Introduction

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Introduction and Motivating Examples

Bayesian Analysis of Multinomial Data

Derivation of the Posterior and Prior Specification
Inference for Parameters of Interest

Analysis of HWE Data

Appendix: BF Detail

Introduction 0000000

- In this lecture we will consider the Bayesian modeling of multinomial data
- The examination of Hardy-Weinberg equilibrium will be used to motivate the multinomial model.
- Again, conjugate priors will be used, though sampling from the posterior will be emphasized as a method for flexible inference.
- Bayes factors will be used as a measure of evidence for hypothesis testing.

Motivating Example: Testing for HWE

- For simplicity we consider a diallelic marker, and suppose we obtain a random sample of genotypes for n individuals.
- The form of the data is

	Genotype			Total
	A_1A_1	A_1A_2	A_2A_2	
Count	n_1	n_2	n_3	n
Population Frequency	q_1	q_2	q_3	1

- So the model contains 3 probabilities (which sum to 1) q₁, q₂, q₃; hence, there are 2 free parameters.
- Suppose the proportions of alleles A_1 and A_2 in a given generation are p_1 and $p_2 = 1 p_1$.
- In terms of q₁, q₂, q₃:

$$p_1 = q_1 + \frac{q_2}{2}$$

$$p_2 = \frac{q_2}{2} + q_3$$

Motivating Example: Testing for HWE

- HWE is the statistical independence of an individual's alleles at a locus.
- Under HWE, the probability distribution for the genotype of an individual in the next generation is:

$$\begin{tabular}{c|cccc} & Genotype \\ \hline A_1A_1 & A_1A_2 & A_2A_2 \\ \hline Proportion & p_1^2 & $2p_1p_2$ & p_2^2 & 1 \\ \hline \end{tabular}$$

 Reasons for deviation from HWE include: small population size, selection, inbreeding and population structure.

A Toy Example

Example:

- Consider the data $n_1 = 88, n_2 = 10, n_3 = 2.$
- Are these frequencies consistent with HWE?
- The MLEs are:

$$\widehat{q}_1 = 0.88 \quad \widehat{q}_2 = 0.10 \quad \widehat{q}_3 = 0.02$$

 $\widehat{p}_1 = 0.93 \quad \widehat{p}_2 = 0.07$

For these data the exact p-value for

$$H_0: q_1 = p_1^2, \quad q_2 = 2p_1p_2, \quad q_3 = p_2^2$$

is 0.0654.

Critique of Non-Bayesian Approach

- Testing for HWE is carried out via (asymptotic, i.e., large sample) χ^2 tests or exact tests.
- χ^2 tests require very large sample sizes for accurate p-values.
- The exact test can be computationally expensive to perform, when there are many alleles/samples.
- Under the null of HWE, the discreteness of the test statistic causes difficulties.
- In general, how to decide on a significance level? The level should be a function of sample size (and in particular should decrease as sample size increases), but how should it be chosen?
- Estimation depends on asymptotic approximations (i.e., large sample sizes).
- Estimation also difficult due to awkward constraints on parameters (particularly with many alleles).

Bayes Theorem

	Genotype			Total
	A_1A_1	A_1A_2	A_2A_2	
Count	<i>n</i> ₁	n_2	n_3	n
Population Frequency	q 1	q 2	9 3	1

- The multinomial with three counts is known as the trinomial distribution.
- We have three parameters, q_1 , q_2 , q_3 , but they sum to 1, so that effectively we have two parameters.
- We write $\mathbf{q} = (q_1, q_2, q_3)$ to represent the vector of probabilities, and $\mathbf{n} = (n_1, n_2, n_3)$ for the data vector.
- Via Bayes Theorem:

$$p(q|n) = \frac{\Pr(n|q) \times p(q)}{\Pr(n)}$$
Posterior \propto Likelihood \times Prior

Elements of Bayes Theorem: The Likelihood

- We assume n independent draws with common probabilities $\mathbf{q} = (q_1, q_2, q_3)$.
- In this case, the distribution of n_1, n_2, n_3 is multinomial:

$$Pr(n_1, n_2, n_3 | q_1, q_2, q_3) = \frac{n!}{n_1! n_2! n_3!} q_1^{n_1} q_2^{n_2} q_3^{n_3}.$$
 (1)

- For fixed n, we may view (1) as a function of q this is the likelihood function.
- The maximum likelihood estimate (MLE) is

$$\widehat{\boldsymbol{q}} = \left(\frac{n_1}{n}, \frac{n_2}{n}, \frac{n_3}{n}\right).$$

 The MLE gives the highest probability to the observed data, i.e. maximizes the likelihood function.

