

TSSL Lab 3 - Nonlinear state space models and Sequential Monte Carlo

In this lab we will make use of a non-linear state space model for analyzing the dynamics of SARS-CoV-2, the virus causing covid-19. We will use an epidemiological model referred to as a Susceptible Exposed Infectious Recovered (SEIR) model. It is a stochastic adaptation of the model used by the The Public Health Agency of Sweden for predicting the spread of covid-19 in the Stockholm region early in the pandemic, see [Estimates of the peak-inp and the number of infected individuals during the covid-19 outbreak in the Stockholm region, Sweden February – April 2020](#).

The background and details of the SEIR model that we will use are available in the document TSSL Lab 3 Predicting Covid-19 Description of the SEIR model on LISAM. Please read through the model description before starting on the lab assignments to get a feeling for what type of model that we will work with.

DISCLAIMER

Even though we will use a type of model that is common in epidemiological studies and analyze real covid-19 data, you should NOT read to much into the results of the lab. The model is intentionally simplified to fit the scope of the lab, it is not validated, and it involves several model parameters that are set somewhat arbitrarily. The lab is intended to be an illustration of how we can work with nonlinear state space models and Sequential Monte Carlo methods to solve a problem of practical interest, but the actual predictions made by the final model should be taken with a big grain of salt.

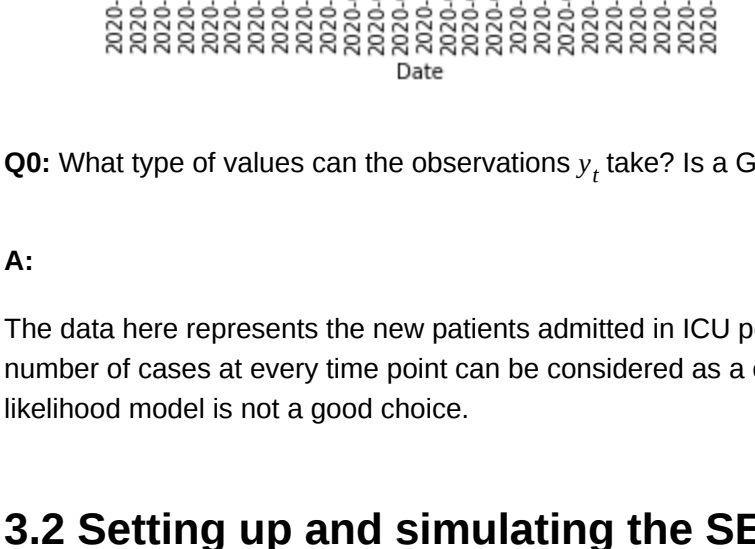
We load a few packages that are useful for solving this lab assignment.

```
In [3]: import pandas # Loading data / handling data frames
import numpy as np
import matplotlib.pyplot as plt
plt.rcParams["figure.figsize"] = (10, 8) # Increase default size of plots
```

3.1 A first glance at the data

The data that we will use in this lab is a time series consisting of daily covid-19 related intensive care cases in Stockholm from March to August. As always, we start by loading and plotting the data.

```
In [2]: data=pandas.read_csv('SR_Stockholm.csv', header=0)
# Read in the data
u_sthln = data['date'].values
n_data = len(y_sthln)
p1=plt.plot(u_sthln, y_sthln)
p1.xticks(range(0, n_data, 7), u_sthln[::7], rotation = 90) # Show only one tick per week for clarity
p1.xlabel('Date')
p1.ylabel('New ICU cases')
p1.show()
```



Q0: What type of values can the observations y_t take? Is a Gaussian likelihood model a good choice if we want to respect the properties of the data?

A:

The data here represents the new patients admitted in ICU per week. There can be same number of cases multiple days. So, it is not a continuous data. The number of cases at every time point can be considered as a class. This makes the data a classification data. Since it's classification problem, Gaussian likelihood model is not a good choice.

