

# Machine learning approaches to assess fouling on field samples

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## Collaborators



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M. Schultz

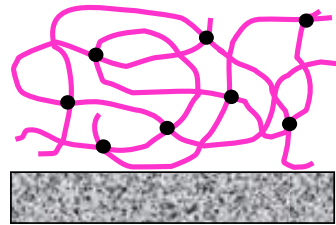


B. Rosenhahn

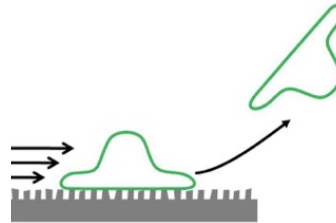
Antifouling conference Gothenburg, Sweden 2025



# Assessment of Fouling release coatings



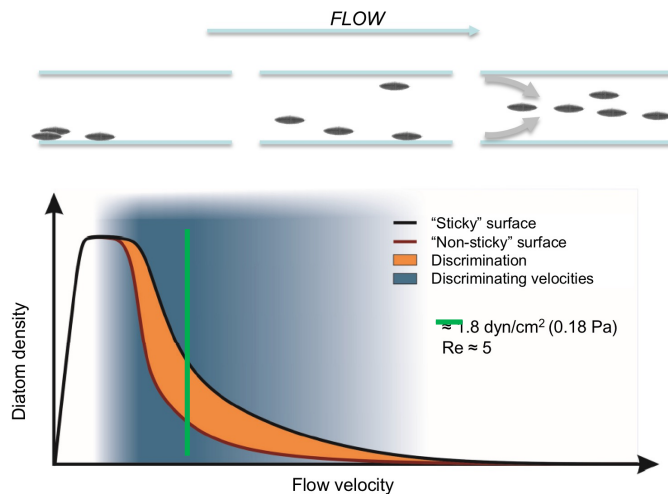
Low-fouling polymer



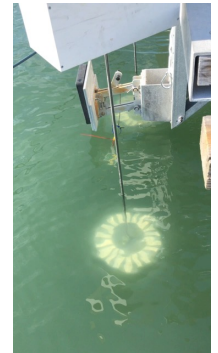
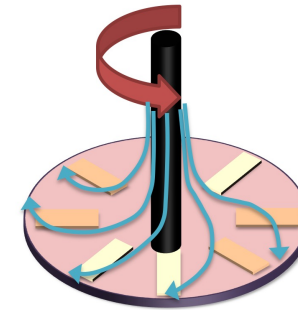
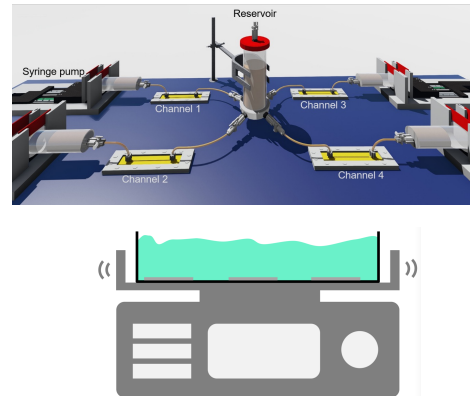
Fouling-release



Courtesy of K. Hunsucker (FIT)



