MA 568 Statistical Analysis of Point Process Data Problem set 1

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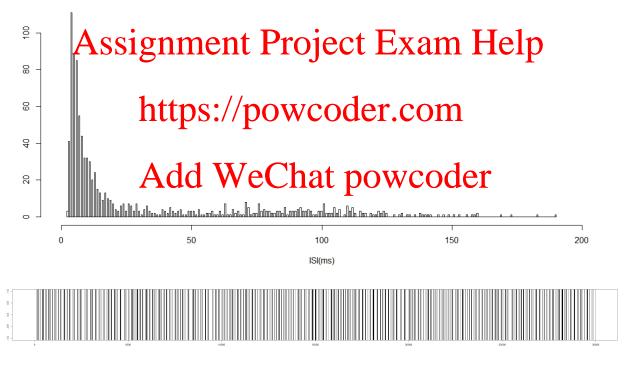
Download the **Retinal_ISIs.mat** file from the course website. This file contains the waiting times, or interspike intervals (ISIs), in milliseconds, of a single retinal ganglion cell over 30 seconds.

```
library(R.matlab)
data <- readMat("C:\\Users\\Long_Tao\\Downloads\\Retinal_ISIs(1).mat")
ISIs <- data$ISIs
hist (ISIs, breaks = 300, xlab='ISI(ms)',xlim=c(0,200), ylab='')

spiketimes <- cumsum (ISIs)
T = max(spiketimes)
plot(1:T, rep(0,T),type='n',xlab='',ylab='')
abline(v=spiketimes)</pre>
```

First we plot the spiking activity as a histogram of the distribution of the times between spiking events and a spike train time series.

Histogram of ISIs

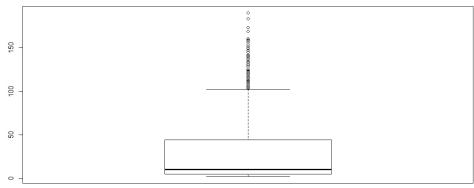


The histogram and spike train time series suggest that this neuron fires most of its spikes with an ISI between 5-40 ms. Features to note include an initial refractory period under 5 ms, a large number of spikes with ISIs around 4-8 ms, a large tail that includes ISIs out to 190 ms, and a small second model around 90-100 ms. This suggest that a simple Poisson model will not be able to fully capture the structure observed in this data.

Question 2

Compute a 5-number summary (min, .25 quantile, median, .75 quantile and max) and a box plot for the ISI distribution.

```
1 quantile(ISIs, probs=seq(0,1,by=.25))
2 0% 25% 50% 75% 100%
3 2 5 10 44 190
boxplot(ISIs)
```

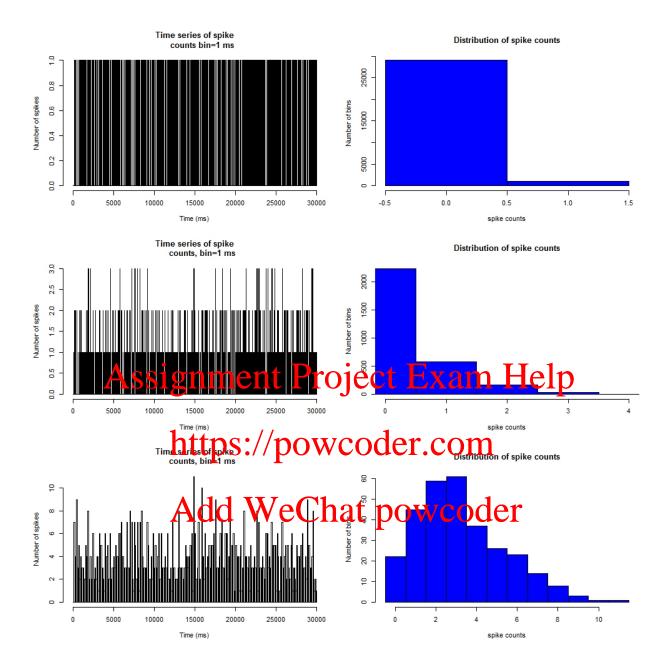


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The 5-number summary and boxplot confirm our earlier observation about the distribution of ISIs. Most spikes occur at small ISIs but to 13-40 mm Out two are the formula ISIs over 100 ms.