3.2 Setting up and simulating the SEIR model

In this section we will set up a SEIR model and use this to simulate a synthetic data set. You should keep these simulated trajectories, we will use them in the following sections.

```
In [3]: from tsstools.lab3 import Param, SEIR

***For Stockholm the population is probably roughly 2.5 million.***
population_size = 2500000

**** Binomial probabilities (p.se, p.ei, p.if, and p.ic) and the transmission rate (rho)****
pse = 1 # This controls the rate of spontaneous s->e transitions. It is set to zero for this lab.
pei = 1 / 5.1 # Based on rfm report
pir = 1 / 1 # Based on rfm report
pic = 1 / 1000 # Quite arbitrary
rho = 0.3 # Quite arbitrary

***The instantaneous contact rate b(t) is modeled as
b(t) = exp(z(t))
z(t) = z(t-1) + epsilon(t), epsilon(t) ~ N(0, sigma_epsilon/2)
****
sigma_epsilon = .1

*** For setting the initial state of the simulation***
l0 = 1000 # Mean number of infectious individuals at initial time point
l0 = 5000 # Mean number of exposed...
rb = 0 # Mean number of recovered
l0 = population_size - l0 - eb - 0 # Mean number of susceptible
init_mean = np.array([l0, eb, 10, 0.1], dtype=np.float64) # The last 0.1 is the mean of z[0]

***All the above parameters are stored in params.***
params = Param(pse, pei, pir, pic, rho, sigma_epsilon, init_mean, population_size)

*** Create a model instance***
model = SEIR(params)
```

Q1: Generate 10 different trajectories of length 200 from the model an plot them in one figure. Does the trajectories look reasonable? Could the data have been generated using this model?

For reproducibility, we set the seed of the random number generator to 0 before simulating the trajectories using np.random.seed(0)

Save these 10 generated trajectories for future use.

(Hint: The SEIR class has a simulate method)

```
In [4]: np.random.seed(0)
help(model.simulate)

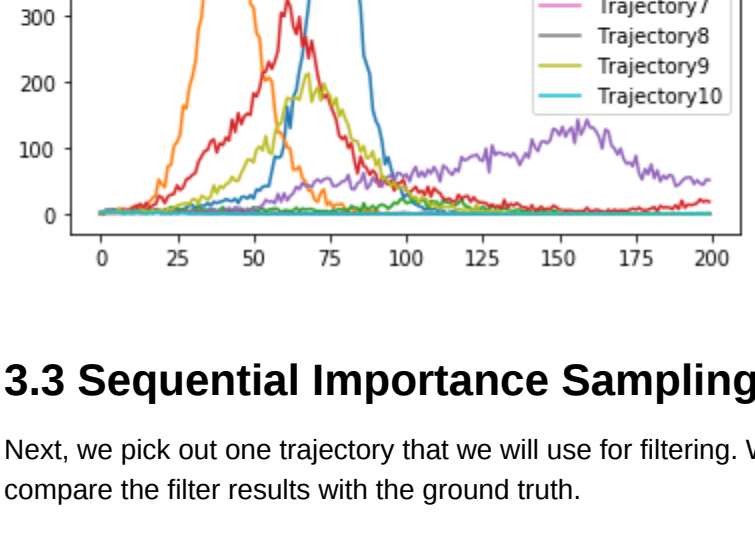
Help on method simulate in module tsstools.lab3:

simulate(t, N=1) method of tsstools.lab3.SEIR instance
Simulate the SEIR model for a given number of time steps. Multiple trajectories
can be simulated simultaneously.

Parameters
-----
t : int
    Number of time steps to simulate the model for.
N : int, optional
    Number of independent trajectories to simulate. The default is 1.

Returns
-----
alpha : ndarray
    Array of size (d,N,t) with state trajectories. alpha[:,i,:] is the i:th trajectory.
y : ndarray
    Array of size (1,N,t) with observations.
```

```
In [5]: genModel = model.simulate(200,10)
for i in range(10):
    plt.plot(genModel[i][0,1,:], label = "Trajectory"+ str(i+1))
    plt.legend()
plt.title("Model Simulation")
plt.rcParams["figure.figsize"] = [5, 5]
```



3.3 Sequential Importance Sampling

Next, we pick out one trajectory that we will use for filtering. We use simulated data to start with, since we then know the true underlying SER states and can compare the filter results with the ground truth.

