# Geographic Population Structure

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In Buri's [1] drift experiment, heterozygosity ($H$) declined. At the same time, the variance ($V$) among populations increased. We already have a model describing the first of these phenomena. Here, we consider the second—the variance among populations—assuming throughout that genetic drift is the only evolutionary force at work.

First a few terms. In the initial generation, the allele frequency was $p_0$. We will treat this as a constant, not a random variable. In generation $t$, the corresponding quantity $p_t$ is a random variable, the randomness having been introduced by the process of genetic drift. For convenience, we set $q_t = 1 - p_t$. The heterozygosity in generation $t$ (also a random variable) is $H_t = 2p_t q_t$. We are interested in the evolutionary change that has happened since generation 0.

## 1 Genetic drift does not change the expected allele frequency

Let us begin with the expected value of $p_1$, the allele frequency in the generation 1. According to the urn model, the number of copies of allele $A$ in each subpopulation is a Binomial random variable with mean $Np_0$. The expected allele frequency is thus $Np_0/N = p_0$. This demonstrates a remarkable fact: the expected allele frequency is unchanged by genetic drift. This is true not only of the first generation, but of each succeeding generation. No matter how many generations are involved, the allele frequency of each subpopulation is a random variable whose expected value is $p_0$. In symbols,

$$E[p_t] = p_0. \tag{1}$$

## 2 The Wahlund Principle and $F_{ST}$

In contrast to $p_t$, the variance $V_t$ has an expected value that does change with time. To model the variance, we begin with its definition:

$$V_t = E[p_t^2] - p_0^2$$

This is just the standard definition of the variance, modified slightly in light of Eqn. 1. Note that, in view of this definition,

$$E[p_t^2] = V_t + p_0^2 \tag{2}$$

We'll need this fact in a minute.

Let us turn now to the heterozygosity. Its expected value in generation $t$ is

$$
\begin{aligned}
E[2p_t q_t] &= 2E[p_t] - 2E[p_t^2] \\
&= 2p_0 - 2(V_t + p_0^2) \quad \text{using Eqns. 1–2} \\
&= 2p_0 q_0 - 2V_t \tag{3}
\end{aligned}
$$

This is another important fact: average heterozygosity in generation $t$ will be smaller than that in generation 0. The amount of the reduction is *exactly twice* the variance of group allele frequencies about their expected value $p_0$. This is known as Wahlund's principle [3]. It shows that there is a close and necessary connection between the decline of heterozygosity (shown on the left in Fig. 1) and the increase in variance (shown on the right). In effect, genetic drift converts heterozygosity into variance among groups.

It is often useful to express the absolute reduction in heterozygosity ($2V_t$) as a proportion of the original heterozygosity ($2p_0 q_0$). This proportional reduction is

$$F_{ST} = \frac{2V_t}{2p_0 q_0} = \frac{V_t}{p_0 q_0} \tag{4}$$

The notation $F_{ST}$ was introduced by Sewall Wright [4] and is now conventional within population genetics. Gillespie defines $F_{ST}$ using different notation
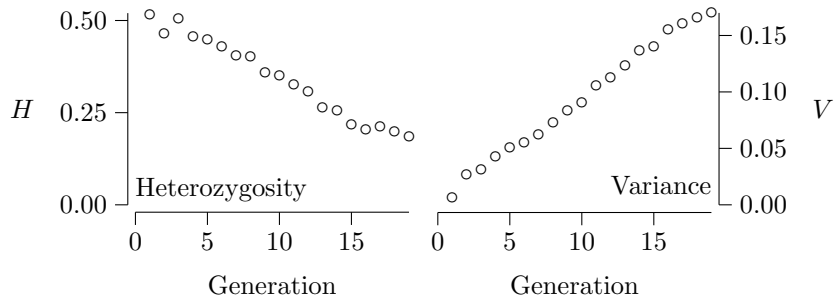
Figure 1: In Buri's [1] drift experiment, heterozygosity ($H$) declined. At the same time, the variance ($V$) among populations increased. Data are from Buri's series I and are tabulated in Table 1.

(his Eqn. 5.3), which was introduced by Nei [2]. Although the two definitions look different, they are interchangeable. In view of Eqn. 4, we can re-express Eqn. 3 as

$$E[2p_t q_t] = 2p_0 q_0 (1 - F_{ST}) \qquad (5)$$

Now $E[2p_t q_t]$ is the expected heterozygosity in generation $t$, and $2p_0 q_0$ is that in generation 0. Thus, Eqn. 5 says that $F_{ST}$ is the proportional reduction in heterozygosity caused by genetic drift. There are corresponding increases in expected homozygosity. The expected frequencies of the three genotypes at a biallelic locus are:

| Genotype | Frequency |
|---|---|
| $AA$ | $E[p_t^2] = p_0^2 + p_0 q_0 F_{ST}$ |
| $Aa$ | $E[2p_t q_t] = 2p_0 q_0 (1 - F_{ST})$ |
| $aa$ | $E[q_t^2] = q_0^2 + p_0 q_0 F_{ST}$ |

These formulas are identical to those for pedigree inbreeding, because genetic drift is a form of inbreeding.

