# 1.3.1 A SAMPLE OF GENES

- Consider a single genetic locus, with two codominant alleles A and B.
- Suppose each independent gene has allelic type A with probability q. We say q is the (population) allele frequency of allele A.
- For a random sample of n genes from the population, the number of A alleles is T ~ Bin(n,q).
- That is Pr(T=t) is proportional to q^t (1-q)^{n-t}.
- The obvious estimator of q is T/n.
- This estimator is unbiased since E(T/n) = nq/n = q.
- Its variance is q(1-q)/n which in fact is the smallest possible variance for any unbiased estimator.

## 1.3.2 Likelihood estimation of q

- The log-likelihood is  $\lambda(q) = t \log (q) + (n-t) \log(1-q)$ .
- · So differentiating the log-likelihood
- $\lambda'(q) = (t/q) (n-t)/(1-q) = n/(q(1-q))((t/n) q)$
- So the maximum likelihood estimator (MLE) is t/n.
- Differentiating again, we find the second derivative:
- $\lambda''(q) = -(t/q^2) (n-t)/(1-q)^2$  and
- $-E(\lambda''(q)) = n/q + n/(1-q) = n/(q(1-q))$
- This is the Fisher information, and the (large-sample) variance of the MLE is  $-1/E(\lambda''(q))$ .
- Here, q(1-q)/n is the variance for any sample size.
- For large n, MLEs are approx unbiased, and have approx the smallest possible variance.

#### **1.3.3 A SAMPLE OF INDIVIDUALS**

- Suppose we sample n individuals, and that n1 have genotype AA, n2 have genotype AB and n3 have genotype BB. n1+n2+n3 =n.
- Then we have (2n1 +n2) genes of allelic type A , in a sample of 2n genes.
- We can estimate q by (2n1 +n2)/2n, but properties of the estimator depend on the model for genotype frequencies:
- The log-likelihood is n1 log(P(AA)) + n2 log(P(AB)) + n3 log(P(BB)).

## 1.3.4 Four examples

- (i) The two genes in an individual must be of the same allelic type (n2=0): complete dependence. The estimator is n1/n and in effect we have a sample of n genes.
- (ii) Hardy-Weinberg equilibrium (HWE); independence of the allelic types of the two genes within an individual. So P(AA) = q<sup>2</sup>, P (AB)= 2q(1-q) and P(BB) = (1-q)<sup>2</sup>.
- (iii) A mixture of (i) and (ii): see 2.2.4.
- (iv) A mixture of subpopulations in HWE: see 1.3.5.

#### **1.3.5 POPULATION STRUCTURE**

- Suppose populations i proportions αi , each in HWE, with qij the freq of allele Aj in population i.
- The overall allele frequencies are weighted average of subpopulation allele frequencies.
- The overall genotype freqs are weighted average of subpopulation HWE frequencies.
- We can show that overall there is excess of each homozygote relative to overall HWE. This excess is known as the Wahlund variance.
- We can show that in total there are fewer heterozygotes than under HWE.
- Details of equations are on the next page.

Genotype frequencies under population structure:

First  $Pr(A_j) = q_{\cdot j} = \sum_i \alpha_i q_{ij}$ , and so

$$\begin{aligned} \Pr(A_{j}A_{j}) - (\Pr(A_{j}))^{2} &= \sum_{i} \alpha_{i}q_{ij}^{2} - q_{.j}^{2} \\ &= \sum_{i} \alpha_{i}(q_{ij} - q_{.j})^{2} \geq 0 \\ \Pr(A_{j}A_{l}) - 2\Pr(A_{j})\Pr(A_{l}) &= 2(\sum_{i} \alpha_{i}q_{ij}q_{il} - q_{.j}q_{.l}) \\ &= 2\sum_{i} \alpha_{i}(q_{ij} - q_{.j})(q_{il} - q_{.l}) \end{aligned}$$

For two alleles, let  $q_{i1} = q_i$ ,  $q_{i2} = 1 - q_i$ ,  $q_i = q$ . If  $\sigma_f^2 = \sum_i \alpha_i (q_i - q)^2$ , then the three genotype freqs are  $q^2 + \sigma_f^2$ ,  $2q(1-q) - 2\sigma_f^2$  and  $(1-q)^2 + \sigma_f^2$ .