Q2: Implement the Sequential Importance Sampling algorithm by filling in the following functions.

The exp_norm function should return the normalized weights and the log average of the unnormalized weights. For numerical reasons, when calculating the weights we should 'normalize' the log-weights first by removing the maximal value.

Let $u_t = \max(\log w_t)$ and take the exponential of $\log w_t - u_t = \log w_t' - u_t$. Normalizing w_t' will yield the normalized weights!

For the log average of the unnormalized weights, care has to be taken to get the correct output, $\log(1/N \sum_{i=1}^N w_t) = \log(1/N \sum_{i=1}^N e^{u_t + \log w_t'}) = \log(1/N \sum_{i=1}^N e^{\log w_t'}) + u_t$. We are going to need this in the future, so best to implement it right away.

(Hint: look at the SEIR model class, it contains all necessary functions for propagation and weighting)

```
In [6]: from tsstools.lab3 import smc_res

def exp_norm(logwgt):
    """
    Exponentiates and normalizes the log-weights.

    Parameters
    -----
    logwgt : ndarray
        Array of size (N,) with log-weights.

    Returns
    -----
    wgt : ndarray
        Array of size (N,) with normalized weights, wgt[i] = exp(logwgt[i])/sum(exp(logwgt)),
        but computed in a numerically robust way!
    logZ : float
        Log of the normalizing constant, logZ = log(sum(exp(logwgt))),
        but computed in a numerically robust way!

    wgtbar = np.max(logwgt)
    wgt = np.exp(logwgt - wgtbar)
    logZ = np.log(1/len(wgt)*sum(wgt)) - wgtbar

    return wgt/sum(wgt), logZ

def ESS(wgt):
    """
    Computes the effective sample size.

    Parameters
    -----
    wgt : ndarray
        Array of size (N,) with normalized importance weights.

    Returns
    -----
    ess : float
        Effective sample size.
    """
    ess = (sum(wgt)**2)/sum(wgt**2)

    return ess

def sis_filter(model, y, N):
    d = model.d
    n = len(y)

    # Allocate memory
    particles = np.zeros((d, N, n), dtype = float) # All generated particles
    logw = np.zeros((1, N, n)) # Unnormalized log-weight
    W = np.zeros((1, N, n)) # Normalized weight
    alpha_filt = np.zeros((d, 1, n)) # Store filter mean
    N_eff = np.zeros(n) # Efficient number of particles
    logZ = 0. # Log-likelihood estimate

    # Filter loop
    for t in range(n):
        # Sample from "bootstrap proposal"
        if t == 0:
            particles[:, :, 0] = model.sample_state(alpha=None, N=N) # Initialize from p(alpha_1)
            logw[:, :, 0] = model.logLik(y[t], particles[:, :, 0]) # Compute weights
        else:
            particles[:, :, t] = model.sample_state(particles[:, :, t-1], N) # Propagate according to dynamics
            logw[:, :, t] = logw[:, :, t-1] + model.logLik(y[t], particles[:, :, t]) # Update weights

            # Normalize the importance weights and compute N_eff
            W[:, :, t] = exp_norm(logw[:, :, t])
            N_eff[t] = ESS(W[:, :, t])

            # Compute filter estimates
            alpha_filt[:, :, t] = np.sum(W[:, :, t]*particles[:, :, t], axis = 1)/np.sum(W[:, :, t])

    return smc_res(alpha_filt, particles, W, logw=logw, N_eff=N_eff, logZ=logZ)
```

```
In [7]: y = genModel[1][0,1,:]
N = 100
sis_filter(model,y,N)

Out[7]: <tsstools.lab3.smc_res at 0x181f9542008>
```

Q3: Choose one of the simulated trajectories and run the SIS algorithm using $N = 100$ particles. Show plots comparing the filter means with the underlying truth of the infected, Exposed and Recovered.

Also show a plot of how the ESS behaves over the run.

