

SEQUENTIAL BAYES ESTIMATION OF THE DIFFERENCE BETWEEN MEANS

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It is desired to estimate the difference between the means of two independent normal distributions as accurately as possible and in a sequential manner when the total number of observations is fixed. The problem is posed in a Bayesian framework with conjugate prior distributions and squared error loss function. It is shown that the optimal sequential design depends on the ratio of the posterior variances of the two means. There exist constants (dependent on the prior parameters, the number of observations taken from each distribution, and the number of observations remaining) such that when the above-mentioned ratio exceeds this constant it is optimal to select the next observation from one distribution; otherwise it is optimal to select it from the other distribution.

1. Introduction. Suppose there are two experiments ε_1 and ε_2 which generate, independently of each other, sequences of i.i.d. random variables which are generically denoted by X and Y respectively. Let each X have a normal distribution $N(\mu_1, P_1^{-1})$ where P_1 is the precision. Let each Y have a normal distribution $N(\mu_2, P_2^{-1})$. Many different sequential schemes for estimating $\mu_1 - \mu_2$ have been proposed. If the observations are taken in pairs $(X_1, Y_1), (X_2, Y_2)$, etc., then $Z_i = X_i - Y_i$ is normally distributed with mean $\mu_1 - \mu_2$ and variance $P_1^{-1} + P_2^{-1}$. Thus all of the results on sequentially estimating the mean of a single normal distribution are applicable. For example, see Anscombe [1], Chow and Robbins [2], Geertsema [4], Ray [6], Robbins [7], Serfling and Wackerly [9], Simons [10], and Starr [12], [13]. Allowing for unequal sample sizes may increase the accuracy of the estimate or decrease the expected total sample size. Srivastava [11] and Robbins, Simons and Starr [8] have proposed a class of sequential rules incorporating both a sampling scheme and a stopping rule which are asymptotically optimal. The approach in this paper is to suppose that the total number of observations is fixed and to concentrate on the sampling scheme. From this point of view the problem closely resembles the two-armed bandit problem. Prior distributions and a loss function are assigned and the concomitant optimal sequential strategy is examined.

2. The Bayesian model. The vector of means and precisions (μ_1, P_1, μ_2, P_2) is assigned a prior distribution so that (μ_1, P_1) and (μ_2, P_2) are priorly independent with normal-gamma distributions $NG(\eta_1, \tau_1, \alpha_1, \beta_1)$ and $NG(\eta_2, \tau_2, \alpha_2, \beta_2)$.

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Properties of this conjugate family of distributions are well known (cf. DeGroot [3], Section 9.6).

The object is to estimate $\theta = \mu_1 - \mu_2$ with squared error loss, so that if $\hat{\theta}$ is an estimator of θ , the loss is $L(\theta, \hat{\theta}) = (\theta - \hat{\theta})^2$. The estimator $\hat{\theta}$ which minimizes the expected loss (risk) with respect to a prior distribution is $E(\mu_1) - E(\mu_2)$ and the minimum risk itself is $\sigma^2(\mu_1) + \sigma^2(\mu_2)$ (cf. Zacks [15], pages 273-274). These are, respectively, the Bayes estimator and the Bayes risk.

Let N be the fixed total number of trials and let m and n be the number of trials allocated to experiments ε_1 and ε_2 respectively. After all N observations have been completed, the Bayes estimator of $\mu_1 - \mu_2$ is the difference in the means of the posterior marginal distributions of μ_1 and μ_2 ; the Bayes risk is the sum of the variances of these two distributions. Once this is established, the whole problem can be modeled as an adaptive control process.

3. Adaptive control process model. For a description of adaptive control processes (ACP) see Yakowitz [14]. In our situation, the state space may be thought of as the set of quadruples (m, n, A_m, B_n) where $m \geq 0, n \geq 0, m + n \leq N, A_m \geq 0, B_n \geq 0$. Here A_m and B_n are the variances of the posterior marginal distributions of μ_1 and μ_2 respectively and are functions of the sample means, sample variances, and prior parameters. From the properties of the normal-gamma distributions, it follows that

$$(3.1) \quad A_m = \frac{2\beta_1 + (m - 1)S_x^2 + \tau_1 m(\tau_1 + m)^{-1}(\bar{X} - \eta_1)^2}{(\tau_1 + m)(m + 2\alpha_1 - 2)}$$

and

$$(3.2) \quad B_n = \frac{2\beta_2 + (n - 1)S_y^2 + \tau_2 n(\tau_2 + n)^{-1}(\bar{Y} - \eta_2)^2}{(\tau_2 + n)(n + 2\alpha_2 - 2)}.$$

In order that these quantities be finite and nonnegative for all m and n , α_1 and α_2 must be larger than 1. The control set consists of two elements: choose ε_1 or choose ε_2 . At time $t < N$ zero loss is incurred while at time $t = N$ a loss of $A_m + B_n$ is incurred. The loss function may then be written as the sum of these losses as t ranges from 0 to N . The statistical law of motion for this ACP is embodied in the following lemma.