The Dirichlet Distribution as a Prior Choice for a Multinomial **q**

- Once the likelihood is specified we need to think about the prior distribution.
- We require a prior distribution over (q₁, q₂, q₃) not straightforward since the three probabilities all lie in [0,1], and must sum to 1.
- A distribution that satisfies these requirements is the dirichlet distribution, denoted dirichlet(v₁, v₂, v₃) and has density:

$$p(q_1, q_2, q_3) = \frac{\Gamma(v_1 + v_2 + v_3)}{\Gamma(v_1)\Gamma(v_2)\Gamma(v_3)} \times q_1^{v_1 - 1} q_2^{v_2 - 1} q_3^{v_3 - 1}$$

$$\propto q_1^{v_1 - 1} q_2^{v_2 - 1} q_3^{v_3 - 1}$$

where $\Gamma(\cdot)$ denotes the gamma function.

The Dirichlet Distribution as a Prior Choice for a Multinomial **q**

• The dirichlet(v_1, v_2, v_3) prior:

$$p(q_1, q_2, q_3) = \frac{\Gamma(v_1 + v_2 + v_3)}{\Gamma(v_1)\Gamma(v_2)\Gamma(v_3)} \times q_1^{v_1 - 1} q_2^{v_2 - 1} q_3^{v_3 - 1}$$

$$\propto q_1^{v_1 - 1} q_2^{v_2 - 1} q_3^{v_3 - 1}.$$

- $v_1, v_2, v_3 > 0$ are specified to reflect prior beliefs about (q_1, q_2, q_3) .
- The dirichlet distribution can be used with general multinomial distributions (i.e. for k = 2, 3, ... categories).
- The beta distribution is a special case of the dirichlet when there are two categories only.

Dirichlet Prior

The mean and variance are

$$E[q_i] = \frac{v_i}{v_1 + v_2 + v_3} = \frac{v_i}{v}$$

$$var(q_i) = \frac{E[q_i](1 - E[q_i])}{v_1 + v_2 + v_3 + 1} = \frac{E[q_i](1 - E[q_i])}{v + 1}$$

for i = 1, 2, 3, where $v = v_1 + v_2 + v_3$.

- Large values of v increase the influence of the prior.
- The dirichlet has a single parameter only (v) to control the spread for all of the dimensions, which is a deficiency.
- The quartiles may be empirically calculated from samples.





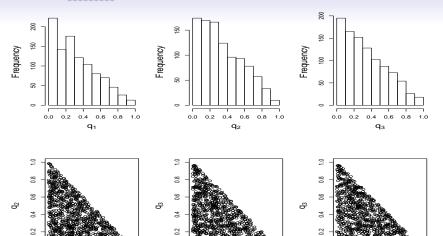


Figure 1: Samples from a dirichlet (1, 1, 1) distribution. The mean is $(\frac{1}{3}, \frac{1}{3}, \frac{1}{3})$.

0.2

0.2 0.4

i 0.6 **q**₁

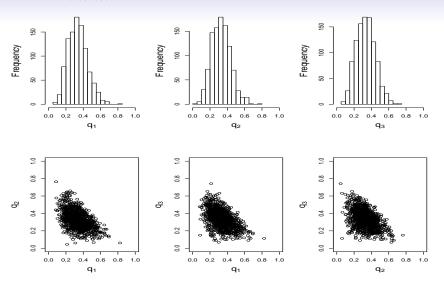


Figure 2: Samples from a dirichlet (6,6,6) distribution. The mean is $(\frac{1}{3},\frac{1}{3},\frac{1}{3})$.

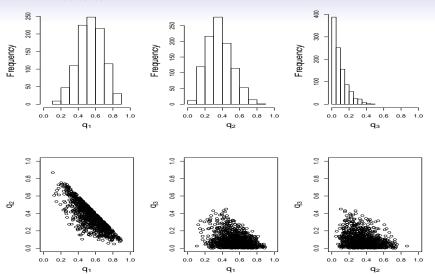


Figure 3: Samples from a dirichlet(6, 4, 1) distribution. The mean is $(\frac{6}{11}, \frac{4}{11}, \frac{1}{11}) = (0.55, 0.36, 0.09)$.