Dynamic accumulation under flow as predictor for fouling release potential



FLORIDA  
TECH

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UNITED STATES NAVAL ACADEMY

Arpa-Sancet, Christophis, Rosenhahn, Biointerphases 2012, 7, 26

Alles, Rosenhahn, Biofouling 2015, 31(5), 469

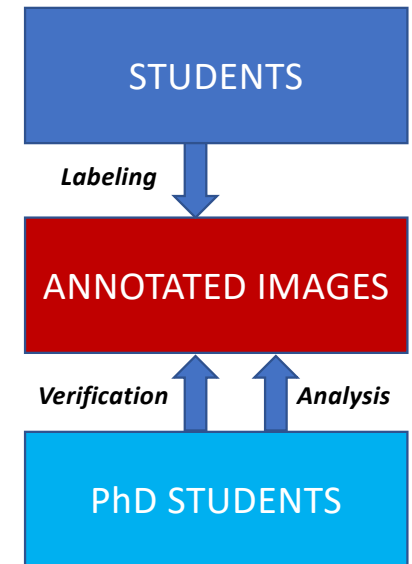
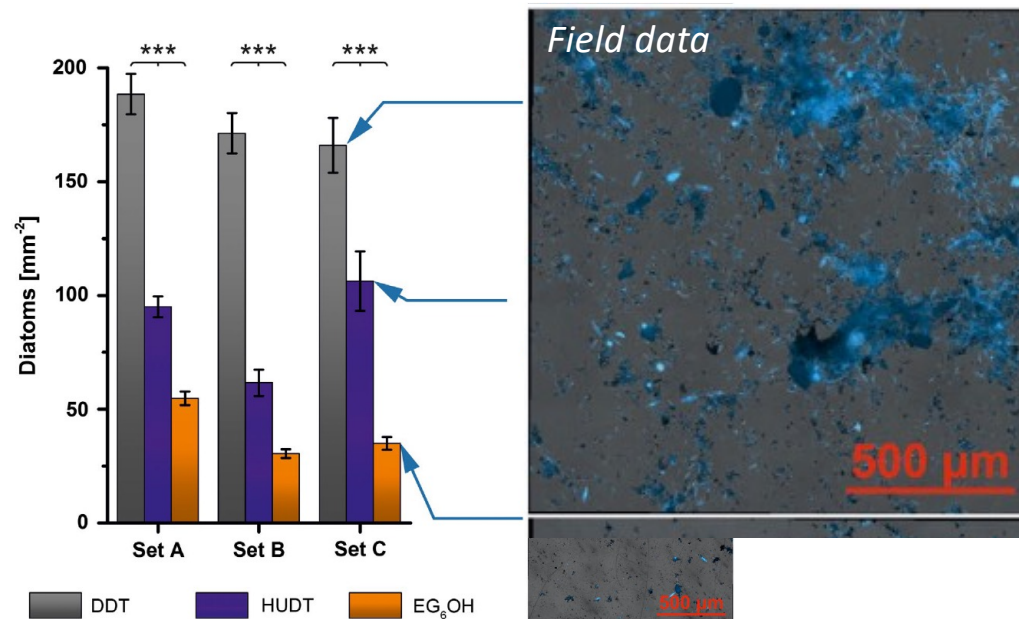
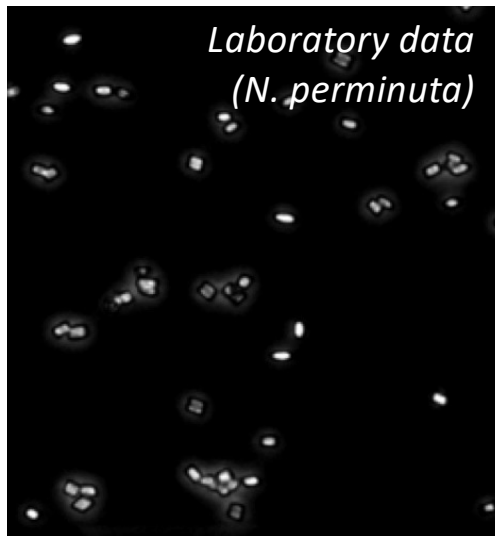
Nolte, Schwarze, Rosenhahn, Biofouling 2017, 33(7), 531

Nolte, Schwarze, Beyer, Özcan, Rosenhahn, Biointerphases 2018, 13(4), 041007

Nolte, Koc, Barros, Hunsucker, Schultz, Swain, Rosenhahn, Biofouling 2018, 34(4), 398