Question 3 Discrete time by of fixed virthand to the Duple Ceches for occur in each time bin. Set bin size to be 1 ms, 10 ms and 100 ms. Plot the time series of spike counts and the distribution of spike counts as a histogram for each bin width. They look similar to the model distribution of increments of Poisson processes.

```
par(mfrow=c(3,2))
           spike1 <- hist(spiketimes, breaks=c(0:30000), main = 'Time_series_of_spike
           counts_bin=1_ms', xlab='Time_(ms)', ylab = 'Number_of_spikes')
           hist (spike1 \$counts, breaks=c(0:2) - .5, col= 'blue', xlab='spike_counts',
           main = 'Distribution_of_spike_counts', ylab='Number_of_bins')
           {\tt spike10} \leftarrow {\tt hist} ({\tt spiketimes} \;,\; {\tt breaks} = {\tt seq} (0,30000, {\tt by} = 10),, \; {\tt main} = {\tt `Time\_series\_of\_spike1} 
           \verb|counts|, \verb|bin=1| ms', | xlab='Time\_(ms)', | ylab='Number\_of\_spikes'|
           hist(spike10$counts, breaks=c(0:5)-.5, col='blue', xlim=c(0,4),, xlab='spike_counts',
           main = 'Distribution_of_spike_counts', ylab='Number_of_bins')
           spike100 \leftarrow hist(spiketimes, breaks = seq(0.30000,by=100), main = 'Time_series_of_spike100', main = 'Time_s
10
           counts, _bin=1_ms', xlab='Time_(ms)', ylab = 'Number_of_spikes')
11
           hist (spike100$counts, breaks=c(0:12)-.5, col= 'blue',, xlab='spike_counts',
12
           main = 'Distribution_of_spike_counts', ylab='Number_of_bins')
```



The log-likelihood for a homogeneous Poisson process with rate parameter λ is given by

$$\log L(\lambda) = N(T)\log(\lambda \Delta t) - \lambda T$$

thus the MLE of λ is

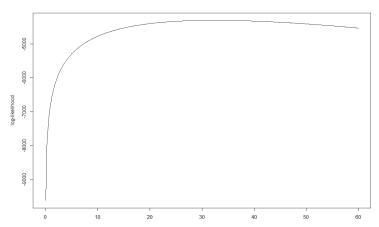
$$\hat{\lambda} = \frac{N(T)}{T} = \frac{972}{30} = 32.4000$$

with

$$s.e.(\hat{\lambda}) = \sqrt{N(T)}/T = 1.039$$

and 95% CI to be

$$[\lambda - 1.96 * s.e., \lambda + 1.96 * s.e.] = [30.3631, 34.4369]$$



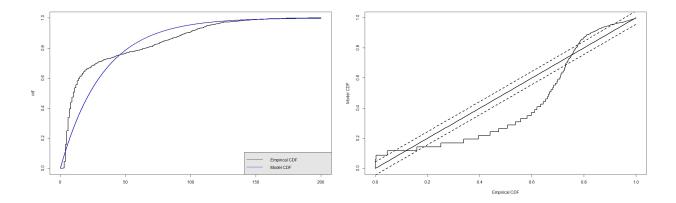
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Question 5