#### 1.4.1 ESTIMATION: case of HWE

- Log-likelihood is λ(q) = log L(q)
  = n1 log(q<sup>2</sup>) + n2 log(2q(1-q)) + n3log((1-q)<sup>2</sup>)
  = (2 n1 + n2) log (q) + (n2 + 2n3) log(1-q)
- The MLE of q is (2 n1 + n2)/2n.
- If T = 2 n1+n2, T ~ Bin(2n,q). --- back to binomial sampling, with a sample size 2n genes.
- Hence, var(T/2n) = q(1-q)/2n.
- Note: One generation of random mating establishes HWE, since, by definition, the two genes in an individual are copies of independently sampled parental genes.

## 1.4.2 Case of a recessive allele

- t = n1 of type AA, and n-t not of type AA.
- Assuming HWE, P(AA) = q<sup>2</sup>, so log-likelihood is λ(q) = t log(q<sup>2</sup>) + (n-t) log (1 -q<sup>2</sup>)
- Differentiating  $\lambda'(q) = 2t/q 2 (n-t) q/(1-q^2)^2$ =  $(2/q(1-q^2)) (t - n q^2)$
- So the MLE of q is  $\sqrt{(t/n)}$ .
- Why should this be expected?
- Now T ~ Bin(n, q^2), but how can we find the variance of this MLE?

#### 1.4.2 ctd: Using Fisher Information

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- $\lambda''(q) = -2t/q^2 2(n-t)/(1-q^2) 4 (n-t) q^2 / (1-q^2)^2$ .
- $E(-\lambda''(q)) = 2n + 2n + 4 q^2 n/(1-q^2)$ =  $4n/(1-q^2)$
- Thus, the variance of the MLE of q is approx. (1-q<sup>2</sup>)/4n.
- Note this is larger than q(1-q)/2n.
- Note (i) We have to make assumptions (HWE),.
  (ii) the variance of the estimator is larger.
  (iii) Using the Fisher information we can measure the information lost.

## 1.4.3 Data on relatives

- We consider just mother-baby pairs and assume HWE.
- See next page for the conditional and joint probabilities.
- $l(q) = n00 log(q^3) + n01 log(q^2 (1-q)) + n10 log (q^2 (1-q)) + n11 log(q(1-q)) + n12 log (q(1-q)^2) + n21 log (q(1-q)^2) + n22 log ((1-q)^3) = (3 n00 + 2 (n01 + n10) + n11 + n12 + n21 )log q + (3 n22 + 2 (n21+n12) + n11 + n10 + n01) log (1-q) = mA log q + mB log (1-q).$
- The MLE of q is mA/(mA +mB), where (mA +mB) = 3n - n11 and mA = (3 n00 + 2 (n01 +n10) + n11 + n12 + n21).

#### Parent and child probabilities

par	prob	ch AA	ch AB	ch BB			
AA	q^2	q	(1-q)	0			
AB	2q(1-q)	q/2	1/2	(1-q)/2			
BB	(1-q)^2	0	q	(1-q)			
	ch AA	ch AB	ch BB	Data counts			
AA	q^3	q^2(1-q)	0	n00	nC	)1	0
AB	q^2(1-q)	q(1-q)	q(1-q)^2	n10	n1	1	n12
BB	0	q(1-q)^2	(1-q)^3	0	n2	21	n22

## 1.4.4 Alternatives to the MLE

- The MLE is ``best", but there are simpler estimators that are not bad.
- One is to use only founders (here the moms): estimate q by (2 nAA + nAB)/2n where nAA and nAB are the numbers of AA and AB moms., (nAA = n00+n01).
- Or, use everyone, disregarding relationship: estimate q by (2 mAA + mAB)/4n, where mAA and mAB are is total numbers of AA and AB individuals. (mAA = 2 n00 + n01 + n10).
- These are both unbiased estimators, but asymptotically the MLE has smaller variance.