In [2]: %matplotlib inline import sys sys.path.append("..") 1. Setup Imports the needed dependencies to run this tutorial. In [3]: import glob import numpy as np import tensorflow as tf import matplotlib.pyplot as plt import deeptrack as dt 2. Loading experimental holographic images Experimental images are located in sample-data folder In [4]: # Load figure 3 data (feeding events) predator sequence = np.load( "../data/data figure3/predator sequence.npy" prey sequence = np.load( "../data/data figure3/prey sequence.npy" # Load figure 4 data (division events) cell1 sequence = np.load( "../data/data\_figure4/Cell1\_sequence.npy" cell2 sequence = np.load( "../data/data\_figure4/Cell2\_sequence.npy" cell3 sequence = np.load( "../data/data\_figure4/Cell3\_sequence.npy" 2.1. Visualising prey and predator planktons In [5]: plt.figure(figsize=(10, 5)) plt.subplot(1,2,1)plt.title("Predator hologram (\$\it{Oxyrrhis\ Marina}\$)") plt.imshow(predator\_sequence[0], cmap="gray") plt.subplot(1,2,2)plt.title("Prey hologram (\$\it{Dunaliella\ tertiolecta}\$)") plt.imshow(prey\_sequence[0], cmap="gray") Out[5]: <matplotlib.image.AxesImage at 0x7fa5a68ef6a0> Predator hologram (Oxyrrhis Marina) Prey hologram (Dunaliella tertiolecta) 10 20 20 30 40 50 60 10 60 10 20 50 60 3. Loading pre-trained WAC-Net model We define the WAC-Net model and load the weights from a pre-trained model. In [6]: # WAC-Net trained for feeding events feeding model = dt.models.WACNet(outputs = 2) # 2 outputs: dry mass and radius feeding model.load weights( ".../pre-trained-models/WACNet dry mass.h5" # WAC-Net models trained for division events division\_event\_models\_loc = glob.glob( "../pre-trained-models/division-event-models/\*.h5" division models = []for model\_loc in division\_event\_models\_loc: empty model = dt.models.WACNet(outputs = 1) empty model.load weights(model loc) division models.append(empty model) 4. Using WAC-Net on experimental sequences 4.1. Example 1 - Feeding events Feeding events are recorded in two sequences: • predator\_sequence contains predator frames • prey\_sequence contains prey frames 4.1.1. Normalising the experimental image Normalise and reshape the image for WAC-Net prediction. In [7]: def Normalise(images, batch = 15): Normalised = [] for i in range(len(images)): Normalised.append(images[i]/np.median(images[i])) Normalised = np.array(Normalised)-1proc = [] #sliding window for i in range(len(Normalised)-batch+1): proc.append(Normalised[i:i+batch]) **return** np.expand dims(proc, axis = -1) In [8]: | # Funtion to convert the predicted dry mass to real dry mass units (pico grams) **def** real dm(p, a=209.16, b=0.28, sp ri inc = 0.21):return (p\*a+b)/sp\_ri\_inc 4.1.2. Predictions on experimental sequences Experimental sequences are normalized with batch size of 1. batch value can be increased to generate a sliding window of the experimental sequence. The WAC-Net assigns weights to the images with in sliding windows to predict the best possible value for dry mass considering all the frames. In [9]: prediction predator = feeding model.predict( Normalise( predator\_sequence, batch = 1prediction prey = feeding model.predict( Normalise( prey sequence, batch = 1) 4.1.3. Converting predictions to real dry mass units In [10]: drymass predator = real dm(prediction predator[:,0]) drymass\_prey = real\_dm(prediction\_prey[:,0]) 4.4. Visualising the dry mass transition in a feeding event In [11]: feeding at = 219 fig, ax = plt.subplots(figsize=(15, 5)) plt.title("Feeding event") plt.plot(np.arange(0, feeding\_at, 1), drymass\_predator[0:feeding\_at], linestyle='dashed', color='orange', marker='.', label='\$\it{Oxyrrhis\ Marina}\$ pre-feeding', alpha = 1, markersize = '5') plt.plot(np.arange(feeding\_at,len(drymass\_predator), 1),drymass\_predator[feeding\_at:], linestyle='dashe d',color='darkorange', marker='.', label='\$\it{Oxyrrhis\ Marina}\$ post-feeding', alpha = 1, markersize = '5') plt.plot(drymass\_prey[:feeding\_at], linestyle='dashed',color='blue', marker='.', label='\$\it{Dunaliella \ Teriolecta}\$', alpha=1, markersize = '5') plt.xlabel('Time (seconds)') plt.ylabel('Drymass (in picograms)') xticks = np.arange(0,550,50)

**Example 2. Using WAC-Net** 

%cd Quantitative-Microplankton-Tracker/examples/

obtain refined values of dry mass.