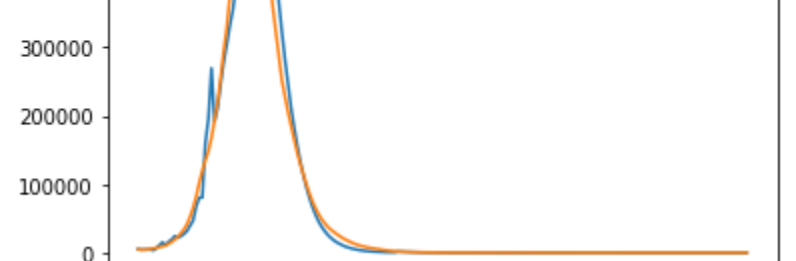
(Hint: In the model we use the S, E, I as states, but S will be much larger than the others. To calculate R, note that $S + E + I + R = \text{Population}$)

```
In [8]: y = genModel[1][0,1,:]
N = 100
sisModel = sis_filter(model,y,N)

In [9]: sisModel.alpha_filt.shape
Out[9]: (4, 1, 200)

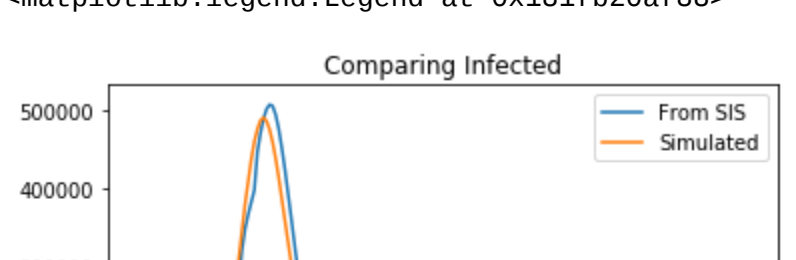
In [10]: plt.plot(sisModel.alpha_filt[0,0,:], label = "From SIS")
plt.plot(genModel[0][0,1,:], label = "Simulated")
plt.title("Comparing Susceptible")
plt.legend()

Out[10]: <matplotlib.legend.Legend at 0x181fa13fe8b>
```



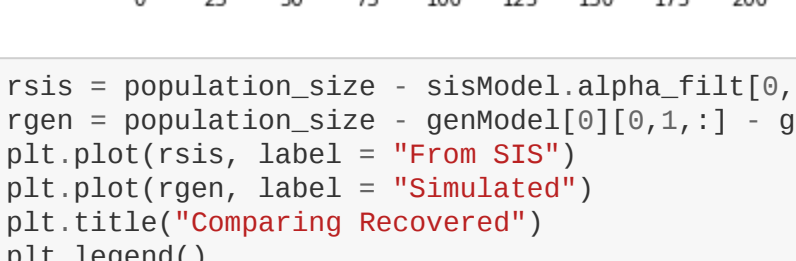
```
In [11]: plt.plot(sisModel.alpha_filt[0,1,:], label = "From SIS")
plt.plot(genModel[0][0,1,:], label = "Simulated")
plt.title("Comparing Exposed")
plt.legend()

Out[11]: <matplotlib.legend.Legend at 0x181fb19108b>
```



```
In [12]: plt.plot(sisModel.alpha_filt[0,2,:], label = "From SIS")
plt.plot(genModel[0][0,1,:], label = "Simulated")
plt.title("Comparing Infected")
plt.legend()

Out[12]: <matplotlib.legend.Legend at 0x181fb20a8fb>
```



```
In [13]: rsis = population_size - sisModel.alpha_filt[0,0,:] - sisModel.alpha_filt[1,0,:] - sisModel.alpha_filt[2,0,:]
rgen = population_size - genModel[0][0,1,:] - genModel[0][1,1,:] - genModel[0][2,1,:]
plt.plot(rsis, label = "From SIS")
plt.plot(rgen, label = "Simulated")
plt.title("Comparing Recovered")
plt.legend()

Out[13]: <matplotlib.legend.Legend at 0x181fb275c8b>
```



3.4 Sequential Importance Sampling with Resampling

Pick the same simulated trajectory as for the previous section.