 Combining the Dirichlet prior, dirichlet(v₁, v₂, v₃), with the multinomial likelihood gives the posterior:

$$p(q_1, q_2, q_3 | \mathbf{n}) \propto \Pr(\mathbf{n} | \mathbf{q}) \times p(\mathbf{q})$$

$$\propto q_1^{n_1} q_2^{n_2} q_3^{n_3} \times q_1^{v_1 - 1} q_2^{v_2 - 1} q_3^{v_3 - 1}$$

$$= q_1^{n_1 + v_1 - 1} q_2^{n_2 + v_2 - 1} q_3^{n_3 + v_3 - 1}.$$

This distribution is another Dirichlet:

$$dirichlet(n_1 + v_1, n_2 + v_2, n_3 + v_3).$$

• Notice: "as if" we had observed counts $(n_1 + v_1, n_2 + v_2, n_3 + v_3)$.

Choosing a Prior

The posterior mean for the expected proportion of counts in cell i is, for i = 1, 2, 3:

$$E[q_{i}|\mathbf{n}] = \frac{n_{i} + v_{i}}{n + v}$$

$$= \frac{n_{i}}{n} \frac{n}{n + v} + \frac{v_{i}}{v} \frac{v}{n + v}$$

$$= MLE \times W + Prior Mean \times (1 - W)$$

where $n = n_1 + n_2 + n_3$, $v = v_1 + v_2 + v_3$.

The weight W is

$$W = \frac{n}{n+v}$$

which is the proportion of the total information (n + v) that is contributed by the data (n).

Choosing a Prior

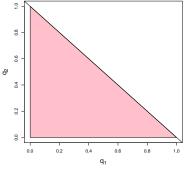
Recall the prior mean is

$$\left(\frac{v_1}{v}, \frac{v_2}{v}, \frac{v_3}{v}\right)$$

- These forms help to choose v₁, v₂, v₃.
- As with the beta distribution we may specify the prior means, and the relative weight that the prior and data contribute: n and v are on a comparable scale.
- For example, suppose we believe that event 1 is four times as likely as each of event 2 or event 3.
- Then we may specify the means in the ratios 4:1:1.
- Suppose n=24 and we wish to allow the prior contribution to be a half of this total (and therefore a third of the complete information). Then the prior sample size is v=12 and the prior mean requirement gives

$$v_1 = 8, v_2 = 2, v_3 = 2.$$

A Uniform Prior



An obvious choice of parameters is $v_1 = v_2 = v_3 = 1$ to give a prior that is uniform over the simplex:

$$\pi(q_1, q_2, q_3) = 2$$

for

$$0 < q_1, q_2, q_3 < 1, \quad q_1 + q_2 + q_3 = 1$$

Note: not uniform over all parameter of interests, as we see shortly

Simple HWE Example

The data is

$$n_1 = 88, n_2 = 10, n_3 = 2.$$

We assume a flat dirichlet prior on the allowable values of q:

$$v_1 = v_2 = v_3 = 1.$$

 This gives the posterior as dirichlet(88 + 1, 10 + 1, 2 + 1) with posterior means:

$$E[q_1|\mathbf{n}] = \frac{1+88}{3+100} = \frac{89}{103}$$

$$E[q_2|\mathbf{n}] = \frac{1+10}{3+100} = \frac{11}{103}$$

$$E[q_3|\mathbf{n}] = \frac{1+2}{3+100} = \frac{3}{103}.$$

Note the similarity to the MLEs of

$$\left(\frac{88}{100}, \frac{10}{100}, \frac{2}{100}\right)$$
.

Simple HWE Example

- We continue with this example and now examine posterior distributions.
- We generate samples from

$$dirichlet(88 + 1, 10 + 1, 2 + 1).$$

- As posterior summaries we display, in Figure 4:
 - Histograms of the 3 univariate marginal distributions $p(q_1|\mathbf{y})$, $p(q_2|\mathbf{y})$, $p(q_3|\mathbf{y})$.
 - Scatterplots of the 3 bivariate marginal distributions $p(q_1, q_2|\mathbf{y})$, $p(q_1, q_3|\mathbf{y})$, $p(q_2, q_3|\mathbf{y})$.
- On each plot we indicate the MLEs for the general model, i.e. the non-HWE model (in red) and under the assumption of HWE (in blue).

