almost two pages long! Results assuming lack of inbreeding, epistasis and assortative mating, but accounting for dominance, linkage, and for the dynamics of the linkage disequilibrium parameter over generations, lead to simpler expressions and are given by Weir et al. (1980).

References

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