Q4: Implement the Sequential Importance Sampling with Resampling or Bootstrap Particle Filter by completing the code below.

```
In [14]: def bpf(model, y, numParticles):
    d = model.d
    n = len(y)
    N = numParticles

    # Allocate memory
    particles = np.zeros((d, N, n), dtype = float) # All generated particles
    logw = np.zeros((1, N, n)) # Unnormalized log-weight
    W = np.zeros((1, N, n)) # Normalized weight
    alpha_filt = np.zeros((d, 1, n)) # Store filter mean
    N_eff = np.zeros(n) # Efficient number of particles
    logZ = 0. # Log-likelihood estimate

    # Filter loop
    for t in range(n):
        # Sample from "bootstrap proposal"
        if t == 0: # Initialize from prior
            particles[:, :, 0] = model.sample_state(alpha=init_mean.reshape(-1), N=N)
        else:
            # Resample and propagate according to dynamics
            ind = np.random.choice(N, N, replace=True, p=W[:, :, t-1])
            resampled_particles = particles[:, ind, t-1]
            particles[:, :, t] = model.sample_state(alpha=resampled_particles, N=N)

        # Compute weights
        logw[:, :, t] = model.logLik(y[t], particles[:, :, t])
        W[:, :, t] = exp_norm(logw[:, :, t])
        logZ += logZ_new + max(logw[:, :, t]) # Update log-likelihood estimate
        N_eff[t] = ESS(W[:, :, t])

        # Compute filter estimates
        alpha_filt[:, :, t] = np.sum(W[:, :, t]*particles[:, :, t], axis = 1)/np.sum(W[:, :, t])

    return smc_res(alpha_filt, particles, W, N_eff = N_eff, logZ = logZ)
```

```
In [15]: bpf(model,y,100)

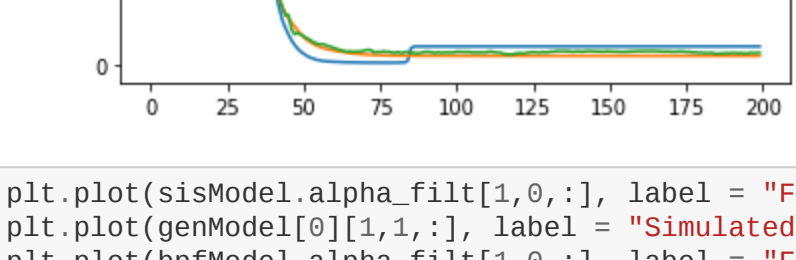
Out[15]: <tsstools.lab3.smc_res at 0x181fb27a208b>
```

Q5: Use the same simulated trajectory as above and run the BPF algorithm using $N = 100$ particles. Show plots comparing the filter means from the Bootstrap Particle Filter algorithm with the underlying truth of the infected, Exposed and Recovered. Also show a plot of how the ESS behaves over the run. Compare this with the results from the SIS algorithm.

```
In [16]: bpfModel = bpf(model,y,100)

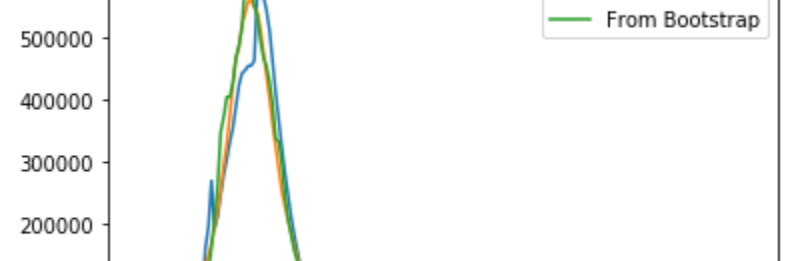
In [17]: plt.plot(sisModel.alpha_filt[0,0,:], label = "From SIS")
plt.plot(genModel[0][0,1,:], label = "Simulated")
plt.plot(bpfModel.alpha_filt[0,0,:], label = "From Bootstrap")
plt.title("Comparing Susceptible")
plt.legend()

Out[17]: <matplotlib.legend.Legend at 0x181fb31d0c8>
```