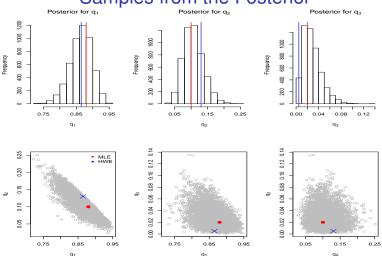


Figure 4: Univariate and bivariate posterior distributions for $\mathbf{n} = (88, 10, 2)$. MLEs in red for the general model and in blue for the HWE model.

- As expected with a sample size of n = 100 and a flat prior, the MLEs lie close to the center of the posteriors.
- Note the asymmetry of the posteriors.
- Asymptotic confidence intervals of the form \$\hat{q}_i \pm 1.96 \times \sec{(\hat{q}_i)}\$ would be symmetric.

- In the context of a binomial sampling model and interest in a particular point (for example, $\theta=0.5$) we could examine intervals for θ .
- In a multinomial context the situation is more complex; shortly we will examine Bayes factors to carry out hypothesis testing.

Parameters of Interest

	Genotype			Total
	A_1A_1	A_1A_2	A_2A_2	
Population Frequency	q_1	q_2	q 3	1

- Rather than q₁, q₂, q₃, we may be interested in other parameters
 of interest.
- In the HWE context: Let X_1 and X_2 be 0/1 indicators of the A_1 allele for the two possibilities at a locus; so $X_1 = X_2 = 1$ corresponds to the genotype A_1A_1 .
- The covariance between X₁ and X₂ is the disequilibrium coefficient:

$$D = q_1 - p_1^2$$

Under HWE $q_1 = p_1^2$, and the covariance is zero.

• Another quantity of interest (Shoemaker et al., 1998) is

$$\psi = \frac{q_2^2}{q_1 q_3}$$

Under HWE, $\psi = 4$.

Parameters of Interest

The inbreeding coefficient is

$$f = \frac{q_1 - p_1^2}{p_1 p_2}$$

- The variance of X_1 and X_2 is $p_1(1 p_1) = p_1p_2$ and so f is the correlation.
- We may express q_1, q_2, q_3 as

$$q_1 = p_1^2 + p_1(1 - p_1)f$$

$$q_2 = 2p_1(1 - p_1)(1 - f)$$

$$q_3 = (1 - p_1)^2 + p_1(1 - p_1)f$$

 Positive values of f indicate an excess of homozygotes (and may indicate inbreeding), while negative values indicate an excess of heterozygotes.

Parameters of Interest

- Each of D, ψ and f are complex functions of q_1, q_2, q_3 and given a Dirichlet prior for the latter do not have known posterior forms.
- The "flat" prior for **q**, dirichlet(1, 1, 1), does not correspond to a flat prior for D, f, ψ , as Figure 5 shows.

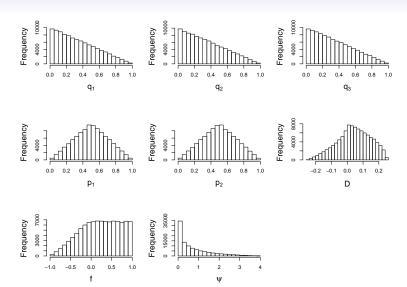


Figure 5: Samples from a dirichlet(1,1,1) for various functions.

Implied Prior on Functions of Interest

- Very important point: As we saw in the binomial development, you can't be "flat" on every scale, but the implications of particular priors can be assessed via simulation.
- We emphasize that we are not uniform on the marginal distributions for q_i , since these follow

$$beta(v_i, v - v_i)$$

distributions, i.e. beta(1,2) if $v_1 = v_2 = v_3 = 1$.

- The priors on the measures of distance from HWE are far from uniform.
- For example, with a "flat" Dirichlet prior dirichlet(1,1,1) the prior probability that f > 0 is 0.67.

Previous Bayesian approaches include:

- Altham (1971).
- Pereira and Rogatko (1984).
- Lindley (1988).
- Shoemaker et al. (1998).
- Montoya-Delgado et al. (2001).
- Consonni et al. (2008).
- Wakefield (2010).
- Consonni et al. (2011).
- Puig et al. (2017).