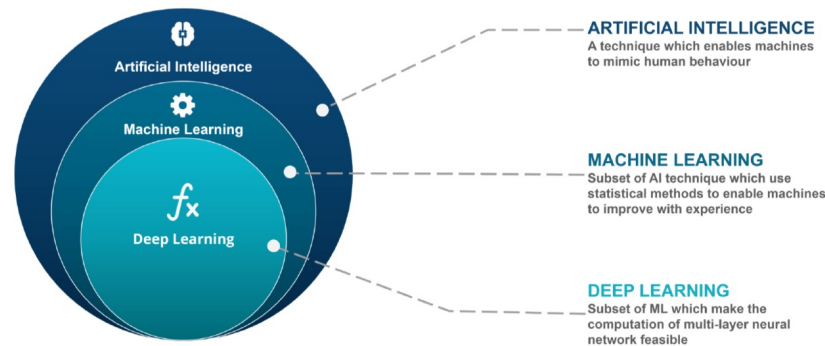
# Analysis of field data

- Counting of diatoms in microfluics works automatically (particle detection)
- Counting of field slides requires manpower as objects on surface are heterogeneous
- Initial DDT/HUdT/EG6 dataset: 4 repl x 9 chem x 60 fov = 2160 Fields of view
- Time for man. analysis per field of view: 5-10 min (depending on coverage, 16 MPix image)  
=> 2160 x 7 min = 252 h of counting (30 days by first and second semester students)
- To be analyzed in 2018/2019: >600 slides, (>36000 FOV  $\approx$  525 days of counting)

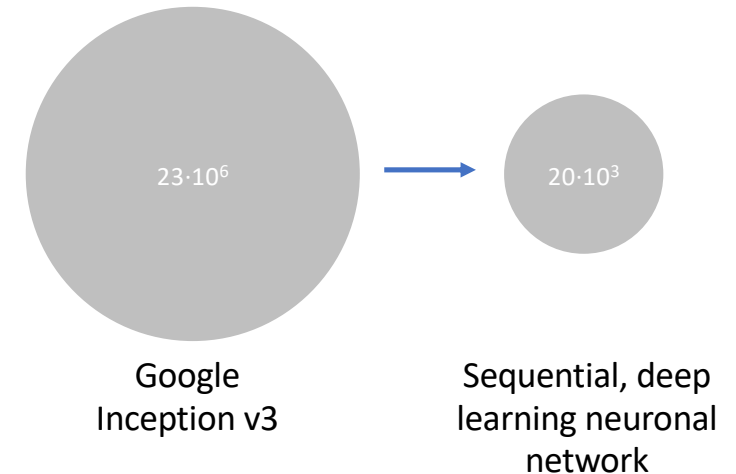


# Deep learning neuronal network

Challenge similar to automated driving



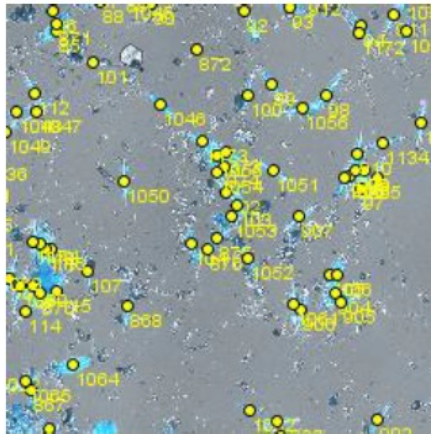
Reduction of free parameters  
(less parameters -> faster analysis)



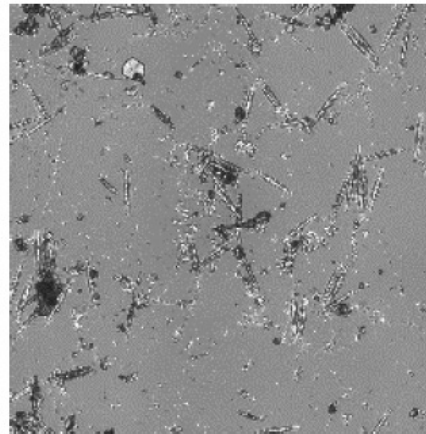
- Goal : Counting diatoms on field samples with particulates and other artifacts present
- Binary classifier -> Not only detection of presence of an object (yes/no) but probability for occurrence in a specified region
- Runs entirely on graphic cards
- Fast analysis ( $\approx 1$ s per image)