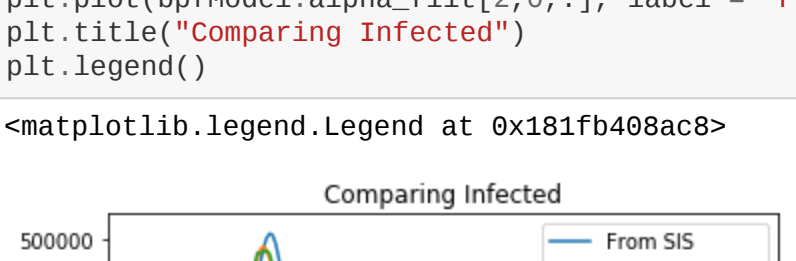
```
In [18]: plt.plot(sisModel.alpha_filt[0,1,:], label = "From SIS")
plt.plot(genModel[0][0,1,:], label = "Simulated")
plt.plot(bpfModel.alpha_filt[0,1,:], label = "From Bootstrap")
plt.title("Comparing Exposed")
plt.legend()

Out[18]: <matplotlib.legend.Legend at 0x181fb386c8b>
```



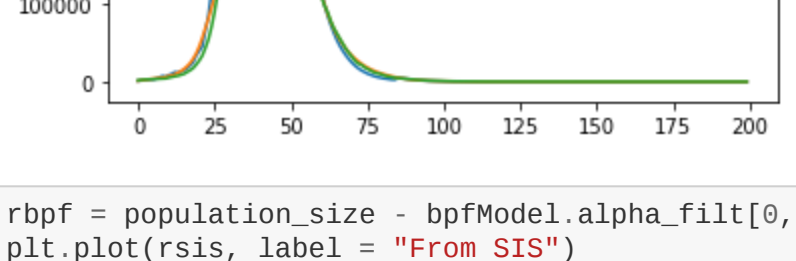
```
In [19]: plt.plot(sisModel.alpha_filt[0,2,:], label = "From SIS")
plt.plot(genModel[0][0,1,:], label = "Simulated")
plt.plot(bpfModel.alpha_filt[0,2,:], label = "From Bootstrap")
plt.title("Comparing Infected")
plt.legend()

Out[19]: <matplotlib.legend.Legend at 0x181fb400ac8b>
```