Bayes factors for HWE

- Recall that Bayes factors measure the evidence in a sample for one hypothesis, as compared to an alternative.
- We derive the Bayes factor for multinomial data in the context of testing for HWE.
- We wish to test

 H_0 : HWE versus H_1 : Not HWE.

 We need to specify priors on the null and alternatives, and then calculate the Bayes factor:

$$\frac{\Pr(\boldsymbol{n}|H_0)}{\Pr(\boldsymbol{n}|H_1)}$$

where p_1 and (q_1, q_2) are the parameters under the null and alternative, respectively.

The HWE Bayes Factor

 The Bayes factor, measuring the evidence in the data for the null, as compared to the alternative is:

$$\mathsf{BF} \ = \ \frac{2^{n_2} \Gamma(w) \Gamma(2n_1 + n_2 + w_1) \Gamma(v_1) \Gamma(v_2) \Gamma(v_3) \Gamma(n_2 + 2n_3 + w_2) \Gamma(n + v)}{\Gamma(w_1) \Gamma(w_2) \Gamma(2n + w) \Gamma(v) \Gamma(n_1 + v_1) \Gamma(n_2 + v_2) \Gamma(n_3 + v_3)}$$

- This appears complex, but is just a function of the observed data, and the prior inputs, and can be easily evaluated¹.
- If BF > 1(< 1) the data are more (less) likely to have come from the null.
- Can be readily extended to k > 2 alleles.
- We next consider a formal decision rule.

 $^{^{1}}$ When we work out a χ^{2} tail area we don't worry about the form of the distribution we just use the relevant function in our favorite software

Bayesian Decision Theory

 Decision as to reject H₀ in favor of H₁ depends on the costs of making the two types of error:

		Decision		
		Report H ₀	Report H ₁	
Truth	H_0	0	C_{I}	
atti	H_1	C_{II}	0	

- Costs of making the two types of error C_I is the cost of a type I error and C_{II} the cost of a type II error.
- The decision theory solution is to report H₁ if:

Posterior Odds of
$$H_0 = \mathsf{BF} \times \mathsf{Prior} \; \mathsf{Odds} < \frac{C_{II}}{C_I} = R$$

so that we only need to consider the ratio of costs *R*.

• If $\frac{C_{II}}{C_{I}} = 4$ (type II errors four times as bad as type I errors) then report H_{1} if

Posterior Odds of
$$H_0 < 4$$
,

i.e. if

$$Pr(H_1 | data) > 0.2.$$

A Simple Example

We again consider the data $n_1 = 88$, $n_2 = 10$, $n_3 = 2$.

These data give a p-value of 0.0654.

With "flat" conjugate Dirichlet priors ($w_1 = w_2 = v_1 = v_2 = v_3 = 1$) we obtain a Bayes factor of 1.54 so that the data are 50% more likely under the null than the alternative, so the evidence in favor of H_0 is not strong.

With a prior probability of the null π_0 , to give a prior odds of $\pi_0/(1-\pi_0)$, we have

Posterior Odds of
$$H_0 = \mathsf{BF} \times \frac{\pi_0}{1 - \pi_0}$$

Hence, with $\pi_0 = 0.5$ the posterior odds equal the Bayes factor, i.e. 1.54.

The posterior probability of the null is

$$\frac{1.54}{1+1.54}=0.61.$$

This probability is very sensitive to the prior on the null.

For example, with $\pi_0=2/3$ we obtain a posterior odds of $1.54 \times 2=3.08$ to give a posterior probability on the null of

$$\frac{3.08}{1+3.08}=0.75.$$

Figure 6 shows functions of interest for the (88, 10, 2) example.

Note that the asymptotic confidence interval for f is (-0.1032, 0.5632).

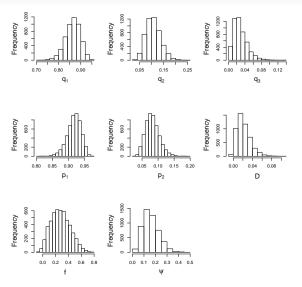


Figure 6: Posterior inference for simple HWE example.

The above prior specifications are convenient analytically, but in some situations we would like to perform Bayesian inference using priors that are based on contextual information.

If we are really interested in the deviations from HWE of a sample from a particular population, then we may have strong prior information which perhaps can be represented through a prior on the inbreeding coefficient *f*.

Under the null we have a single probability p_1 , the probability of an A_1 allele.