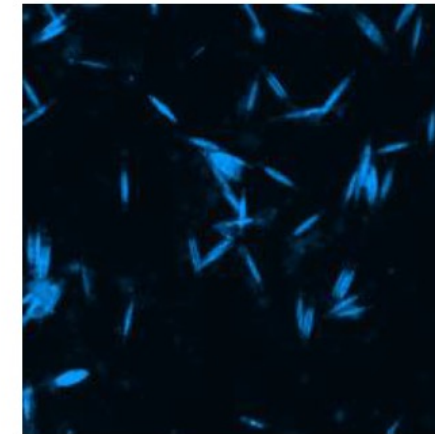
## Microscopy data and learning sets



Coordinates



Phase Contrast

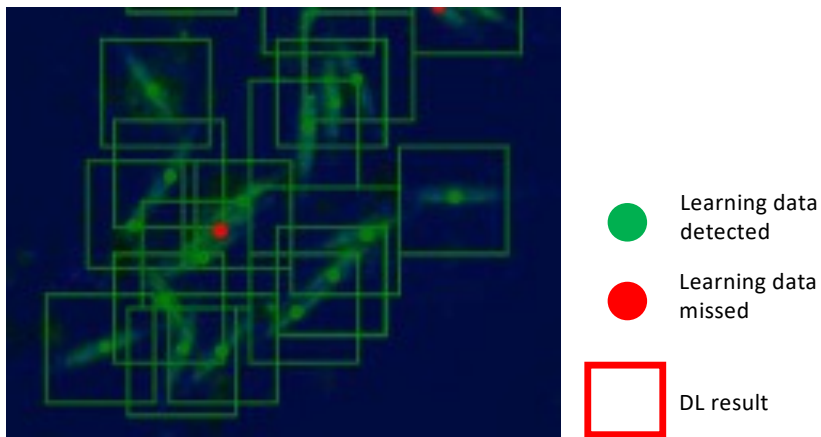
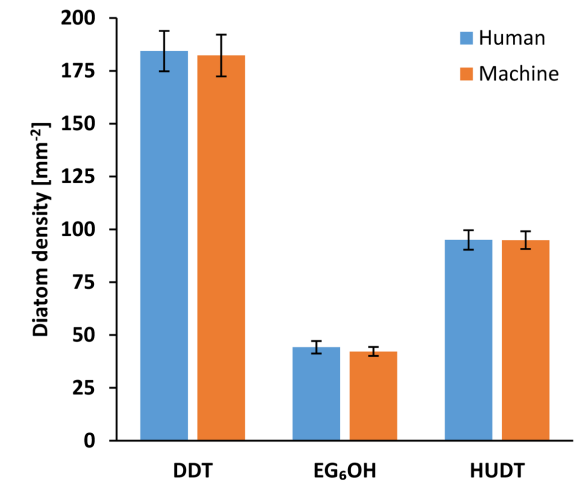
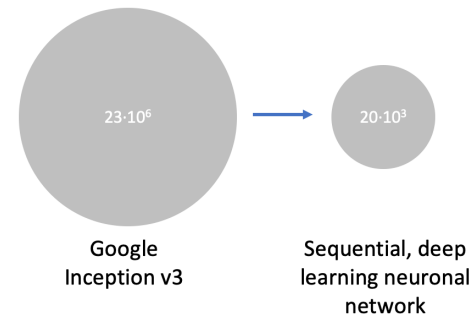
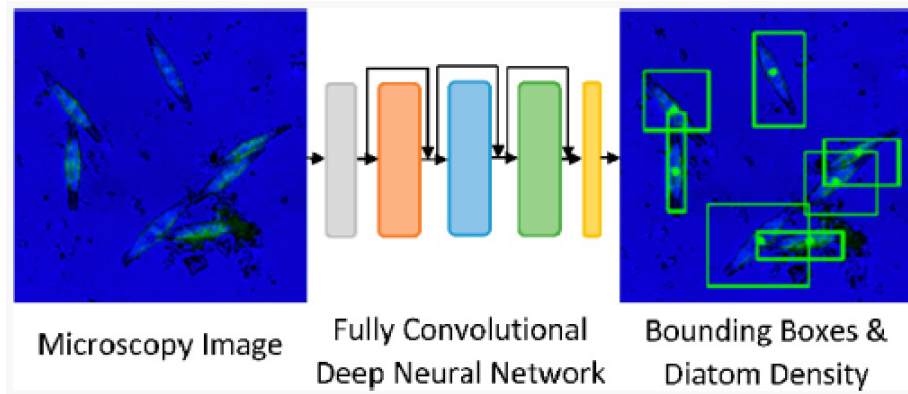