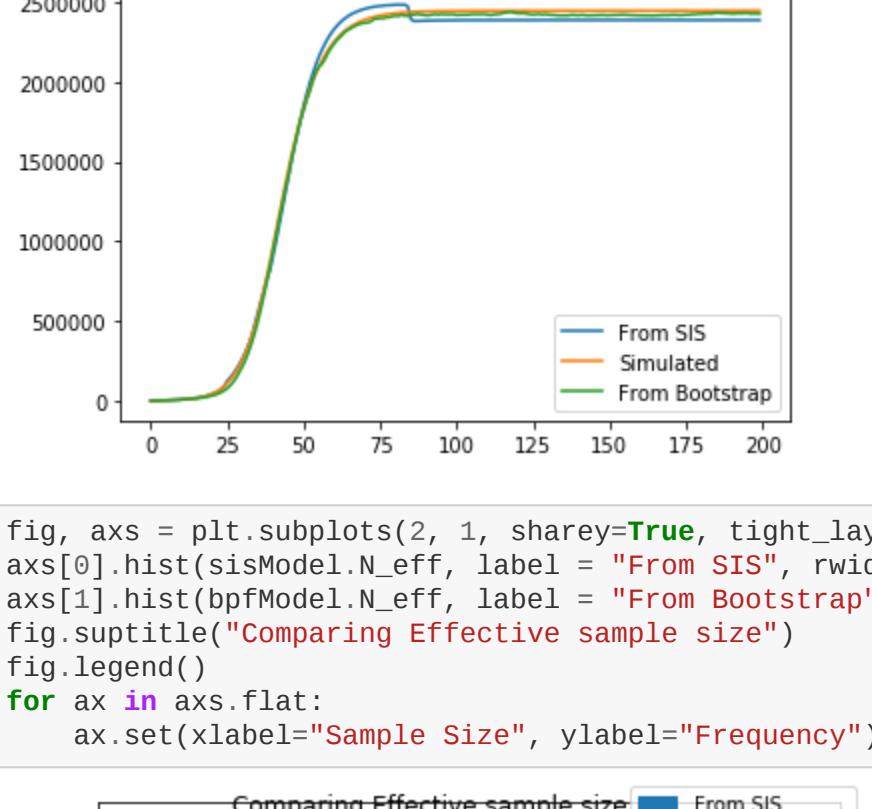


```
In [20]: rbpf = population_size - bpfModel.alpha_filt[0,0,:] - bpfModel.alpha_filt[1,0,:] - bpfModel.alpha_filt[2,0,:]
rgen = population_size - genModel[0][0,1,:] - genModel[0][1,1,:] - genModel[0][2,1,:]
plt.plot(rbpf, label = "From Bootstrap")
plt.plot(rgen, label = "Simulated")
plt.title("Comparing Recovered")
plt.legend()

Out[20]: <matplotlib.legend.Legend at 0x181fb488ec8b>
```



```
In [21]: fig, axs = plt.subplots(2, 2, sharey=True, tight_layout=True)
axs[0].hist(sisModel.N_eff, label = "From SIS", rwidth=0.85, bins = 20)
axs[1].hist(bpfModel.N_eff, label = "From Bootstrap", rwidth=0.85, bins = 20, color='red', alpha = 0.5)
fig.legend(['Comparing Effective sample size'])
for ax in axs.flat:
    ax.set(xlabel="Sample Size", ylabel="Frequency")
```



For the particle filter without resampling, out of 200, the effective samples were 0 for more than 150 times. While in the Bootstrap particle filter, the samples are resampled according to weights where the sample with higher weights is given more prominence. As we can see the size is 100 for most of the time.

Since we have considered a better sample size, the alpha values of Bootstrap sample are almost equal to the true values compared with the SIS model.

3.5 Estimation of the data likelihood and learning a model parameter

In this section we consider the real data and learning the model using this data. For simplicity we will only look at the problem of estimating the ρ parameter and assume that others are fixed.

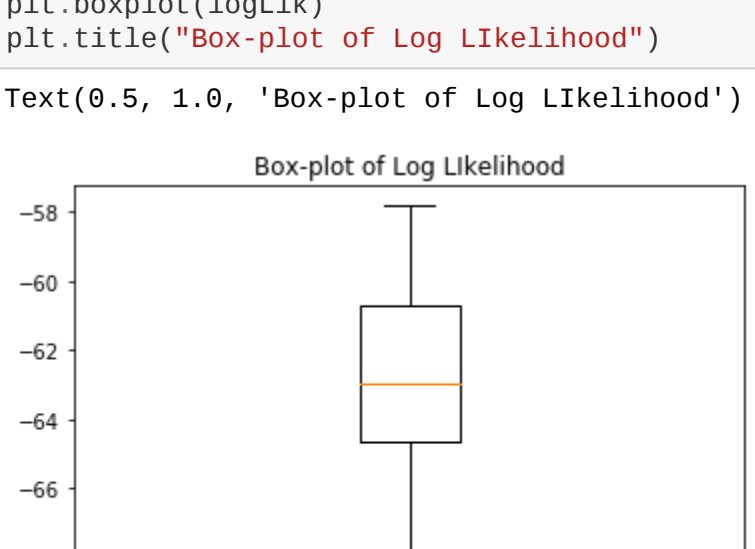
You are more than welcome to also study the other parameters.