LEMMA 3.1. *Given A_m and B_n , $U = a_m A_m/A_{m+1}$ and $V = b_n B_n/B_{n+1}$ have independent beta distributions $B(\alpha_1 + \frac{1}{2}m, \frac{1}{2})$ and $B(\alpha_2 + \frac{1}{2}n, \frac{1}{2})$ respectively where*

$$(3.3) \quad a_m = \frac{(\tau_1 + m)(m + 2\alpha_1 - 2)}{(\tau_1 + m + 1)(m + 2\alpha_1 - 1)}$$

and

$$(3.4) \quad b_n = \frac{(\tau_2 + n)(n + 2\alpha_2 - 2)}{(\tau_2 + n + 1)(n + 2\alpha_2 - 1)}.$$

PROOF. Tedious algebraic manipulation leads to the fact that

$$(3.5) \quad A_{m+1} = a_m(A_m + (m + 2\alpha_1 - 2)^{-1}(\tau_1 + m + 1)^{-1}(X_{m+1} - \eta_1')^2)$$

where

$$(3.6) \quad \eta_1' = (\tau_1 \eta_1 + m \bar{X}_m) / (\tau_1 + m).$$

Let \mathcal{F}_m be the σ -field generated by X_1, \dots, X_m . From the properties of the normal-gamma family of distributions, X_{m+1} given \mathcal{F}_m is distributed as $C^{\frac{1}{2}}W + \eta_1'$ where

$$(3.7) \quad C = (m + 2\alpha_1 - 2)(\tau_1 + m + 1)(m + 2\alpha_1)^{-1}A_m,$$

η_1' is defined as in (3.6) and W is a random variable having a Student's t -distribution with $2\alpha_1$ df. This means that, given \mathcal{F}_m , A_{m+1} is distributed as $a_m A_m (1 + W^2(m + 2\alpha_1)^{-1})$. Thus U has a beta distribution $B(\alpha_1 + \frac{1}{2}m, \frac{1}{2})$. An analogous argument can be applied to B_{n+1} to get the distribution of V . Furthermore, A_{m+1} and B_{n+1} are independent, so U and V are independent. \square

Because of the Markovian nature of the statistical law of motion and the separability of the loss function for this ACP, the dynamic programming algorithm yields the optimal strategy (cf. Yakowitz [14], Theorem 3.3).

4. The optimal sequential design. Suppose there are k trials remaining where m trials have already been allocated to ε_1 and n trials to ε_2 . Let $\mathcal{F}_{m,n}$ denote the sigma field generated by $(X_1, \dots, X_m, Y_1, \dots, Y_n)$. Unless otherwise stated, all expectations in this section are conditional on $\mathcal{F}_{m,n}$. Also for notational convenience, the arguments A_m, B_n will be suppressed from the functions in this section unless needed. Denote the anticipated risk over the remaining k trials by $R_k(m, n)$. Then

$$(4.1) \quad R_1(m, n) = \min \{E(A_{m+1} + B_n), E(A_m + B_{n+1})\}.$$

From Lemma 3.1 it follows that

$$(4.2) \quad R_1(m, n) = A_m + B_n - \max \{A_m(\tau_1 + m + 1)^{-1}, B_n(\tau_2 + n + 1)^{-1}\}.$$

In general there exist functions $F_k(m, n)$ and $G_k(m, n)$ such that

$$(4.3) \quad R_k(m, n) = A_m + B_n - \max \{F_k(m, n), G_k(m, n)\}.$$

These functions may be defined recursively as follows.

