## BPL YEAST AIR Fedbatch demo

This notebook demonstrate yeast fedbatch cultivation. We look at impact of changes in the glucose feeding. We also take a look at tuning of the DO-control system. Both liquid- and gasphase are included in the model.

Interaction with the compiled model as FMU is mainly through the simplified commands: par(), init(), newplot(), simu() etc. The last simulation is always available in the workspace and called 'sim\_res'. The command describe() brings mainly up description infomration from the actual Modelica code from the FMU but is complemented with information given in the dedicated Python setup-file.

The idea is to demonstrate how simulations and varyiing conditions can provide some process insight that can support the experimetnal work. I hope that at the end of this session you are ready to formulate your own questions you want to address with simulations - and you can just go on in this notebook! Just press the field "+Code" in the upper left part of notebook interface and you get a new "cell" where you write your own code. You can copy and paste from cells above using ctrol-c and ctrl-p as usual and edit the cell. When your are ready to execute the cell just press the "play button" to the left in the cell or press shift-enter as in "ordinary" Jupyter notebooks.

After a session you may want to save your own notebook. That you can do on your Google Drive account and I refer to Colab instructions for how to do this. It is easy.

## Enjoy!

Now specific installation and the run simulations. Start with connecting to Github. Then upload the two files:

- FNU BPL YEAST AIR Fedbatch linux jm cs.fmu
- Setup-file BPL YEAST AIR Fedbatch explore

## Filter out DepracationWarnings for 'np.float as alias' is needed - wish I could make filter more narrow

import warnings warnings.filterwarnings("ignore")

%%bash git clone https://github.com/janpeter19/BPL\_YEAST\_AIR\_Fedbatch

%cd BPL YEAST AIR Fedbatch

```
In [1]: run -i BPL_YEAST_AIR_Fedbatch_DOcontrol_explore.py
```

```
Linux - run FMU pre-comiled JModelica 2.4
```

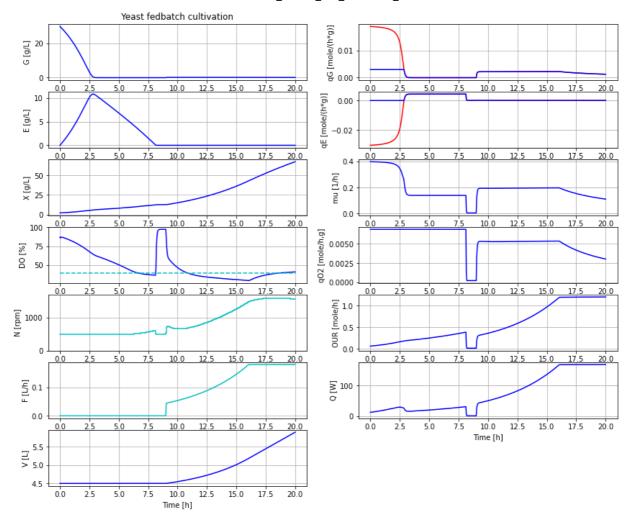
```
Model for bioreactor has been setup. Key commands:
- par() - change of parameters and initial values
```

init()change initial values only

- simu() - simulate and plot - newplot() - make a new plot

show()show plot from previous simulation

```
- display parameters and initial values from the last simulati
         - disp()
        on
         - describe() - describe culture, broth, parameters, variables with values /
        units
        Note that both disp() and describe() takes values from the last simulation
        Brief information about a command by help(), eg help(simu)
        Key system information is listed with the command system info()
        plt.rcParams['figure.figsize'] = [36/2.54, 30/2.54]
In [3]: describe('culture'); print(); describe('liquidphase'); print(); describe('gas
        Saccharomyces cerevisae - default parameters for strain H1022
        Reactor broth substances included in the model
        Cells
                            = 1 - molecular weight = 24.6 Da
                index
        Glucose index
                            = 2 - molecular weight = 180.0 Da
        Ethanol index = 3 - molecular weight = 46.0 Da Dissolved 02 index = 4 - molecular weight = 32.0 Da
        Dissolved CO2 index = 5 - molecular weight = 44.0 Da
        Reactor gasphase substances included in the model
        N2 etc index = 1 - molecular weight = 28.0 Da
        02 index = 2 - molecular weight = 32.0 Da
        CO2 index
                     = 3 - molecular weight = 44.0 Da
        Ethanol index = 4 - molecular weight = 46.0 Da
        # Culture parameters and others at default values
In [4]:
         par(q02lim=0.0069)
         # Process initial conditions
         init(V 0=4.5, VG 0=4.5*30, VX 0=4.5*2, VE 0=4.5*0)
         # Feed profile
         par(t start=9, F start=0.044, mu feed=0.20, F max=0.18)
         # DO-control parameters
         par(samplePeriod=1/60, K=10, Ti=0.5, I 0=500)
         # Simulate and plot
         newplot(title='Yeast fedbatch cultivation', plotType='Extended 2')
         simu(20)
```

