Markov-modulated (marked) Poisson processes

Jan-Ole Koslik

The functionalities of Lcpp can also be called to fit so-called Markov-modulated Poisson processes. These are doubly stochastic Poisson point processes where the intensity is directed by an underlying continuous-time Markov chain. Such processes are useful for modelling arrival times, for example of calls in a call center, or patients in the hospital.

Example 1: Markov-modulated Poisson processes

Setting parameters

We choose to have a considerably higher rate and shorter stays of the underlying Markov chain in state 2, i.e. state 2 is **bursty**.

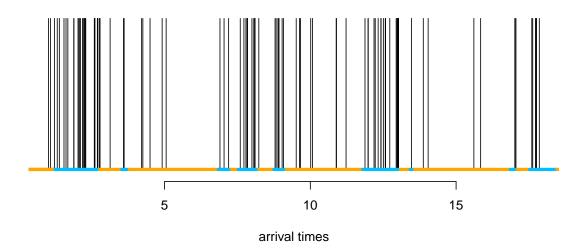
```
# state-dependent rates
lambda = c(2, 15)
# generator matrix of the underlying Markov chain
Q = matrix(c(-0.5,0.5,2,-2), nrow = 2, byrow = TRUE)
```

Simulating an MMPP

```
set.seed(123)
k = 200 # number of state switches
trans_times = s = rep(NA, k) # time points where the chain transitions
s[1] = sample(1:2, 1) \# initial \ distribution \ c(0.5, 0.5)
# exponentially distributed waiting times
trans_times[1] = rexp(1, -Q[s[1],s[1]])
# in a fixed interval, the number of arrivals is Pois(lambda * interval_length)
n_arrivals = rpois(1, lambda[s[1]]*trans_times[1])
# arrival times within fixed interval are uniformly distributed
arrival_times = runif(n_arrivals, 0, trans_times[1])
for(t in 2:k){
  s[t] = c(1,2)[-s[t-1]]
  # exponentially distributed waiting times
  trans_times[t] = trans_times[t-1] + rexp(1, -Q[s[t], s[t]])
  # in a fixed interval, the number of arrivals is Pois(lambda * interval_length)
  n_arrivals = rpois(1, lambda[s[t]]*(trans_times[t]-trans_times[t-1]))
  # arrival times within fixed interval are uniformly distributed
  arrival_times = c(arrival_times,
                    runif(n_arrivals, trans_times[t-1], trans_times[t]))
}
arrival_times = sort(arrival_times)
```

Let's visualize the simulated MMPP

```
--- state 1
--- state 2
```



What makes the MMPP special compared to a regular Poisson point process is its **burstiness** when the Markov chain is in the second state.

Writing the negative log-likelihood function

The likelihood of a stationary MMPP for waiting times x_1, \ldots, x_n is (Meier-Hellstern (1987), Langrock, Borchers, and Skaug (2013))

$$L(\theta) = \delta \left(\prod_{i=1}^{n} \exp((Q - \Lambda)x_i)\Lambda \right) 1,$$

where Q is the generator matrix of the continuous-time Markov chain, Λ is a diagonal matrix of state-dependent Poisson intensities, δ is the stationary distribution of the continuous-time Markov chain, and 1 is a column vector of ones. We can easily calculate the log of the above expression using the standard implementation of the general forward algorithm forward_g() when choosing the first matrix of state-dependent densities to be the identity (i.e.) the first row of the allprobs matrix to be one and all other matrices of state-dependent density matrices to be Λ .

```
mllk = function(theta.star, timediff, N=2){
  lambda = exp(theta.star[1:N]) # state specific rates
  Q = diag(N) # generator matrix
  Q[!Q] = exp(theta.star[N+1:(N*(N-1))])
  diag(Q) = 0
  diag(Q) = -rowSums(Q)
  Qube = Lcpp::tpm_cont(Q-diag(lambda), timediff) # exp((Q-Lambda)*deltat)
  allprobs = matrix(lambda, nrow = length(timediff+1), ncol = N, byrow = T)
  allprobs[1,] = 1
  delta = solve(t(Q+1), rep(1,N), tol = 1e-20)
  -Lcpp::forward_g(delta, Qube, allprobs)
}
```