Before we begin to tweak the parameters we run the particle filter using the current parameter values to get a benchmark on the log-likelihood.

```
In [22]: loglik = []
for i in range(20):
    logZ = bpf(model, y, 200).logZ
    loglik.append(logZ)
```

```
In [23]: plt.boxplot(loglik)
plt.title("Box-plot of Log Likelihood")

Out[23]: Text(0.5, 1.0, "Box-plot of Log Likelihood")
```



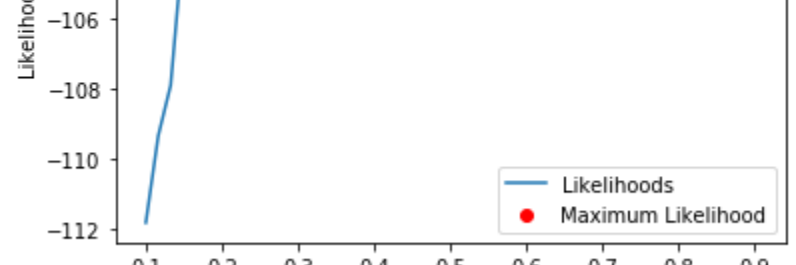
Q7: Make a grid of the ρ parameter in the interval [0.1, 0.5]. Use the bootstrap particle filter to calculate the log-likelihood for each value. Run the bootstrap particle filter using $N = 100$ multiple times (at least 20 per value) and use the average as your estimate of the log-likelihood. Plot the log-likelihood function and mark the maximal value.

(Hint: use np.linspace to create a grid of parameter values)

```
In [24]: rhos = np.linspace(start=0.1, stop=0.4, steps=4)
avgs = []
for rho in rhos:
    params = Param(pse, pei, pir, pic, rhos[0], sigma_epsilon, init_mean, population_size)
    model = SEIR(params)
    loglik = [bpf(model, y_sthln, 100).logZ for n in range(20)]
    avgs.append(np.mean(loglik))
```

```
In [25]: rhoOpt = rhos[np.argmax(avgs)]
plt.plot(rhos, avgs, label = "Log Likelihoods")
plt.scatter(rhoOpt, max(avgs), color = "red", label = "Maximum Likelihood")
plt.xlabel("rho value")
plt.ylabel("Likelihood")
plt.legend()
```

```
Out[25]: <matplotlib.legend.Legend at 0x181fb0b488b>
```



```
In [26]: print("The Maximum Likelihood is {:.4f} and the optimal rho value with maximum Likelihood is {:.4f}".format(max(avgs), rhoOpt))

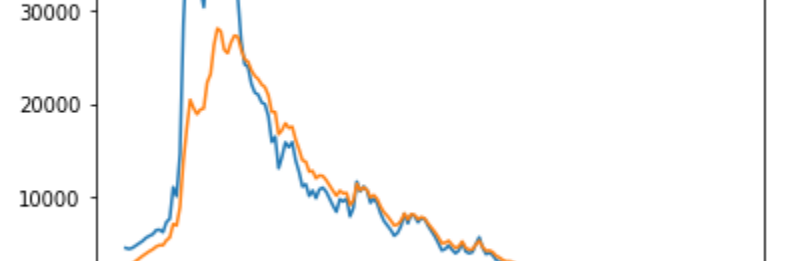
The Maximum Likelihood is -186.6764 and the optimal rho value with maximum Likelihood is 0.4265
```

Q8: Run the bootstrap particle filter on the full dataset with the optimal ρ value. Present a plot of the estimated infected, Exposed and Recovered states.

```
In [27]: params = Param(pse, pei, pir, pic, rhoOpt, sigma_epsilon, init_mean, population_size)
model = SEIR(params)
optModel = bpf(model, y_sthln, 200)
```

```
In [28]: rExp = plt.subplots(2, 2, sharey=True, tight_layout=True)
plt.plot(optModel.alpha_filt[0,0,:], label = "Exposed")
plt.plot(optModel.alpha_filt[0,1,:], label = "Infected")
plt.legend()

Out[28]: <matplotlib.legend.Legend at 0x181fb0b488b>
```



```
In [29]: rRec = population_size - optModel.alpha_filt[0,0,:] - optModel.alpha_filt[1,0,:] - optModel.alpha_filt[2,0,:]
plt.plot(rRec, label = "Recovered")
plt.legend()

Out[29]: <matplotlib.legend.Legend at 0x181fb0b488b>
```