$$(4.4) \quad F_0(m, n) = G_0(m, n) = 0$$

and for $k \geq 0$

$$(4.5) \quad F_{k+1}(m, n) = A_m(\tau_1 + m + 1)^{-1} + E(\max \{F_k(m + 1, n), G_k(m + 1, n)\})$$

$$(4.6) \quad G_{k+1}(m, n) = B_n(\tau_2 + n + 1)^{-1} + E(\max \{F_k(m, n + 1), G_k(m, n + 1)\}).$$

Once these functions have been determined, the optimal policy is to select experiment ε_1 provided $F_k(m, n) \geq G_k(m, n)$ and experiment ε_2 otherwise. In order to gain more information about the optimal policy let $D_k(m, n) = F_k(m, n) - G_k(m, n)$, the relative advantage of ε_1 over ε_2 . These functions may be defined recursively in the following manner.

$$(4.7) \quad D_1(m, n) = A_m(\tau_1 + m + 1)^{-1} + B_n(\tau_2 + n + 1)^{-1}.$$

Let $H_k(m, n) = E(\max \{F_k(m, n), G_k(m, n)\})$, then from (4.5) and (4.6) it follows that

$$(4.8) \quad D_{k+1}(m, n) = D_1(m, n) + H_k(m + 1, n) - H_k(m, n + 1).$$

Since $\max \{F, G\} = G + (F - G)^+$ where x^+ denotes $\max \{x, 0\}$,

$$(4.9) \quad H_k(m + 1, n) = E(G_k(m + 1, n)) + E(D_k^+(m + 1, n)).$$

Since $\max \{F, G\} = F + (G - F)^+ = F - (F - G)^-$ where x^- denotes $\min \{x, 0\}$,

$$(4.10) \quad H_k(m, n + 1) = E(F_k(m, n + 1)) - E(D_k^-(m, n + 1)).$$

Now using (4.5) and (4.6) again we find that

$$(4.11) \quad E(G_k(m + 1, n)) = B_n(\tau_2 + n + 1)^{-1} + H_k(m + 1, n + 1),$$

and also that

$$(4.12) \quad E(F_k(m, n + 1)) = A_m(\tau_1 + m + 1)^{-1} + H_k(m + 1, n + 1).$$

Thus $E(G_k(m + 1, n) - F_k(m, n + 1)) = -D_1(m, n)$ independent of k , so that for $k \geq 1$

$$(4.13) \quad D_{k+1}(m, n) = E(D_k^+(m + 1, n)) + E(D_k^-(m, n + 1)).$$

Similar recursive relations have been derived by the author [5] for the Bernoulli two-armed bandit problem. These relations are now used to prove the first result about the optimal procedure.

THEOREM 4.1. *For any positive constant c ,*

$$D_k(m, n, cA_m, cB_n) = cD_k(m, n, A_m, B_n).$$

PROOF. By not suppressing the arguments A_m, B_n the proof is by induction on k and follows directly from (4.13). \square

Since the optimal strategy depends on the sign of $D_k(m, n)$, the decision rule is a function of (A_m, B_n) only through the ratio A_m/B_n . This means that a new function $\Delta_k(m, n, r)$ can be defined as follows.

$$(4.14) \quad \Delta_k(m, n, A_m/B_n) = B_n D_k(m, n, A_m, B_n).$$

LEMMA 4.1. *The functions $\Delta_k(m, n, r)$ are defined recursively through the following equations.*

$$(4.15) \quad \Delta_1(m, n, r) = r(\tau_1 + m + 1)^{-1} - (\tau_2 + n + 1)^{-1}$$

and for $k = 1, \dots, N - 1$,

$$(4.16) \quad \Delta_{k+1}(m, n, r) = E(\Delta_k^+(m + 1, n, a_m U^{-1}r)) \\ + E(b_n V^{-1} \Delta_k^-(m, n + 1, b_n^{-1} V r))$$

where U and V have independent beta distributions $B(\alpha_1 + \frac{1}{2}m, \frac{1}{2})$ and $B(\alpha_2 + \frac{1}{2}n, \frac{1}{2})$ respectively and a_m and b_n are given in (3.3) and (3.4).

PROOF. This follows directly from (4.7), (4.13), and Lemma 3.1. \square

LEMMA 4.2. For the functions $\Delta_k(m, n, r)$ defined in Lemma 4.1, there exist two sequences $Q_k(m)$ and $P_k(n)$ such that $Q_k(m) > 0$ and $P_k(n) < 0$ and for all $r \geq 0$, $P_k(n) < \Delta_k(m, n, r) < Q_k(m)r$.

PROOF. The proof is by induction on k and uses Lemma 4.1. \square

The main use of this lemma is in the proof of the following theorem giving a characterization of the optimal sequential decision rule.

THEOREM 4.2. For the functions $\Delta_k(m, n, r)$ defined in Lemma 4.1, there exists a unique sequence of constants $\gamma_k(m, n)$ such that $\Delta_k(m, n, r) \geq 0$ if and only if $r \geq \gamma_k(m, n)$.

PROOF. Using Lemma 4.2, the following properties of $\Delta_k(m, n, r)$ may be established by induction:

- (i) $\Delta_k(m, n, r)$ is a strictly increasing function of r ,
- (ii) $\Delta_k(m, n, r)$ is a continuous function of r ,
- (iii) $\Delta_k(m, n, r)$ is negative for sufficiently small values of r , and
- (iv) $\Delta_k(m, n, r)$ is positive and arbitrarily large for sufficiently large values of r .

Then the existence of $\gamma_k(m, n)$ is guaranteed by the intermediate value theorem and the uniqueness of $\gamma_k(m, n)$ is guaranteed by property (i). \square

The optimal sequential procedure is then of the following form. If m trials have been allocated to ε_1 , n trials have been allocated to ε_2 , and there are k trials remaining, then on the basis of the prior parameters and the posterior variances A_m, B_n it is optimal to allocate the next trial to ε_1 if and only if A_m/B_n exceeds $\gamma_k(m, n)$.

5. The myopic decision rule. Unfortunately, little is known about the constants $\gamma_k(m, n)$. In approximating the optimal decision rule, one needs only to choose a set of constants $g_k(m, n)$ "close" to $\gamma_k(m, n)$. An appealing choice is

$$(5.1) \quad g_k(m, n) = \gamma_1(m, n) = (\tau_1 + m + 1)(\tau_2 + n + 1)^{-1},$$

and the resulting decision rule, which is independent of k , is called the myopic rule. It "acts" as if there were always but one more trial remaining.

This decision rule also appears when one obtains the limiting form of the optimal strategy as more and more information is assumed to be available concerning the precisions P_1 and P_2 . Consider prior distributions in the normal-gamma family where the marginals of P_i ($i = 1, 2$) are gamma distributions with $\beta_i = \sigma_i^2 \alpha_i$. Suppose σ_i^2 is a constant and α_i is allowed to increase to $+\infty$. These marginals approach singular distributions with the entire probability mass concentrated at σ_i^{-2} . The prior marginal distributions of μ_i approach normal distributions with mean η_i and precision $\tau_i \sigma_i^{-2}$. Note also that as $\alpha_i \rightarrow +\infty$, $a_m \rightarrow (\tau_1 + m)(\tau_1 + m + 1)^{-1}$ and $b_n \rightarrow (\tau_2 + n)(\tau_2 + n + 1)^{-1}$. The random variables A_m, B_n, U and V all approach singular distributions with probability mass concentrated at $\sigma_1^2(\tau_1 + m)^{-1}, \sigma_2^2(\tau_2 + n), 1$ and 1 respectively. The limiting form

of $\Delta_k(m, n, r)$, from the Helly-Bray theorem, satisfies the recursive equations.

$$(5.2) \quad \Delta_1(m, n, r) = r(\tau_1 + m + 1)^{-1} - (\tau_2 + n + 1)^{-1},$$

and for $k = 1, \dots, N - 1$

$$(5.3) \quad \Delta_{k+1}(m, n, r) = \Delta_k^+(m + 1, n, a_m r) + b_n \Delta_k^-(m, n + 1, b_n^{-1} r).$$

THEOREM 5.1. *For the functions $\Delta_k(m, n, r)$ defined in (5.2) and (5.3), $r > \gamma_1(m, n) \Rightarrow \Delta_k(m, n, r) \geq 0$.*

PROOF. The proof is by induction on k . \square

So the limiting optimal decision rule is to allocate the next trial to ϵ_1 provided

$$(5.4) \quad \sigma_1^2 \sigma_2^{-2} \geq (\tau_1 + m)(\tau_1 + m + 1)(\tau_2 + n)^{-1}(\tau_2 + n + 1)^{-1}.$$

Since this decision rule does not depend on the outcomes of the various trials, the sampling may be done in one stage, choosing as the total number of trials to be allocated to ϵ_1 the largest value of m such that (5.9) is satisfied.

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