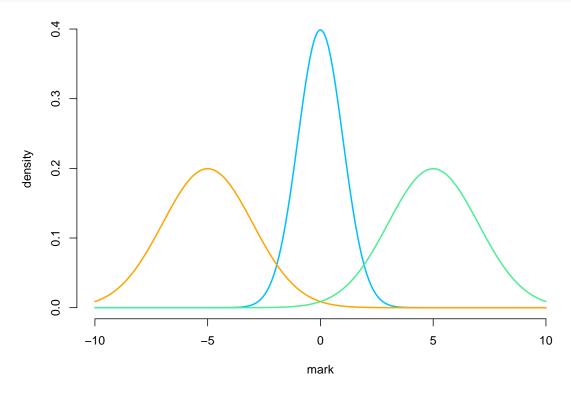
Fitting an MMPP to the data

Results

```
exp(mod$estimate)
#> [1] 1.9496891 15.0831215 1.8998849 0.4003955
```

Example 2: Markov-modulated marked Poisson processes

Such processes can also carry additional information, so called **marks**, at every arrival time when we also observe the realization of a different random variable that only depends on the underlying states of the continuous-time Markov chain. For example for patient arrivals in the hospital we could observe a biomarker at every arrival time. Information on the underlying health status is then present in both, the arrival times (because sick patients visit more often) and the biomarkers.



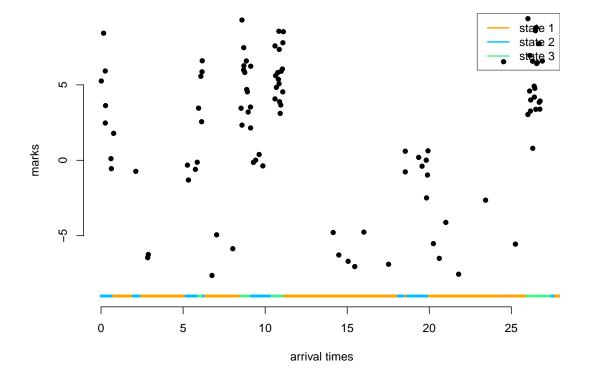
Simulating an MMMPP

We now show how to simulate an MMMPP and additionally how to generalize to more than two hidden states.

```
set.seed(123)
k = 200 # number of state switches
trans_times = s = rep(NA, k) # time points where the chain transitions
s[1] = sample(1:3, 1) # initial distribuion uniformly
# exponentially distributed waiting times
trans_times[1] = rexp(1, -Q[s[1], s[1]])
# in a fixed interval, the number of arrivals is Pois(lambda * interval_length)
n_arrivals = rpois(1, lambda[s[1]]*trans_times[1])
# arrival times within fixed interval are uniformly distributed
arrival_times = runif(n_arrivals, 0, trans_times[1])
# marks are iid in interval, given underlying state
marks = rnorm(n_arrivals, mu[s[1]], sigma[s[1]])
for(t in 2:k){
  # off-diagonal elements of the s[t-1] row of Q divided by the diagonal element
  # give the probabilites of the next state
  s[t] = sample(c(1:3)[-s[t-1]], 1, prob = Q[s[t-1],-s[t-1]]/-Q[s[t-1],s[t-1]])
  # exponentially distributed waiting times
```

Let's visualize the simulated MMMPP

```
n = length(arrival_times)
plot(arrival_times[1:100], marks[1:100], pch = 16, bty = "n",
        ylim = c(-9,9), xlab = "arrival times", ylab = "marks")
segments(x0 = c(0,trans_times[1:98]), x1 = trans_times[1:99],
        y0 = rep(-9,100), y1 = rep(-9,100), col = color[s[1:49]], lwd = 4)
legend("topright", lwd = 2, col = color,
        legend = c("state 1", "state 2", "state 3"), box.lwd = 0)
```