Fluorescence

## Learning dataset :

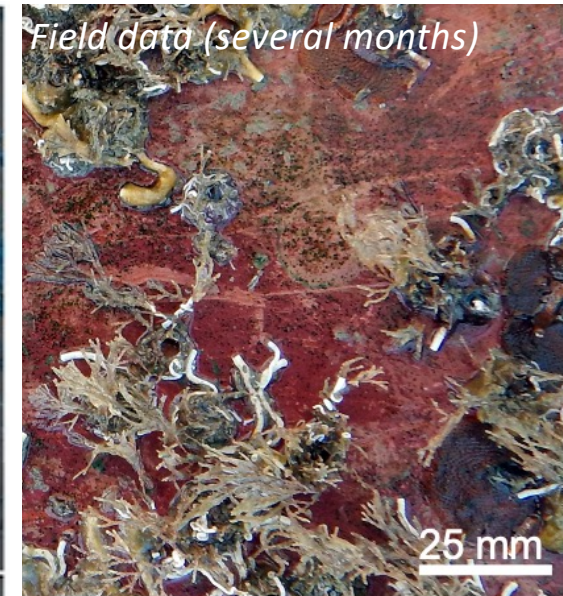
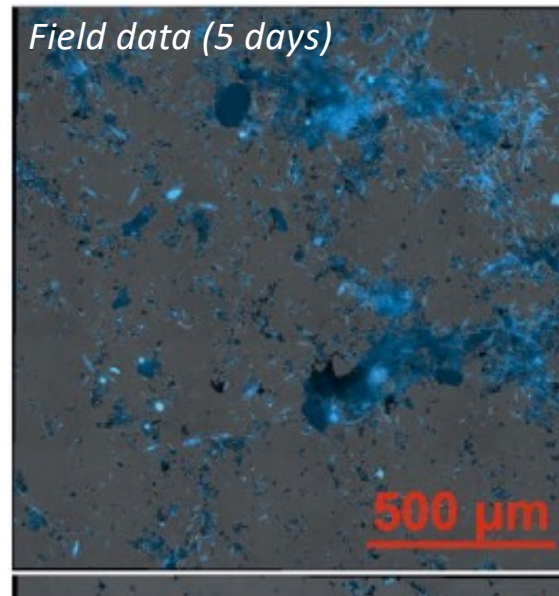
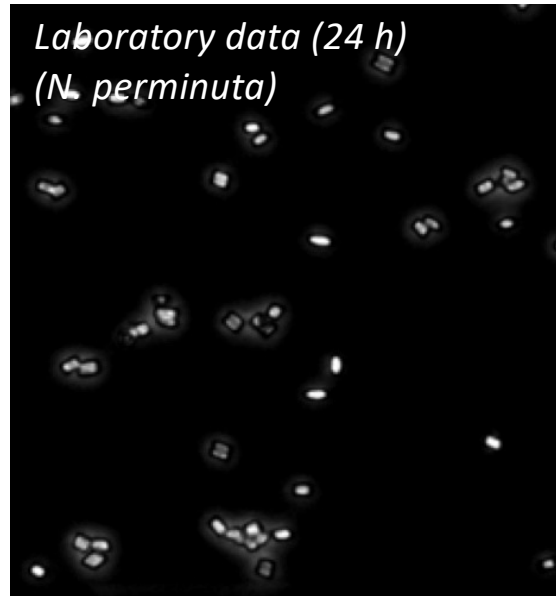
- Initial labeling could not be used due to inaccurate labels (edge of diatom or even next to diatom in close vicinity)
- Diatom positions assigned by different users (different accuracy, required careful selection)
- Expert relabeling with markers in the center of the diatoms were necessary
- Data augmentation (Contrast, brightness, mirroring, rotation)

# Counting of field data with deep learning



- Deep learning counts as good as humans
- Human counting depends on person
- Time required for 600 images was 8 minutes (0.8 s/image) (would take 70 h by human) and DL works non-supervised!