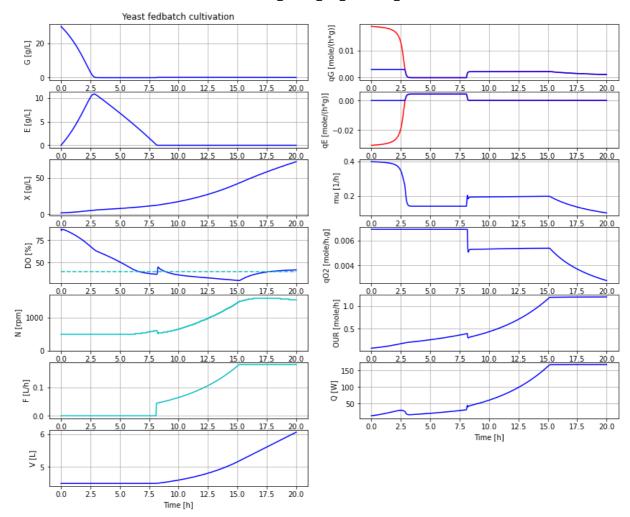


```
In [5]: disp('culture', decimals=4)
```

qGmax : 0.02 Ks : 0.01 q02lim : 0.0069

```
In [6]: # Let us start the feeding just after the batch phase has ended and keep othe
par(t_start=8.1)

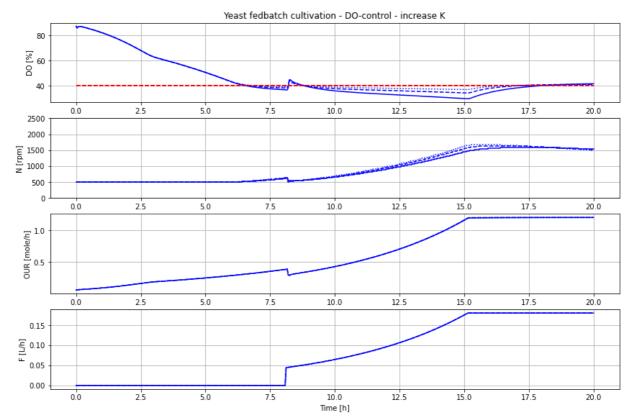
# Simulate and plot
newplot(title='Yeast fedbatch cultivation', plotType='Extended_2')
simu(20)
```



The increase of DO to about 50 % at end of batch phase should be possible to detect easily. This simulation is more realistic and we use these settings from now on.

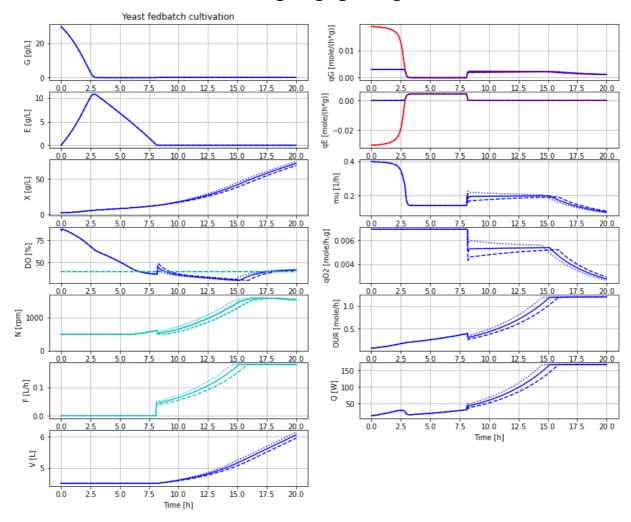
```
In [7]: # Let us take a closer look at the DO-control system and try to make control
   newplot(title='Yeast fedbatch cultivation - DO-control - increase K', plotType
   for value in [10, 20, 40]: par(K=value); simu(20)

# Reset K to the original value
   par(K=10)
```



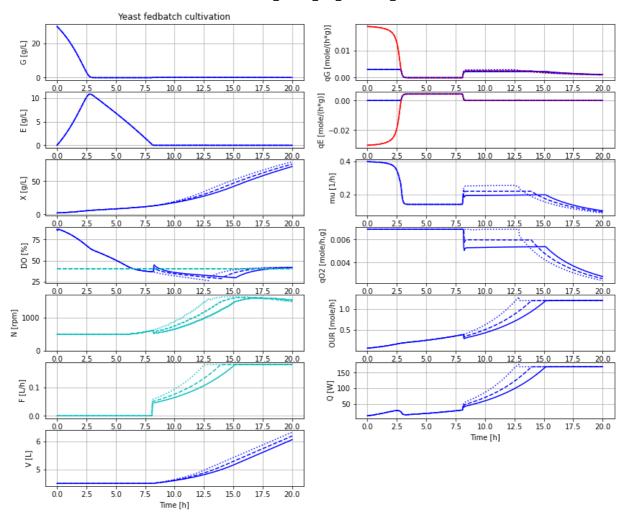
We see that by a higher control gain K the DO-control error get smaller and the stability of the control system is maintained.

```
In [8]: # Let us check the sensitivity to changes in the feed profile design
newplot(title='Yeast fedbatch cultivation', plotType='Extended_2')
for value in [0.044, 0.038, 0.050]: par(F_start=value); simu(20)
```

