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What Are the Odds? Modern Relevance and Bayes Factor Solutions for MacAlister’s Problem from the 1881 Educational Times

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Abstract

In 1881, Donald MacAlister posed a problem in the *Educational Times* that remains relevant today. The problem centers on the statistical evidence for the effectiveness of a treatment based on a comparison between two proportions. For the data presented by MacAlister, the $p$ value is not significant and from a classical perspective it therefore remains unclear whether and to what extent the data favor the null hypothesis. A brief historical sketch is followed by a discussion of two default Bayesian solutions, one based on a one-sided test between independent rates, and one based on a one-sided test between dependent rates. We demonstrate the current-day relevance of MacAlister’s original question with a modern-day example about the effectiveness of an educational program.

**Keywords:** Contingency tables; Bayes factor; Evidence.
What Are the Odds? Modern Relevance and Bayes Factor

Solutions for MacAlister’s Problem from the 1881 *Educational Times*

In 1881, Donald MacAlister posed a famous problem in the *Educational Times*, a problem that represents one of the earliest instances concerning the comparison of two proportions in small samples:

Of 10 cases treated by Lister’s method, 7 did well and 3 suffered from blood-poisoning; of 14 cases treated with ordinary dressings, 9 did well and 5 had blood-poisoning; what are the odds that the success of Lister’s method was due to chance?

It is clear that the answer to this question is of considerable statistical relevance, far exceeding the specifics of the problem at hand. In modern-day educational research, one often wants to quantify the evidence for the effectiveness of a new program or instruction; if the new program seems to result in a beneficial outcome, the immediate question is identical that posed by MacAlister: “what are the odds that the success of the new method is due to chance”?

Before proceeding, a few remarks are in order. First and foremost, the traditional $p$ value cannot be used to address MacAlister’s question, as the $p$ value is based on a single hypothesis (i.e., the null hypothesis) and therefore does not produce an odds. Moreover, for the MacAlister data the $p$ value is not even close to being significant ($p > .70$ for all standard classical methods); based on this $p$ value, one might be tempted to believe that the evidence supports the null hypothesis. But to what degree? Second, MacAlister did not pose his question with mathematical exactness, and this requires that it has to be interpreted at least to some extent. The solutions offered in 1882 demonstrate how easy it
is to misunderstand the problem (Dale, 1999, pp. 435–438; Winsor, 1948). Third, the problem as posed cannot be solved without involving the prior odds that Lister’s method is effective (another issue that the \( p \) value ignores). To appreciate the importance of the prior odds, consider the fact that Lister was a famous scientist who had advocated the use of antiseptic dressings to reduce the possibility of post-surgical infection, based on the theory that these infections were caused by germs (Lister, 1867/1967). The idea that antiseptic dressings fail to reduce the rate of post-surgical infection will strike the modern reader as absurd; consequently, the prior odds that the method’s success is due to chance are extremely low. In MacAlister’s example, we have the rare case that we know the answer –that Lister was correct– before we begin, so we can focus without distraction on the evidence presented the data provide about the odds of success. These can then be multiplied by the prior odds as we like in order to obtain the posterior odds, as explained below. Fourth, the results may be presented in familiar form using a contingency table, as presented in Table 1.

The solution proposed by MacAlister was based on a procedure developed by Liebermeister (1877). Denote the probability of recovery by Lister’s method and by the traditional method as \( \theta_l \) and \( \theta_t \), respectively. It is clear that the interest partly concerns the probability of Lister’s method outperforming the traditional method, that is,

\[ \mathcal{H}_+ : p(\theta_l > \theta_t \mid y), \]

where \( y \) denotes the observed data. But what should this probability be compared to? MacAlister assumed independent uniform priors for \( \theta_l \) and \( \theta_t \) and computed \( p(\theta_l > \theta_t \mid y) = 0.59825 \). MacAlister compared this proportion to its complement, \( p(\theta_l < \theta_t \mid y) \), and concluded “we may wager nearly 3 to 2 that the difference in the results is not due to chance”. We may understand “due to chance” as “due to mere chance.” Note that MacAlister’s solution quantifies evidence in favor of the effectiveness of the treatment, despite the fact that the \( p \) value is not even close to being significant.