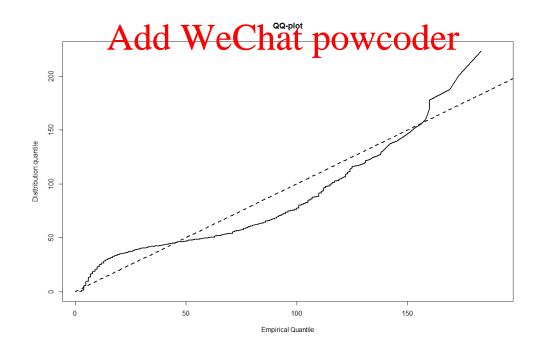
```
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    lam = N/30
    w \leftarrow seq(0, 200, by=.01)
    Femp <- numeric(length(w))
    Qs <- ISIs [order (ISIs)]
    for (i in 1:length(w)) (Fem i < Vine by plot(w, Femp, type='l', xlab ='', ylab =
    lines(w, 1-exp(-w/lam), col='blue', lwd=2)
    legend('bottomright', c('Empirical_CDF', "Model_CDF"), col = c('black','blue'),
    lty = c(1, 1), merge = TRUE, bg = "gray90")
9
10
    \mathrm{KSstat} \, < - \, \max(\, \mathrm{abs}(1 \! - \! \exp(-w/\mathrm{lam}\,) \, - \, \mathrm{Femp/N})\,)
11
    \# \text{ KSstat} = 0.2518
12
13
    plot (Femp, 1-exp(-w/lam), type='l', ylab='Model_CDF', xlab='Empirical_CDF', lwd=2)
14
15
    x = seq(0,1,by = .01); y=x
16
    lines(x,y,lty=1, lwd=2)
    x=seq(0,1,by=.01); y=x+1.36/sqrt(N)
17
18
    lines(x,y,lty=2, lwd=2)
    x=seq(0,1,by=.01); y=x-1.36/sqrt(N)
19
    lines(x,y,lty=2, lwd=2)
20
```

$$KS - statistic = \max |\hat{F}_{Emp} - F_{dis}| = 0.2518$$

Note that the KS-statistic value may be different if you choose different precision of the Empirical cdf here.



From QQ-plot, we see that the expect.



```
FF1 <- var(spike1$counts)/ mean(spike1$counts); FF1 # 0.9676

qgamma(c(.025,.975), length(spike1$counts)/2, scale=2/length(spike1$counts)) #0.9841 1.0161

FF10 <- var(spike10$counts)/ mean(spike10$counts); FF10 # 1.1661

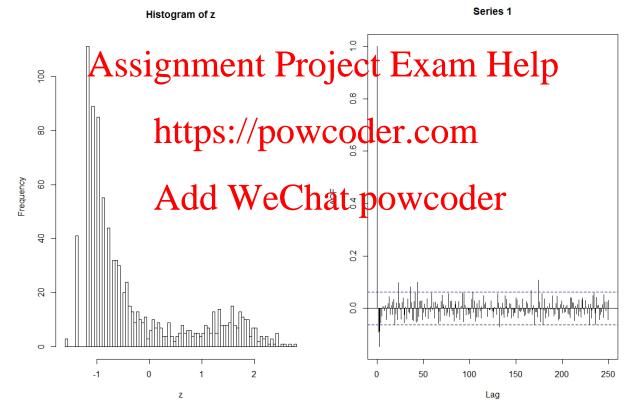
qgamma(c(.025,.975), length(spike10$counts)/2, scale=2/length(spike10$counts))# 0.9500 1.0512

FF100 <- var(spike100$counts)/mean(spike100$counts); FF100 # 1.4583

qgamma(c(.025,.975), length(spike100$counts)/2, scale=2/length(spike100$counts)) # 0.8464 1.1662
```

Question 8

Plot the rescaled ISIs and the autocorrelation function of the observed ISIs with 95% confidence bounds. Note the 95% confidence interval for the correlation coefficient of independent Gaussian rv is $1.96/\sqrt{n}$.



The rescaled ISIs are not normally distributed and at small lags, the autocorrelation function falls outside of the 95% confidence bounds more often than expected by chance alone for small lags, suggesting that nearby spikes have ISIs that are not independent.

Question 9

From the above analysis, we can conclude that under a simple Poisson model the firing rate of this neuron is likely somewhere between 30-34 Hz. However, this Poisson model fails to account for many aspects of the spiking activity including its refractoriness, bursting, and long ISIs. Our analysis of the Fano Factor suggests that there is less variability in the 1 ms increments and more in the 20 ms and 100 ms increments than what would be expected from a Poisson process. The autocorrelation analysis suggests a dependence structure on past spiking activity. Therefore, describing the structure of this data will require history dependent point process models.

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