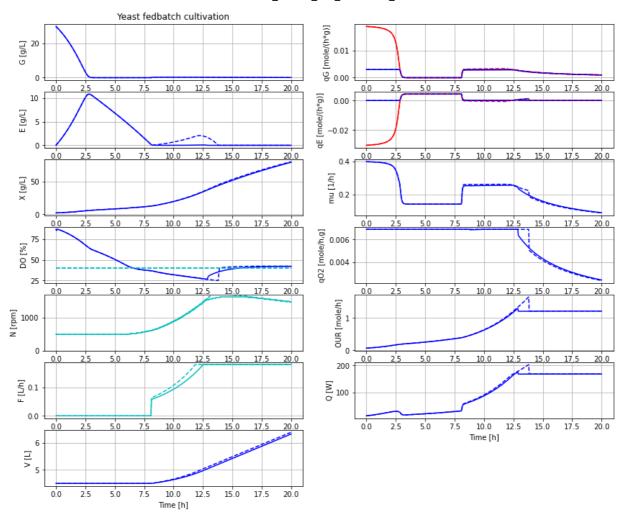


The variation in F\_start has an impact and we see that the actual growth rate during fedbatch phase do converge to the set growth rate of the feed, but i takes more than 5 hours.

```
In [9]: # Let us investigate a feedprofile that is closer to the maximal capacity
    newplot(title='Yeast fedbatch cultivation', plotType='Extended_2')
    par(F_start=0.044, mu_feed=0.20); simu(20)
    par(F_start=0.050, mu_feed=0.22); simu(20)
    par(F_start=0.057, mu_feed=0.26); simu(20)
```



```
In [10]: # And let us see what happens if the feedprofile exceed the culture capacity
newplot(title='Yeast fedbatch cultivation', plotType='Extended_2')
par(F_start=0.057, mu_feed=0.26); simu(20)
par(F_start=0.063, mu_feed=0.28); simu(20)
par(F_start=0.044, mu_feed=0.20)
```



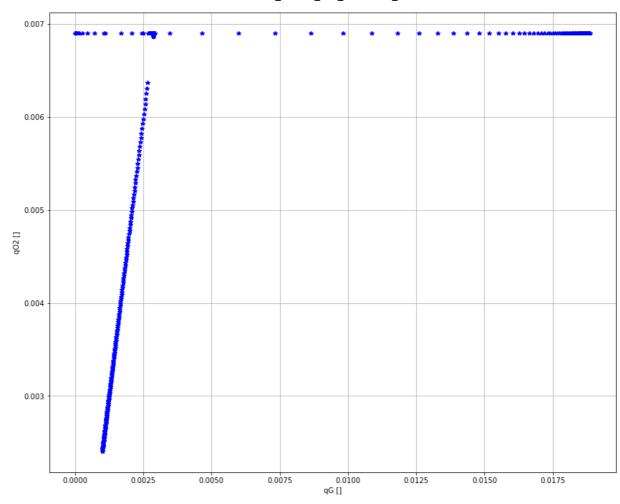
Note that with the feedprofile that exceed culture respiratory capacity, ethanol is accumulated during time 8-12.5 hours. When the feedprofile then is constant from time 12.5 hours and on, then the accumulated ethanol is consumed over about an hour. This leads to a higher oxygen demand and heat production during this time. The specific cell growth rate is also slightly higher during this period.

```
In [12]: # Improvise and make your own diagram - study the relation q02 vs qG(G)

plt.figure()
ax1 = plt.subplot(1,1,1)
ax1.set_ylabel('q02 []')
ax1.set_xlabel('qG []')
ax1.grid()

setLines()

diagrams.clear()
diagrams.append("ax1.plot(sim_res['bioreactor.culture.qGm'], sim_res['bioreactor.culture.qGm'], sim_res['bioreactor.culture.qGm'], sim_res['bioreactor.culture.qGm']
```



In [14]: # List of components in the process setup and also a couple of other things
describe('parts')

['airFlow\_setpoint', 'airtube', 'atmosphere', 'bioreactor', 'bioreactor.cultu re', 'bioreactor.gas\_liquid\_transfer', 'compressor', 'DO\_setpoint', 'dosagesc heme', 'DOsensor', 'feedtank', 'gasphase', 'liquidphase', 'MSL', 'N\_high', 'N\_low', 'PIreg', 'pump']

In [15]: system\_info()

System information

-OS: Linux

-Python: 3.8.2

-PyFMI: 2.7.4

-FMU by: JModelica.org

-FMI: 2.0

-Type: FMUModelCS2

-Name: BPL YEAST AIR.Fedbatch DOcontrol

-Generated: 2022-08-26T11:07:26

-MSL: 3.2.2 build 3

-Description: Bioprocess Library version 2.1.0 beta

-Interaction: FMU-explore ver 0.9.2

In [ ]: