

## BAYES SEQUENTIAL ESTIMATION OF A POISSON RATE: A DISCRETE TIME APPROACH<sup>1</sup>

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This paper provides explicit solutions to the problem of estimating the arrival rate  $\lambda$  of a Poisson process using a Bayes sequential approach. The loss associated with estimating  $\lambda$  by  $d$  is assumed to be of the form  $(\lambda - d)^2\lambda^{-p}$  and the cost of observation includes both a time cost and an event cost. A discrete time approach in which decisions are made at the end of time intervals having length  $t$ . Limits of the procedures as  $t$  approaches zero are discussed and related to the continuous time Bayes sequential procedure.

**1. Introduction.** Suppose that one observes a continuous-time Poisson process in order to estimate its arrival rate  $\lambda$  using a Bayes sequential approach. The observation cost is assumed to be  $c_1$  per unit time and  $c_2$  per event observed where  $c_1$  and  $c_2$  are nonnegative constants. The loss associated with estimating  $\lambda$  by  $d$  is assumed to be of the form  $L(\lambda, d) = (\lambda - d)^2\lambda^{-p}$  where we will discuss values of  $p$  in the interval  $0 < p < 3$ . Thus the total loss resulting from estimating  $\lambda$  by  $d$  after having observed the process for time  $t$  during which  $X_t$  events occurred is  $L(\lambda, d) + c_1t + c_2X_t$ . Prior information about  $\lambda$  is assumed to be represented by a gamma distribution.

In this paper a discrete time approach is considered in which the process is observed continuously but decisions are made at the end of time intervals having length  $t$ . The optimal Bayes sequential decision procedure for estimating  $\lambda$  is determined explicitly. Furthermore, the solutions will be presented in a simple form which makes them appealing for practical applications. This approach contrasts with one taken by Shapiro and Wardrop [6] in which decisions can only be made at the time an event occurs. El-Sayyad and Freeman [4] have also attacked the problem using both approaches and various loss and cost functions. Shapiro and Wardrop [7] have also solved the problem in continuous time using the notion of "monotone case" for continuous time problems and employing Dynkin's identity.

If the prior gamma distribution over  $\lambda$  has density function  $g(\lambda) = \Gamma(\alpha)^{-1}\beta^\alpha\lambda^{\alpha-1}e^{-\beta\lambda}$  and one observes  $X_t$  events in time  $t$ , the posterior distribution at time  $t$  will also be a gamma distribution with parameters  $(\alpha + X_t, \beta + t)$ . It will be convenient then to represent the results of experimentation by a plot of the posterior parameters. The optimal stopping region will be determined using backward induction; see, e.g., DeGroot (1970) or Chow, Robbins and Siegmund (1971).

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**2. Observation cost proportional to observation time ( $c_2 = 0$ ).** Define  $\rho_0(\alpha, \beta)$  as the risk, or minimum expected loss, of estimating  $\lambda$  without any observation of the process when  $(\alpha, \beta)$  are the parameters of the prior distribution. Define  $\rho_i(\alpha, \beta, t)$  as the risk, or minimum expected total loss, of the optimal procedure when  $i$  sampling intervals of length  $t$  are available. Also define  $\rho^*(\alpha, \beta, t)$  as the risk of the optimal procedure in the infinite horizon case. Then

$$(2.1) \quad \begin{aligned} \rho_1(\alpha, \beta, t) &= \min\{\rho_0(\alpha, \beta), E[\rho_0(\alpha + X_t, \beta + t)] + c_1 t\} && \text{and} \\ \rho_i(\alpha, \beta, t) &= \min\{\rho_0(\alpha, \beta), E[\rho_{i-1}(\alpha + X_t, \beta + t, t)] + c_1 t\} \end{aligned}$$

where the expected value is taken with respect to the marginal distribution of  $X_t$ . The functions  $\rho_i$  determine the optimal procedure in a finite horizon problem. The role they play here will be their use in determining the nature of the function  $\rho^*(\alpha, \beta, t)$ . Since  $\rho^*(\alpha, \beta, t) = \min\{\rho_0(\alpha, \beta), E[\rho^*(\alpha + X_t, \beta + t, t)] + c_1 t\}$ , the optimal stopping region in the  $(\alpha, \beta)$  plane is  $\{(\alpha, \beta) | \rho^*(\alpha, \beta, t) = \rho_0(\alpha, \beta)\}$  and the continuation region is  $\{(\alpha, \beta) | \rho^*(\alpha, \beta, t) < \rho_0(\alpha, \beta)\}$ .