Under the alternative we may specify the prior

$$\pi(p_1, f) = \pi(p_1) \times \pi(f|p_1)$$

where the conditioning allows the constraints on f:

$$f_{\min} = \max\left(-\frac{p_1}{1 - p_1}, -\frac{1 - p_1}{p_1}\right) < f < 1$$

Unfortunately there is no closed form Bayes factor available, instead we describe a simulation-based technique — the rejection algorithm.

Let θ denote the parameters with prior distribution $\pi(\theta)$, and let $\widehat{\theta}$ be the MLE and $p(\mathbf{y}|\widehat{\theta})$ the maximized likelihood.

Then the rejection algorithm (e.g., Wakefield, 2013, Chapter 3) proceeds as follows:

- 1. Generate $U \sim U(0,1)$ and $\theta \sim \pi(\theta)$, independently.
- 2. Accept θ if

$$U < \frac{p(\mathbf{y}|\boldsymbol{\theta})}{p(\mathbf{y}|\widehat{\boldsymbol{\theta}})},$$

otherwise reject θ .

3. Return to 1.

The resultant $\theta^{(s)}$, s = 1, ..., S, are an independent sample from the posterior $p(\theta|y)$.

The rejection algorithm may be very inefficient if the prior and likelihood differ substantially (e.g., prior is dispersed and/or likelihood is peaked).

An estimate of the normalizing constant (required for Bayes factor calculation) is given by

$$\widehat{p}(\mathbf{y}) = \frac{1}{S} \sum_{s=1}^{S} p(\mathbf{y}|\boldsymbol{\theta}^{(s)})$$

where $\theta^{(s)} \sim \pi(\cdot)$.

Note that this only requires samples from the prior — the rejection algorithm is not needed.

In the HW context the maximized likelihood is available in closed form.

Specific Non-Conjugate Priors

Recall the prior is

$$\pi(p_1, f) = \pi(p_1) \times \pi(f|p_1)$$

Two components:

- For $\pi(p_1)$ we take a Be (w_1, w_2) prior.
- For $\pi(f|p_1)$ we transform to

$$\phi = \log((f - f_{\min})/(1 - f))$$

and assume $\phi|p_1$ is normal.

HWE Example Revisited

We again consider the data $n_{11} = 88$, $n_{12} = 10$, $n_{22} = 2$.

These data give a *p*-value of 0.0654. The MLE for *f* is 0.23 with asymptotic standard error 0.17. MLE of HWE proportions: (0.865, 0.130, 0.05).

With flat conjugate Dirichlet priors we obtained a Bayes factor of 1.54 so that the data are 50 % more likely under the null, but the evidence is low.

HWE Example Revisited

We assume that the 50% point of the prior on f is 0, and the 95% point is 0.5

We obtain a Bayes factor of 0.29 so that the data are 3.4 times as likely under the alternative, but the evidence is again weak.

The posterior probability that f > 0 is 0.98.

The difference between the priors is that the non-conjugate version gives more weight close to where the data are located.

Graphical Summaries

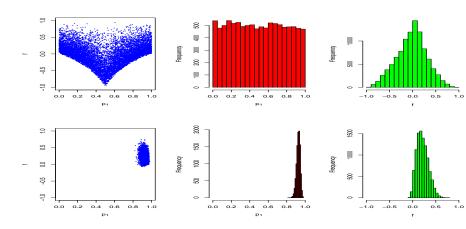


Figure 7: Prior (top) and Posterior (bottom). Notice the clear constraint in the top left plot.

In contrast to estimation, in which the prior influence generally disappears with increasing sample size, the Bayes factor remains influenced by the prior.

To illustrate we multiply the data of the previous example by different factors.

Factor	Conj BF	Non-conj BF	Post prob $f > 0$	<i>p</i> -value
1	1.54	0.50	0.984	0.0654
2	0.40	0.070	0.997	0.0089
5	0.0039	0.000639	1	3.6×10^{-5}
10	1.2×10^{-6}	1.8×10^{-7}	1	5.3×10^{-9}

The conjugate and non-conjugate Bayes factors remain different (though the conclusions are the same).

 The R package HWEBayes implements the rejection algorithm and importance sampling (a numerical integration technique), for testing and estimation in the HWE context:

http://cran.r-project.org/web/packages/HWEBayes/index.html

- The vignette contains a worked example.
- Code for a four-allele example is here:
 http://faculty.washington.edu/jonno/HWEBayesFourAllele.R
- More details of the methodology: Wakefield (2010).