The hypothesis that the difference is due to mere chance, however, plays no role in
MacAlister’s solution, as no prior mass is assigned to the invariance or general law that
the treatments are equally effective: that is $\mathcal{H}_0 : \theta_l = \theta_t$ (Wrinch & Jeffreys, 1921). By
assigning no zero prior mass to mere chance (i.e., the hypothesis that the treatments are
in fact equally effective), the question at hand cannot be answered. MacAlister’s odds of
“nearly 3 to 2” address a different question, namely “what are the odds that the success of
Lister’s method is due to its superiority versus its inferiority over the traditional
method?”.

**Two Default Bayes Factor Solutions**

In order to address MacAlister’s problem, we contrast two hypotheses. The first
hypothesis represents the assertion that both treatments are equally effective, that is,
$\mathcal{H}_0 : \theta_l = \theta_t$; the second hypothesis represents the assertion that Lister’s treatment is
superior to the standard treatment, that is, $\mathcal{H}_+ : \theta_l > \theta_t$.

We now wish to compute the evidence that the data provide for $\mathcal{H}_+$ over $\mathcal{H}_0$. Recall
that Bayes’ rule can be recast as follows:

$$
\frac{p(\mathcal{H}_+ \mid y)}{p(\mathcal{H}_0 \mid y)} = \frac{p(\mathcal{H}_+)}{p(\mathcal{H}_0)} \times \frac{p(y \mid \mathcal{H}_+)}{p(y \mid \mathcal{H}_0)} = \text{Bayes factor}.
$$

Thus, data $y$ are used to update the prior odds to posterior odds. The assessment of prior
odds is inherently subjective and depends on background information that informs one’s
initial skepticism about the hypotheses under consideration. Indeed, in commenting on
MacAlister’s solution to his own problem, Miss Elizabeth Blackwood stated –quite
correctly, in our view– “I will merely remark that Dr. MacAlister would probably feel less
satisfied as to the correctness of his result, if Lister were not the eminent man of science
he is, but some superstitious old woman who, while really expert in dressing wounds,
relied for protection against blood-poisoning mainly upon some mysterious charms and
incantations”. MacAlister’s response made it clear that he did not consider prior odds to
factor into the problem at all: “Miss Elizabeth Blackwood has perhaps not read my solution: there is no symbol in it representing Mr. Lister’s science. For algebraical purposes I might substitute Mumbo Jumbo for Lister throughout, as I substituted the letter A, and no step of the reasoning on which alone the result depends would be altered”. Here we adhere to the intention from MacAlister and focus on the Bayes factor $BF_{+0}$, that is, the change from prior to posterior model odds brought about by the data (Jeffreys, 1961).

The Bayes factor

$$BF_{+0} = \frac{p(y \mid H_+)}{p(y \mid H_0)}$$

expresses the evidence in the data for the one-sided hypothesis $H_+ : \theta_l > \theta_t$, asserting that Lister’s treatment is superior to the standard treatment, against the point hypothesis $H_0 : \theta_l = \theta_t$, asserting that both treatments are equally effective. In order to compute $p(y \mid H_+)$ and $p(y \mid H_0)$ we need to assign priors to the model parameters $\theta_l$ and $\theta_t$. This can be accomplished in many ways. Here we explore two default solutions: a model in which $\theta_l$ and $\theta_t$ are independent, and a model in which $\theta_l$ and $\theta_t$ are dependent. Both models yield a similar outcome.

**Solution I: Prior independence of $\theta_l$ and $\theta_t$**

The default Bayes factor approach contrasts the single-rate model $H_0$ to the dual-rate model $H_1$. The dual-rate model usually does not include information about the predicted direction of the effect. However, with any two-sided Bayes factor in hand a simple correction produces the desired one-sided version (see appendix for details).

To obtain the default two-sided Bayes factor $BF_{10}$ we assume that under the dual-rate model, each rate has an independent uniform prior distribution ranging from 0 to 1 (Gunel & Dickey, 1974; Jeffreys, 1935; de Braganca Pereira & Stern, 1999). Based on this default prior specification, the one-sided Bayes factor can be computed easily in JASP
(jasp-stats.org), a free and open-source statistical software program with a graphical user interface familiar to users of SPSS. The same result is available for R users through the BayesFactor package (Morey & Rouder, 2015). Figure 1 shows a JASP screenshot that displays the analysis input options as well as the statistical output.

As shown in Figure 1, the $BF_{0+} \approx 1.8$, which means that the observed data are almost twice as likely under the single rate model $H_0$ than under the dual-rate model $H_+$. According to the classification scheme proposed by Jeffreys (1961, Appendix B), this level of evidence is “not worth more than a bare mention”. Assuming that the single rate model and the dual-rate model are equally likely a priori, we can use MacAlister’s terminology and state that “we may wager nearly 2 to 1 that the difference in the results is not due to mere chance”. Regardless of the inconclusive nature of the evidence in this particular instance, this result does answer MacAlister’s question.

**Solution II: Prior dependence of $\theta_l$ and $\theta_t$**

An alternative model specification views the two rates as dependent (e.g., Howard, 1998). Such a dependence is reasonable in many such problems; the probabilities of the two groups are typically similar because effects are not arbitrarily large. Instead of thinking about the separate probabilities at which the two groups recover, it is convenient to instead frame the problem in terms of an overall recovery rate, and the difference of the two groups from that overall rate (see also Kass & Vaidyanathan, 1992). This induces a reasonable dependency between the two groups.

The two parameters—the overall rate, and the difference between the two groups—is best expressed on the probit scale, in order to avoid the common problem of compression of the probability scale at the extremes:

$$
\Phi^{-1}(\theta_l) = \Phi (\mu + \delta/2), \quad \text{and}
$$

$$
\Phi^{-1}(\theta_t) = \Phi (\mu - \delta/2),
$$
where $\Phi^{-1}$ denotes the the probit transformation; that is, the inverse of the standard normal cumulative distribution function. Parameter $\mu$ is the overall recovery rate on the probit scale, and $\delta$ is one-half the difference between the two groups and represents the effect of interest. Next, $\mu$ and $\delta$ are assigned normal priors. For demonstration, we assign a $\text{Normal}(0,.707)$ prior distribution to $\mu$ and $\delta$ is assigned a folded (i.e., positive-only, to incorporate knowledge about the hypothesized direction of the effect) normal prior with mean 0 and standard deviation $\sigma$. The test will not be very sensitive to the prior choice on $\mu$; however, a reasonable prior on $\delta$ is important, as it is the parameter of interest. For demonstration we choose $\sigma = \sqrt{2}$ as a default value. We choose these settings because they yield the same marginal priors on $\theta_1$ and $\theta_2$ as under Solution I.

The Bayes factor of interest is based on a comparison between two models, $H_0 : \delta = 0$ versus $H_+ : \delta > 0$. To obtain $BF_{+0}$, we use Gaussian quadrature. There are a number of other ways to obtain the Bayes factor, including importance sampling and Markov chain Monte Carlo sampling. An interactive application to compute and visualize the Bayes factor can be found at richarddmorey.shinyapps.io/probitProportions, and R code to compute the Bayes factor and plots can be downloaded at gist.github.com/richarddmorey/4c7a408a45c3045ab949.

Analyzing the MacAlister data, the top panel of Figure 2 shows the prior and posterior distributions for the difference $\delta$ under $H_+$. At the value of interest, $\delta = 0$, the posterior distribution is about 2.3 times as high as the prior distribution, and, hence, $BF_{0+} \approx 2.3$ (Dickey & Lientz, 1970; Wagenmakers, Lodewyckx, Kuriyal, & Grasman, 2010). As for the default analysis using the independent priors, the Bayes factor indicates that the data are more likely to occur under $H_0$ than under $H_+$, but the strength of this evidence is not impressive.
A Modern Example from Education Research

To underscore the relevance of MacAlister’s problem for current-day research we turn to a study by Tuckman and Kennedy (2011) published in *The Journal of Experimental Education*. These authors investigated the effect of a learning strategies course on students’ academic performance as quantified by several dependent variables including retention rate, that is, the proportion of students that return to college the following year. The data showed that from a total of $n_1 = 351$ first-year students who took the course, 93.4% returned to college the next year; from a total of $n_2 = 351$ matched students who did not take the course, 85.5% returned. Table 2 shows the data in the form of a contingency table. For these data, MacAlister’s question is again relevant: *What are the odds that the success of the learning strategies course was due to mere chance?*

We address this question as we did before, by contrasting two hypotheses. The null hypothesis states that the course has no effect, $H_0: \theta_1 = \theta_2$. The alternative hypothesis has direction and states that the course increases the retention rate, $H_+: \theta_1 > \theta_2$. As before, the change from prior to posterior odds for $H_0$ versus $H_+$ is expressed through the Bayes factor $BF_{+0}$.