Writing the negative log-likelihood function

The likelihood of a stationary MMMPP for waiting times x_1, \ldots, x_n and marks y_0, y_1, \ldots, y_n only changes slightly from the MMPP likelihood, as we include the matrix of state-specific densities (Lu (2012), Mews et al. (2023)):

$$L(\theta) = \delta P(y_0) \Big(\prod_{i=1}^n \exp((Q - \Lambda)x_i) \Lambda P(y_i) \Big) 1,$$

where Q, Λ and δ are as above and $P(y_i)$ is a diagonal matrix with state-dependent densites for the observation at time t_i . We can again easily calculate the log of the above expression using the standard implementation of the general forward algorithm forward_g() when first calculating the allprobs matrix with state-dependent densities for the marks (as usual for HMMs) and then multiplying each row except the first one element-wise with the state-dependent rates.

```
mllk = function(theta.star, y, timediff, N){
  lambda = exp(theta.star[1:N]) # state specific rates
  mu = theta.star[N+1:N]
  sigma = exp(theta.star[2*N+1:N])
  Q = diag(N) # generator matrix
  Q[!Q] = \exp(\text{theta.star}[3*N+1:(N*(N-1))])
  diag(Q) = 0
  diag(Q) = -rowSums(Q)
  delta = solve(t(Q+1), rep(1,N), tol = 1e-20)
  Qube = Lcpp::tpm_cont(Q-diag(lambda), timediff) # exp((Q-Lambda)*deltat)
  allprobs = matrix(1, length(y), N)
  for(j in 1:N){
    allprobs[,j] = dnorm(y, mu[j], sigma[j])
  allprobs[-1,] = allprobs[-1,] * matrix(lambda, length(y)-1, N, byrow = T)
  -Lcpp::forward_g(delta, Qube, allprobs)
}
```

Fitting an MMPP to the data

Results

```
N = 3
(lambda = exp(mod2$estimate[1:N]))
#> [1]     0.9646715     4.8640549     19.5008964
(mu = mod2$estimate[N+1:N])
#> [1]     -5.18540506     -0.09097056     4.80538783
(sigma = exp(mod2$estimate[2*N+1:N]))
#> [1]     1.7931063     0.9644485     2.0093266
Q = diag(N)
Q[!Q] = exp(mod2$estimate[3*N+1:(N*(N-1))])
diag(Q) = 0
diag(Q) = -rowSums(Q)
Q
```

```
#> [,1] [,2] [,3]

#> [1,] -0.5909290  0.2788642  0.3120648

#> [2,]  0.9258937 -1.1798376  0.2539439

#> [3,]  1.1933693  1.2097186 -2.4030879
```

References

- Langrock, Roland, David L Borchers, and Hans J Skaug. 2013. "Markov-Modulated Nonhomogeneous Poisson Processes for Modeling Detections in Surveys of Marine Mammal Abundance." *Journal of the American Statistical Association* 108 (503): 840–51.
- Lu, Shaochuan. 2012. "Markov Modulated Poisson Process Associated with State-Dependent Marks and Its Applications to the Deep Earthquakes." Annals of the Institute of Statistical Mathematics 64: 87–106.
- Meier-Hellstern, Kathleen S. 1987. "A Fitting Algorithm for Markov-Modulated Poisson Processes Having Two Arrival Rates." European Journal of Operational Research 29 (3): 370–77.
- Mews, Sina, Bastian Surmann, Lena Hasemann, and Svenja Elkenkamp. 2023. "Markov-Modulated Marked Poisson Processes for Modeling Disease Dynamics Based on Medical Claims Data." *Statistics in Medicine*.