Conclusions

- The dirichlet distribution is convenient but quite inflexible as a prior distribution.
- Alternative priors are more difficult to specify since they are on scales that are more difficult to interpret (e.g. the logistic-normal distribution).
- Bayes factors are sensitive to the prior.
- Monte Carlo sampling is a powerful tool for inference.
- For multiple alleles computation is slow whether the approach is frequentist or Bayesian.

Derivation of Bayes Factor for Assessing HWE

 We need to specify priors on the null and alternatives, and then calculate the Bayes factor:

$$\frac{\Pr(\mathbf{n}|H_0)}{\Pr(\mathbf{n}|H_1)} = \frac{\int \Pr(\mathbf{n}|p_1)p(p_1)dp_1}{\int \Pr(\mathbf{n}|q_1,q_2)p(q_1,q_2)dq_1dq_2}$$

where p_1 and (q_1, q_2) are the parameters under the null and alternative, respectively.

- Under the null we have a single parameter, and under the alternative two.
- Important point: When Bayes factors are evaluated we need to include the normalizing constants.

- Under H₀ and H₁ we must take care to evaluate the probability of the same data, n₁, n₂, n₃.
- Under the null,

$$\Pr(\mathbf{n}|p_1) = \Pr(n_1, n_2, n_3|p_1) = \frac{n!2^{n_1}}{n_2! n_{12}! n_3!} p_1^{2n_1+n_2} (1-p_1)^{n_2+2n_3}.$$

• With a Be(w_1, w_2) prior on p_1 :

$$Pr(n_{1}, n_{2}, n_{3}|H_{0}) = \int Pr(\mathbf{n}|p_{1}) \times p(p_{1})dp_{1}$$

$$= \frac{n!2^{n_{2}}\Gamma(w)\Gamma(2n_{1} + n_{2} + w_{1})\Gamma(n_{2} + 2n_{3} + w_{2})}{n_{1}!n_{2}!n_{3}!\Gamma(w_{1})\Gamma(w_{2})\Gamma(2n + w)}$$
(2)

This is the probability of the observed data under the null.

HWE Bayes Factor

The Bayes factor is

$$\frac{\Pr(\boldsymbol{n}|H_0)}{\Pr(\boldsymbol{n}|H_1)}$$

and we have just given the form of the numerator.

- We now turn to the denominator.
- Under the alternative we assume *q* ∼ dirichlet(*v*₁, *v*₂, *v*₃).
- The probability of the data under the alternative is:

$$\Pr(n_{1}, n_{2}, n_{3} | H_{1}) = \int \Pr(\mathbf{n} | q_{1}, q_{2}) \times p(q_{1}, q_{2}) dq_{1} dq_{2}$$

$$= \frac{n! \Gamma(v) \Gamma(n_{1} + v_{1}) \Gamma(n_{2} + v_{2}) \Gamma(n_{3} + v_{3})}{n_{1}! n_{2}! n_{3}! \Gamma(v_{1}) \Gamma(v_{2}) \Gamma(v_{3}) \Gamma(n + v)}.$$
(3)

 Again, just a probability distribution, which we may evaluate for any realization of (n₁, n₂, n₃).

The HWE Bayes Factor

 Hence, the Bayes factor, measuring the evidence in the data for the null, as compared to the alternative is:

$$BF = \frac{\Pr(n_1, n_2, n_3 | H_0)}{\Pr(n_1, n_2, n_3 | H_1)}$$

$$= \frac{2^{n_2} \Gamma(w) \Gamma(2n_1 + n_2 + w_1) \Gamma(v_1) \Gamma(v_2) \Gamma(v_3) \Gamma(n_2 + 2n_3 + w_2) \Gamma(n + v)}{\Gamma(w_1) \Gamma(w_2) \Gamma(2n + w) \Gamma(v) \Gamma(n_1 + v_1) \Gamma(n_2 + v_2) \Gamma(n_3 + v_3)}$$

which is (2) divided by (3).

- This appears complex, but is just a function of the observed data, and the prior inputs, and can be easily evaluated.
- If BF > 1(< 1) the data are more (less) likely to have come from the null.
- Can be readily extended to k > 2 alleles.

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