The stopping risk  $\rho_0(\alpha, \beta)$  associated with the Bayes estimate  $d = \beta^{-1}(\alpha - p)$  is given by  $\rho_0(\alpha, \beta) = \beta^{p-2}\Gamma(\alpha - p + 1)/\Gamma(\alpha)$ , where we assume  $\alpha > p$ . By (2.1),  $\rho_1(\alpha, \beta, t) = \min\{\rho_0(\alpha, \beta), \rho_0(\alpha, \beta) - [t\beta^{p-2}\Gamma(\alpha - p + 1)/(\beta + t)\Gamma(\alpha) - c_1 t]\}$ . Thus, the optimal procedure when at most one sampling period is available for observation is characterized by the following stopping region  $R_{s,t}$  and continuation region  $R_{c,t}$ :

$$(2.2) \quad \begin{aligned} R_{s,t} &= \{(\alpha, \beta) | \Gamma(\alpha)/\Gamma(\alpha - p + 1) \geq \beta^{p-2}/c_1(\beta + t), \alpha > p, \beta > 0\} \\ R_{c,t} &= \{(\alpha, \beta) | \Gamma(\alpha)/\Gamma(\alpha - p + 1) < \beta^{p-2}/c_1(\beta + t), \alpha > p, \beta > 0\}. \end{aligned}$$

**3. Characterization of the optimal stopping region for  $1 \leq p < 3$ ;  $c_2 = 0$ .** Note that for  $1 \leq p \leq 2$ ,  $\Gamma(\alpha)/\Gamma(\alpha - p + 1)$  is nondecreasing in  $\alpha$  and  $\beta^{p-2}/c_1(\beta + t)$  is decreasing in  $\beta$ . This is also true for  $2 < p < 3$  if  $\beta > t(p - 2)/(3 - p)$ . In either case once the point  $(\alpha, \beta)$  enters  $R_{s,t}$ , the posterior parameters  $(\alpha + X_t, \beta + t)$  must remain there. Consider any point  $(\alpha, \beta) \in R_{s,t}$ . Then  $\rho_1(\alpha, \beta, t) = \rho_0(\alpha, \beta)$  and  $\rho_1(\alpha + X_t, \beta + t, t) = \rho_0(\alpha + X_t, \beta + t)$  for all  $t$  and  $X_t$ . Hence it follows from (2.1) by induction that  $\rho_n(\alpha, \beta, t) = \rho_0(\alpha, \beta)$  for any  $n$  and  $(\alpha, \beta) \in R_{s,t}$ . Furthermore, it follows from [3], page 296, that  $\lim_{n \rightarrow \infty} \rho_n(\alpha, \beta, t) = \rho^*(\alpha, \beta, t)$ . Hence,  $\rho^*(\alpha, \beta, t) = \rho_0(\alpha, \beta)$  for  $(\alpha, \beta) \in R_{s,t}$  and  $\rho^*(\alpha, \beta, t) < \rho_0(\alpha, \beta)$  in  $R_{c,t}$ . This proves that  $R_{s,t}$  and  $R_{c,t}$  are the optimal stopping and continuation regions for the infinite horizon problem when  $1 \leq p < 3$ .