First, the results from the independent prior analysis (Gunel & Dickey, 1974; Jeffreys, 1935) are displayed in Figure 3. The output shows that $BF_{+0} = 45.83$, meaning that the observed data are 45.83 times more likely to occur under $H_+$ than under $H_0$. According to Jeffreys’ classification scheme, this constitutes “very strong” evidence in favor of the effectiveness of the course on retention rate.

Second, the results from the dependent prior analysis with the probit model are displayed in the bottom panel of Figure 2. As before, the distributions are for the difference $\delta$ under $H_+$. At the value of interest, $\delta = 0$, the prior distribution is about .0142 times as high as the posterior distribution, and, hence, $BF_{+0} \approx 70$. Even though the two methods give slightly different results, they agree that the data provide considerable
support in favor of $H_+$. 

**Conclusion**

We have outlined a Bayesian method to quantify the support that the data provide for the equality or inequality of two rates. In contrast to classical or frequentist procedures based on the $p$ value, the Bayesian method allows one to address the key problem posed by MacAlister in 1881: what are the odds that the success of a particular treatment is based on mere chance? In our solution to MacAlister’s problem, we compared a single rate model $H_0$ against an order-restricted default dual-rate model $H_1$, using two fundamentally different prior specifications. As usual, it should be acknowledged that the default prior distributions can often be enriched and adjusted by incorporating substantive knowledge about the problem at hand. Moreover, in applied settings, one might extend the current framework and use model-averaging to obtain superior predictions (e.g., Hoeting, Madigan, Raftery, & Volinsky, 1999); in addition, one might specify utilities and combine these with the fundamental unknowns in order make the best possible decision in a coherent manner (e.g., Lindley, 1985). Both prediction and decision-making require the consideration of the prior odds for the competing hypotheses, an endeavor that is often inherently subjective.

Despite these reservations, we believe that in many situations the default prior specifications provide an appropriate reference analysis that can be used to replace or supplement the currently dominant framework of $p$-value hypothesis testing. For a range of standard statistical models, such reference analyses can be easily conducted using the R BayesFactor package (Morey & Rouder, 2015) or the free and open-source program JASP (jasp-stats.org). Prominent advantages of the default Bayesian analysis include the possibility to monitor evidence as the data accumulate and the ability to discriminate evidence of absence from the absence of evidence. The problem posed by MacAlister in
1881 is still relevant today, and Bayesian methods such as the one outlined in this article constitute a solution that is theoretically elegant and practically relevant.
References


Liebermeister, C. (1877). Ueber Wahrscheinlichkeitsrechnung in Anwendung auf
therapeutische Statistik. In R. Volkmann (Ed.), *Sammlung klinischer Vortrage no. 110 (innere Medicin no. 39)* (pp. 935–962).


Appendix

Obtaining the One-Sided Bayes Factor

The Bayes factor BF_{+0} can be easily obtained by decomposing it into two parts (Morey & Wagenmakers, 2014; Pericchi, Liu, & Torres, 2008):

\[
BF_{+0} = \frac{p(y \mid \mathcal{H}_+)}{p(y \mid \mathcal{H}_0)} = \frac{p(y \mid \mathcal{H}_+)}{p(y \mid \mathcal{H}_1)} \times \frac{p(y \mid \mathcal{H}_1)}{p(y \mid \mathcal{H}_0)} = BF_{+1} \times BF_{10},
\]

where \(BF_{+1}\) quantifies the evidence for the hypothesis \(\mathcal{H}_+\) that Lister’s method is superior against the undirected hypothesis \(\mathcal{H}_1 : \theta_l \neq \theta_t\) that simply asserts that the two treatments have a different effect.