## 3 Model of completely isolated subpopulations

In Eqn. 5, we express the expected heterozygosity ($2p_t q_t$) in generation $t$ as a function of that in generation 0. Early in the semester, we derived a similar result:

$$E[2p_t q_t] = 2p_0 q_0 \left(1 - 1/2N\right)^t \qquad (6)$$

To refresh your memory, see Gillespie's Eqn. 2.3. These equations look very different, yet both are correct. They provide a simple way to derive the rule by which $F_{ST}$ changes with time. Set Eqns. 5 and 6 equal to each other, and solve for $F_{ST}$. You will discover that

$$F_{ST} = 1 - \left(1 - 1/2N\right)^t$$

$$\approx 1 - e^{-t/2N} \qquad (7)$$

The last line above uses the approximation that $e^x \approx 1 + x$ if $x$ is near zero. Eqn. 7 applies when the populations are totally separated. It shows that $F_{ST}$ increases according to a very simple rule, increasing gradually toward its maximal value, 1.0.

## 4 Migration in addition to drift

If there is migration between populations, we cannot use any of the results in section 3. This case demands a different theory, which is discussed by Gillespie. Under the "island-model" of population structure, $F_{ST}$ converges toward an equilibrium at which

$$F_{ST} = \frac{1}{4Nm + 1} \qquad (8)$$

as shown on Gillespie's p. 136.

We have two equations for $F_{ST}$. Eqn. 8 refers to equilibrium under the island model, and Eqn. 7 to the case of totally isolated populations. In the exercises below, we will consider human data under these two extreme cases.

## Exercises

1. Plot the $V_t$ data in Table 1, to make a graph like that on the right side of Fig. 1.

2. Combine Eqns. 4 and 7 to obtain a formula for the variance, $V_t$. Plot this formula as a line in your graph, assuming as Buri did that $2N = 18$. How well do Buri's variance data fit the model?

3. In the continental human populations, $F_{ST} \approx 1/9$. Use this value to estimate $Nm$ (under the equilibrium model).

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Table 1: Data from series I of Buri's [1] drift experiment, as plotted in Fig. 1. Key: $t$, generation; $H_t$, mean heterozygosity ($2pq$) within subpopulations; $V_t$, variance among group allele frequencies.

| $t$ | $H_t$ | $V_t$ | $t$ | $H_t$ | $V_t$ |
|---|---|---|---|---|---|
| 0 | 1.000 | 0.000 | 10 | 0.348 | 0.090 |
| 1 | 0.514 | 0.006 | 11 | 0.325 | 0.105 |
| 2 | 0.464 | 0.026 | 12 | 0.305 | 0.112 |
| 3 | 0.504 | 0.031 | 13 | 0.263 | 0.123 |
| 4 | 0.456 | 0.042 | 14 | 0.255 | 0.136 |
| 5 | 0.448 | 0.050 | 15 | 0.216 | 0.140 |
| 6 | 0.428 | 0.055 | 16 | 0.202 | 0.155 |
| 7 | 0.403 | 0.062 | 17 | 0.210 | 0.160 |
| 8 | 0.402 | 0.072 | 18 | 0.197 | 0.165 |
| 9 | 0.358 | 0.083 | 19 | 0.183 | 0.170 |

4. Now assume that the human continental populations have been totally isolated, and use the observed $F_{ST}$ to estimate $t/2N$. Then convert this into an estimate of the time in years since the human populations separated. (Assume that $N = 10,000$ and that generations are 25 years long.)

# References

[1] P. Buri. Gene frequency in small populations of mutant *Drosophila*. *Evolution*, 10:367–402, 1956.

[2] Masatoshi Nei. Analysis of gene diversity in subdivided populations. *Proceedings of the National Academy of Sciences, USA*, 70(12):3321–3323, 1973.

[3] Sten Wahlund. Composition of populations and of genotypic correlations from the viewpoint of population genetics. In Kenneth M. Weiss and Paul A. Ballanoff, editors, *Demographic Genetics*, pages 224–263. Wiley, New York, 1975. [Translation].

[4] Sewall Wright. The genetical structure of populations. *Annals of Eugenics*, 15:323–354, 1951.