NOTE:

Open in Colab

This example demonstrates how to use a pre-trained WAC-Net (Weighted average convolutional neural network) on experimental images to

If you're running this notebook on your local machine, please comment the code in the cell below

In [1]: | git clone https://github.com/softmatterlab/Quantitative-Microplankton-Tracker.git



ax.set\_xticks(xticks)

plt.show()

700

600

500

400

300

200

100

0

4.2.1. Normalising

dm\_vals

dm cell1=[]

)[:,0]

dm cell2=[]

)[:,0]

dm cel13=[]

)[:,0]

In [12]:

In [13]:

In [14]:

In [15]:

Drymass (in picograms)

plt.legend(prop={'size': 16})

ax.set\_xticklabels([int(x\*(1/10)) for x in xticks])

Oxyrrhis Marina pre-feeding

Oxyrrhis Marina post-feeding

10

15

• cell1\_sequence contains frames of parent cell, Gen 1 daughter cell 1, and Gen 2 daughter cell 1

• cell3\_sequence contains frames of Gen 2 daughter cell 2 (along with parent cell and Gen 1 daugher cell 1)

• cell2\_sequence contains frames of Gen 1 daughter cell 2 (along with parent cell)

m = lambda rad, ri: ((4\*np.pi)/3) \* ((rad\*1e+6)\*\*3) \* (ri-1.33)return m(rad range[0], ri range[0]), m(rad range[1], ri range[1])

pred = real dm(pred, dm vals[1]-dm vals[0], dm vals[0])

pred = real dm(pred, dm vals[1]-dm vals[0], dm vals[0])

pred = real\_dm(pred, dm\_vals[1]-dm\_vals[0], dm\_vals[0])

4.2.2. Visualising the dry mass dynamics in a double division event

20

Dunaliella Teriolecta

4.2. Example 2 - Double division event

Division events are recorded in three sequences:

Function to de-normalise the dry mass values

def dm\_range(rad\_range, ri\_range):

for i in range(len(division models)):

cell1\_sequence, batch = 15

dm cell1 = np.mean( dm cell1, axis=0)

for i in range(len(division\_models)):

cell2 sequence,

dm cell2 = np.mean( dm cell2, axis=0)

for i in range(len(division models)):

cell3 sequence,

dm\_cell3 = np.mean(\_dm\_cell3, axis=0)

pred = division models[i].predict(

pred = division models[i].predict(

pred = division\_models[i].predict(

Out[12]: (5.361651462126584, 152.6814029644634)

Predictions on cell 1 sequence

Normalise(

\_dm\_cell1.append(pred)

Predictions on cell 2 sequence

Normalise(

dm cell2.append(pred)

Predictions on cell 3 sequence

Normalise(

\_dm\_cell3.append(pred)

dm vals = dm range([4e-6, 9e-6], [1.35, 1.38])

Feeding event

Time (seconds)

dt2 = 335end1 = 600 $daughter_lim = 10$ plt.plot(time[10:dt1], dm\_cell1[10:dt1], linestyle='dashed', color="midnightblue", marker='.', label='Parent Cell', alpha=1, markersize=10, linewidth=3) # parentcell1 plt.plot(time[dt1:dt2], dm\_cell1[dt1:dt2], linestyle='dashed', color="orange", marker='.', label='\nGeneration 1 - Daughter Cell 1', alpha=1, markersize='10', linewidth=3) # stationary cell plt.plot(time[dt1:dt2-daughter\_lim], dm\_cell2[dt1:dt2-daughter\_lim], linestyle='dashed', color="orange" marker='.', label='Generation 1 - Daughter Cell 2', alpha=1, markersize='10', linewidth=3) #d isplaced cell # Second generation---plt.plot(time[dt2:end1], dm\_cell1[dt2:end1], linestyle='dashed', color="darkcyan", marker='.', label='\nGeneration 2 - Daughter cell 1', alpha=1, markersize='10', linewidth=3) # stationary cell plt.plot(time[dt2:end1], dm\_cell3[dt2:end1], linestyle='dashed', color="darkcyan", marker='.', label='Generation 2 - Daughter cell 2', alpha=1, markersize='10', linewidth=3) # daughtercell1 **def** place\_ticks(x = 10, y = 10): plt.locator\_params(axis="x", nbins=x) plt.locator\_params(axis="y", nbins=y) xticks = np.arange(0,700,14.5)ax.set\_xticks(xticks) ax.set\_xticklabels([np.round(x\*(5/6/60),1) for x in xticks]) place\_ticks(x=10, y=10) fs = 30plt.xticks(fontsize=fs) plt.yticks(fontsize=fs) plt.legend(prop={'size': fs}) plt.ylabel("Dry mass (pg)", fontsize=fs) plt.xlabel("Time (hours)", fontsize=fs) plt.ylim([0, 700]) plt.show() 700<sub>T</sub> ---- Parent Cell 600 Generation 1 - Daughter Cell 1 ---- Generation 1 - Daughter Cell 2 500 Generation 2 - Daughter cell 1 Ory mass (pg) ---- Generation 2 - Daughter cell 2 200 100 4.0 Time (hours)

In [16]: | time = np.arange(0, len(dm\_cell1), 1)

dt1 = 128

fig,ax = plt.subplots(figsize=(40, 14))