One can partition the continuation region  $R_{c,t}$  into a number of disjoint regions such that in each region the optimal procedure is explicitly described. The  $k$ th region is defined as:

$$(3.1) \quad \begin{aligned} R_k &= \{(\alpha, \beta) | (\beta + kt)^{p-2}/c_1[\beta + (k + 1)t] \leq \Gamma(\alpha)/\Gamma(\alpha - p + 1) \\ &< [\beta + (k - 1)t]^{p-2}/c_1(\beta + kt), \alpha > p, \beta > 0\}. \end{aligned}$$

The optimal procedure in the  $k$ th region is to observe the process for at least one

more period. No more than  $k$  periods will ever be needed, however, since the posterior parameters must lie in  $R_{s,t}$  after  $k$  further sampling periods. Since the prior parameters must belong to one of the above regions for some  $k = k_0 < \infty$ , there is an upper bound  $k_0 t$  on the total observation time.

Note that the optimal procedure was derived by realizing that for certain values of  $p$  monotonicity arises and the myopic rule is optimal. When  $c_2 = 0$  we only find this to be true for  $1 < p < 3$ , and for the case in which  $2 < p < 3$  we also need to choose  $t$  such that the prior parameter  $\beta$  is greater than  $t(p - 2)/(3 - p)$ . If this condition is not met, one can partition  $R_{s,t}$  into  $R_1 \cup R_2 \cup R_3$  where  $R_1 = \{(\alpha, \beta) | \Gamma(\alpha)/\Gamma(\alpha - p + 1) \geq \beta^{p-3}/c_1, \alpha > p, \beta > 0\}$ ,  $R_2 = \{(\alpha, \beta) | \beta^{p-2}/c_1(\beta + t) \leq \Gamma(\alpha)/\Gamma(\alpha - p + 1) < \beta^{p-3}/c_1, \alpha > p, \beta > t(p - 2)/(3 - p)\}$ , and  $R_3 = \{(\alpha, \beta) | \beta^{p-2}/c_1(\beta + t) \leq \Gamma(\alpha)/\Gamma(\alpha - p + 1) < \beta^{p-3}/c_1, \alpha > p, \beta \leq t(p - 2)/(3 - p)\}$ . In  $R_1 \cup R_2$  it is optimal to stop sampling and in  $R_{c,t}$  one should continue. The optimal procedure in  $R_3$  is unknown, however.

Note that when  $p = 1$  the optimal stopping region becomes

$$R_{s,t} = \left\{ (\alpha, \beta) \mid \beta \geq \frac{1}{2}(t^2 + 4/c_1)^{\frac{1}{2}} - \frac{t}{2}, \alpha > 1 \right\}.$$

If  $(\alpha_0, \beta_0)$  are the parameters of the prior distribution, the optimal procedure is a fixed time procedure in which one samples for exactly  $\tau = \max \{0, \lfloor \frac{1}{2}(t^2 + 4/c_1)^{\frac{1}{2}} + \frac{t}{2} - \beta_0 \rfloor\}$  units of time, where  $\lfloor \cdot \rfloor$  is the greatest integer function. Note when  $t = 1$  (ordinary Poisson sampling), that the optimal procedure is equivalent to taking a sequential random sample of exactly  $\tau = \max \{0, \lfloor \frac{1}{2}(1 + 4/c_1) + \frac{1}{2} - \beta_0 \rfloor\}$  Poisson random variables.

When  $p = 2$  the optimal stopping region becomes

$$R_{s,t} = \left\{ (\alpha, \beta) \mid \beta \geq \frac{1}{c_1(\alpha - 1)} - t, \alpha > 2, \beta > 0 \right\}.$$

The simple form of the boundary is appealing for practical applications.

**4. The optimal procedure for  $c_2 > 0$ .** The cost function for observation time  $t$  has the form  $c(X_t) = c_1 t + c_2 X_t$ . Thus  $\rho_1(\alpha, \beta, t) = \min\{\rho_0(\alpha, \beta), E[\rho_0(\alpha + X_t, \beta + t) + c(X_t)]\} = \min\{\rho_0(\alpha, \beta), \rho_0(\alpha, \beta) - [t\beta^{p-2}\Gamma(\alpha - p + 1)/(\beta + t)\Gamma(\alpha) - c_1 t - c_2 t\alpha/\beta]\}$ . Hence the optimal procedure when at most one sampling period is available is characterized by the following stopping region  $\mathcal{C}_{s,t}$  and continuation region  $\mathcal{C}_{c,t}$ :

$$\mathcal{C}_{s,t} = \left\{ (\alpha, \beta) \mid 1 - \frac{c_1 \Gamma(\alpha)(\beta + t)}{\Gamma(\alpha - p + 1)\beta^{p-2}} - \frac{c_2 \Gamma(\alpha + 1)(\beta + t)}{\Gamma(\alpha - p + 1)\beta^{p-1}} < 0, \alpha > p, \beta > 0 \right\} \tag{4.1}$$

$$\mathcal{C}_{c,t} = \left\{ (\alpha, \beta) \mid 1 - \frac{c_1 \Gamma(\alpha)(\beta + t)}{\Gamma(\alpha - p + 1)\beta^{p-2}} - \frac{c_2 \Gamma(\alpha + 1)(\beta + t)}{\Gamma(\alpha - p + 1)\beta^{p-1}} > 0, \alpha > p, \beta > 0 \right\}.$$

Note that for  $1 < p < 2$  and  $\beta > t(p - 1)/(2 - p)$  the sum

$$\frac{c_1\Gamma(\alpha)(\beta + t)}{\Gamma(\alpha - p + 1)\beta^{p-2}} + \frac{c_2\Gamma(\alpha + 1)(\beta + t)}{\Gamma(\alpha - p + 1)\beta^{p-1}}$$

is nondecreasing in  $\alpha$  and  $\beta$ . It follows that once  $\mathcal{C}_{s,t}$  is entered by posterior parameters it cannot be escaped. By the same arguments given for the case in which  $c_2 = 0$ ,  $\mathcal{C}_{s,t}$  is the optimal stopping region and  $\mathcal{C}_{c,t}$  the optimal continuation region when  $1 < p < 2$  and  $\beta > t(p - 1)/(2 - p)$ .

If we let  $c_1 = 0$  the above arguments hold additionally for  $0 \leq p \leq 1$  and all  $t$ . This follows by noting that

$$\frac{c_2\Gamma(\alpha + 1)(\beta + t)}{\Gamma(\alpha - p + 1)\beta^{p-1}}$$

is increasing in  $\alpha$  and  $\beta$  for  $0 \leq p \leq 1$ . The optimal stopping and continuation regions in this case are

$$(4.2) \quad \begin{aligned} \mathcal{D}_{s,t} &= \left\{ (\alpha, \beta) \mid 1 - \frac{c_2\Gamma(\alpha + 1)(\beta + t)}{\Gamma(\alpha - p + 1)\beta^{p-1}} \leq 0, \alpha > p, \beta > 0 \right\} \quad \text{and} \\ \mathcal{D}_{c,t} &= \left\{ (\alpha, \beta) \mid 1 - \frac{c_2\Gamma(\alpha + 1)(\beta + t)}{\Gamma(\alpha - p + 1)\beta^{p-1}} > 0, \alpha > p, \beta > 0 \right\}. \end{aligned}$$

**5. Convergence to continuous time.** One can use the discrete time approach as an approximation to a continuous time problem, the approximation improving as the length  $t$  of the interval is suitably decreased. In viewing continuous time as a limiting version of the discrete time problem as  $t \rightarrow 0$ , one obtains the stopping regions

$$(5.1) \quad \begin{aligned} \mathcal{R}_s &= \lim_{t \rightarrow 0} \mathcal{R}_{s,t} = \{(\alpha, \beta) \mid \Gamma(\alpha)/\Gamma(\alpha - p + 1) \\ &\geq \beta^{p-3}/c_1, \alpha > p, \beta > 0\}, \\ \mathcal{C}_s &= \lim_{t \rightarrow 0} \mathcal{C}_{s,t} = \{(\alpha, \beta) \mid 1 - c_1\beta^{3-p}\Gamma(\alpha)/\Gamma(\alpha - p + 1) \\ &- c_2\beta^{2-p}\Gamma(\alpha + 1)/\Gamma(\alpha - p + 1) \leq 0, \alpha > p, \beta > 0\}, \\ \mathcal{D}_s &= \lim_{t \rightarrow 0} \mathcal{D}_{s,t} = \{(\alpha, \beta) \mid 1 - c_2\beta^{2-p}\Gamma(\alpha + 1)/\Gamma(\alpha - p + 1) \\ &\leq 0, \alpha > p, \beta > 0\}, \end{aligned}$$

which are the corresponding optimum boundaries derived by Shapiro and Wardrop in their continuous time approach. It can be shown that as  $t \rightarrow 0$  the Bayes risk of the optimum discrete time procedure converges to the risk of the continuous time Bayes sequential procedure, a result needed to rigorously prove that the limit of the optimal discrete time solution yields the solution to the limiting problem.

When  $p = 3$  and  $c_2 = 0$ , or when  $p = 2$  and  $c_2 > 0$ , an exact solution to the discrete time problem was not obtainable. However, one can identify in each case a stopping region, a continuation region, and a region separating the above two in which the optimal procedure is unknown. It should be noted that as the length  $t$  of

the sampling interval decreases, the size of the region within which the procedure is unknown decreases. Taking the limit as  $t \rightarrow 0$  this region vanishes and one again obtains the optimum boundaries for the continuous time problem given by Shapiro and Wardrop.

When  $p = 1$  and  $c_2 = 0$  the optimal procedure is again a fixed time procedure with optimum stopping region

$$R_s = \{(\alpha, \beta) | \beta \geq (c_1)^{-\frac{1}{2}}, \alpha > 1\}.$$

If  $(\alpha_0, \beta_0)$  is the prior parameter point, the optimal procedure in this case is to sample for exactly  $\tau = \max\{0, (c_1)^{-\frac{1}{2}} - \beta_0\}$  units of time.

In contrast, the optimal procedure when  $p = 3$  and  $c_2 = 0$  is an "inverse" sampling scheme with optimum stopping region.

$$R_s = \{(\alpha, \beta) | (\alpha - 1)(\alpha - 2) \geq 1/c_1, \alpha > 3, \beta > 0\}.$$

Hence, the optimal procedure in this case is to sample until exactly  $k$  events occur, where  $k$  is the smallest nonnegative integer such that  $(\alpha_0 + k - 1)(\alpha_0 + k - 2) \geq 1/c_1$ .

When  $p = 2$  the optimal procedure is characterized by the stopping region

$$R_s = \left\{ (\alpha, \beta) \mid \beta \geq \frac{1}{c_1(\alpha - 1)}, \alpha > 2, \beta > 0 \right\}.$$

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#### REFERENCES

- [1] CHOW, Y. S., ROBBINS, H. and SIEGMUND, D. (1971). *Great Expectations: The Theory of Optimal Stopping*. Houghton Mifflin, Boston.
- [2] COX, D. R. and HINKLEY, D. V. (1974). *Theoretical Statistics*. Chapman and Hall, London.
- [3] DEGROOT, M. H. (1970). *Optimal Statistical Decisions*. McGraw-Hill, New York
- [4] EL-SAYYAD, G. M. and FREEMAN, P. R. (1973). Bayesian sequential estimation of a Poisson rate. *Biometrika* **60** 289-296.
- [5] NOVIC, B. (1977). Bayes sequential estimation of a Poisson rate. Technical Report #134, Department of Statistics, Carnegie-Mellon Univ.
- [6] SHAPIRO, C. P. and WARDROP, R. L. (1977a). Large sample properties of the Bayes' sequential procedure for estimating the arrival rate of a Poisson process with invariant loss. Technical Report, Department of Statistics, Univ. Wisconsin.
- [7] SHAPIRO, C. P. and WARDROP, R. L. (1977b). Dynkin's Identity applied to Bayes' sequential estimation of a Poisson process. Technical Report, Department of Statistics, Univ. Wisconsin.

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