Equation 1 demonstrates that in order to compute the one-sided Bayes factor \(BF_{+0}\) we multiply the two-sided \(BF_{10}\) by the factor \(BF_{+1}\). Klugkist, Laudy, and Hoijtink (2005) noted that \(BF_{+1}\) equals the ratio of posterior and prior mass under \(\mathcal{H}_1\) that is consistent with the restriction postulated by \(\mathcal{H}_+\). That is,

\[
BF_{+1} = \frac{p(\theta_l > \theta_t \mid y, \mathcal{H}_1)}{p(\theta_l > \theta_t \mid \mathcal{H}_1)} \tag{2}
\]

which simplifies to \(BF_{+1} = 2 \times p(\theta_l > \theta_t \mid y, \mathcal{H}_1)\) whenever the prior distributions do not express any knowledge or preference for the ordering of \(\theta_l\) and \(\theta_t\), meaning that \(p(\theta_l > \theta_t) = 1/2\).
Author Note

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Footnotes

1In a later rejoinder, Miss Blackwood retorted as follows: “As to the right which Dr. MacAlister claims to substitute, if he chooses, Mumbo Jumbo for Lister in his solution, I am inclined to think he has already exercised that right. But, granting this mathematical license of substitution ‘for algebraical purposes’ (a euphemism apparently for juggling purposes) is Lister to be arbitrarily valued at $\frac{1}{2}$ merely because we don't know what other value to assign to him? Poor Lister!”.
Table 1

*MacAlister’s 1881 data displayed as a contingency table.*

<table>
<thead>
<tr>
<th>Method</th>
<th>Did well</th>
<th>Blood poisoning</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Lister</td>
<td>7</td>
<td>3</td>
<td>10</td>
</tr>
<tr>
<td>Traditional</td>
<td>9</td>
<td>5</td>
<td>14</td>
</tr>
<tr>
<td>Total</td>
<td>16</td>
<td>8</td>
<td>24</td>
</tr>
</tbody>
</table>
Table 2

Number of students retained as a function of having attended a learning strategies course.

Data reported in Tuckman and Kennedy (2011).

<table>
<thead>
<tr>
<th>Group</th>
<th>Retained</th>
<th></th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Yes</td>
<td>No</td>
<td></td>
</tr>
<tr>
<td>Course takers</td>
<td>328</td>
<td>23</td>
<td>351</td>
</tr>
<tr>
<td>Non-Course takers</td>
<td>300</td>
<td>51</td>
<td>351</td>
</tr>
<tr>
<td>Total</td>
<td>628</td>
<td>74</td>
<td>702</td>
</tr>
</tbody>
</table>
Figure Captions

Figure 1. A screenshot of JASP input and output for the MacAlister data. The corresponding .jasp file with data and analyses is available at www.ejwagenmakers.com/Code/2015/Listerism.jasp.

Figure 2. Prior and posterior distributions for the difference parameter $\delta$ after applying the dual-rate dependent probit model to the data from MacAlister (top panel) and Tuckman and Kennedy (bottom panel). The Bayes factor equals the ratio of posterior and prior ordinates at $\delta = 0$ (e.g., Wagenmakers et al., 2010).

Figure 3. A screenshot of JASP input and output for the Tuckman and Kennedy (2011) data. The corresponding .jasp file with data and analyses is available at www.ejwagenmakers.com/Code/2015/Tuckman.jasp.
Reply to MacAlister (1881), Figure 1
Reply to MacAlister (1881), Figure 2
Bayesian Contingency Tables

<table>
<thead>
<tr>
<th>Group</th>
<th>Retained</th>
<th>Not Retained</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Course Takers</td>
<td>328</td>
<td>23</td>
<td>351</td>
</tr>
<tr>
<td>Non-Course Takers</td>
<td>300</td>
<td>51</td>
<td>351</td>
</tr>
<tr>
<td>Total</td>
<td>628</td>
<td>74</td>
<td>702</td>
</tr>
</tbody>
</table>

Bayesian Contingency Tables Tests

- $BF_{01}$ independent multinomial: 45.83
- $N$: 702

Note: All tests, hypothesis is group Course Takers greater than group Non-Course Takers.

Odds ratio

- $BF_{01} = 45.83$
- $BF_{10} = 0.022$

Density plot of log(odds ratio)