# From image recognition to semantic segmentation



COMPLEXITY



ASTM D6990-20 

## Standard Practice for Evaluating Biofouling Resistance and Physical Performance of Marine Coating Systems

### Significance and Use

5.1 This practice is designed to provide guidance to a panel inspector for quantitative and consistent evaluation of coating performance from test panels coated with marine antifouling coating systems. The practice assesses performance of coating systems based on both antifouling and physical properties.

5.2 The user is cautioned that the results are representative for the specific region and time of year in which the specimens are immersed. It shall be noted that interpretation of results will depend on the geographical location where the test is conducted, whether the coated specimens are exposed either totally or partially immersed, under static or dynamic conditions, and position and orientation.

5.3 Simultaneous testing of a proven standard antifouling coating system (known to minimize fouling accumulation, for example, containing biocide or active agent(s) to prevent fouling settlement/growth) in the specific marine environment shall be included as a reference to assist in interpretation of results. In addition, a negative control (inert surface susceptible to heavy fouling) shall be included on a regular basis. For the exposure to be valid, the surface of the negative control should show heavy fouling relative to the standard system(s).

5.4 Marine coating systems that produce positive results relevant to the standard system(s) show potential for use in protecting underwater marine structures.

5.5 The format can be utilized independent of exposure protocol and coating type, and provides the end user with a consistent practice and format for reporting of performance rating.

- Quantitative and consistent visual evaluation of coatings
- Appropriate consideration of positive and negative controls
- Method should work for a broad range of coating types

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Computers & Geosciences 32 (2006) 1259–1269

[www.elsevier.com/locate/cageo](http://www.elsevier.com/locate/cageo)

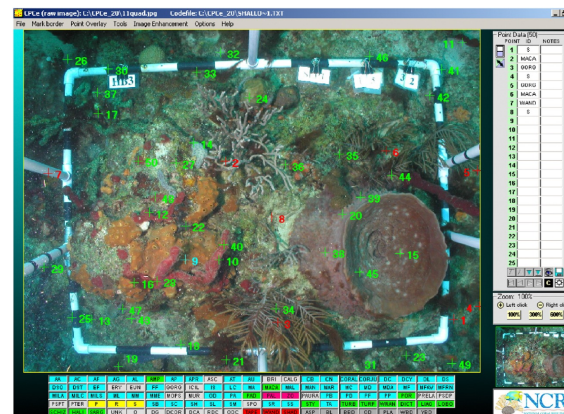
## Coral Point Count with Excel extensions (CPCe): A Visual Basic program for the determination of coral and substrate coverage using random point count methodology☆

Kevin E. Kohler\*, Shaun M. Gill

National Coral Reef Institute, Nova Southeastern University Oceanographic Center, Dania Beach, FL 33004, USA

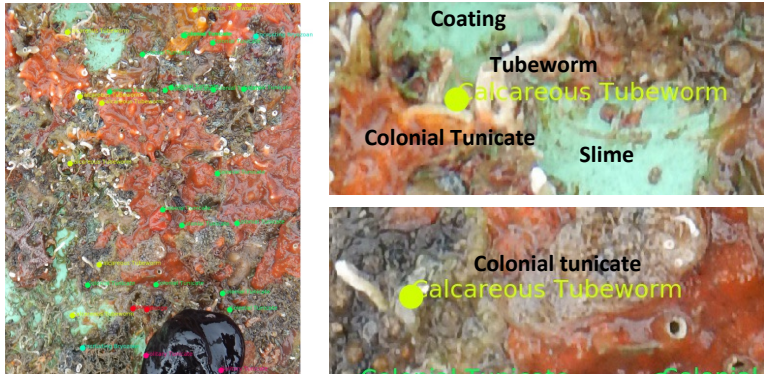
Received 19 July 2005; received in revised form 17 November 2005; accepted 21 November 2005

### Occurrence based (random point annotation)



### Area based (manual segmentation)

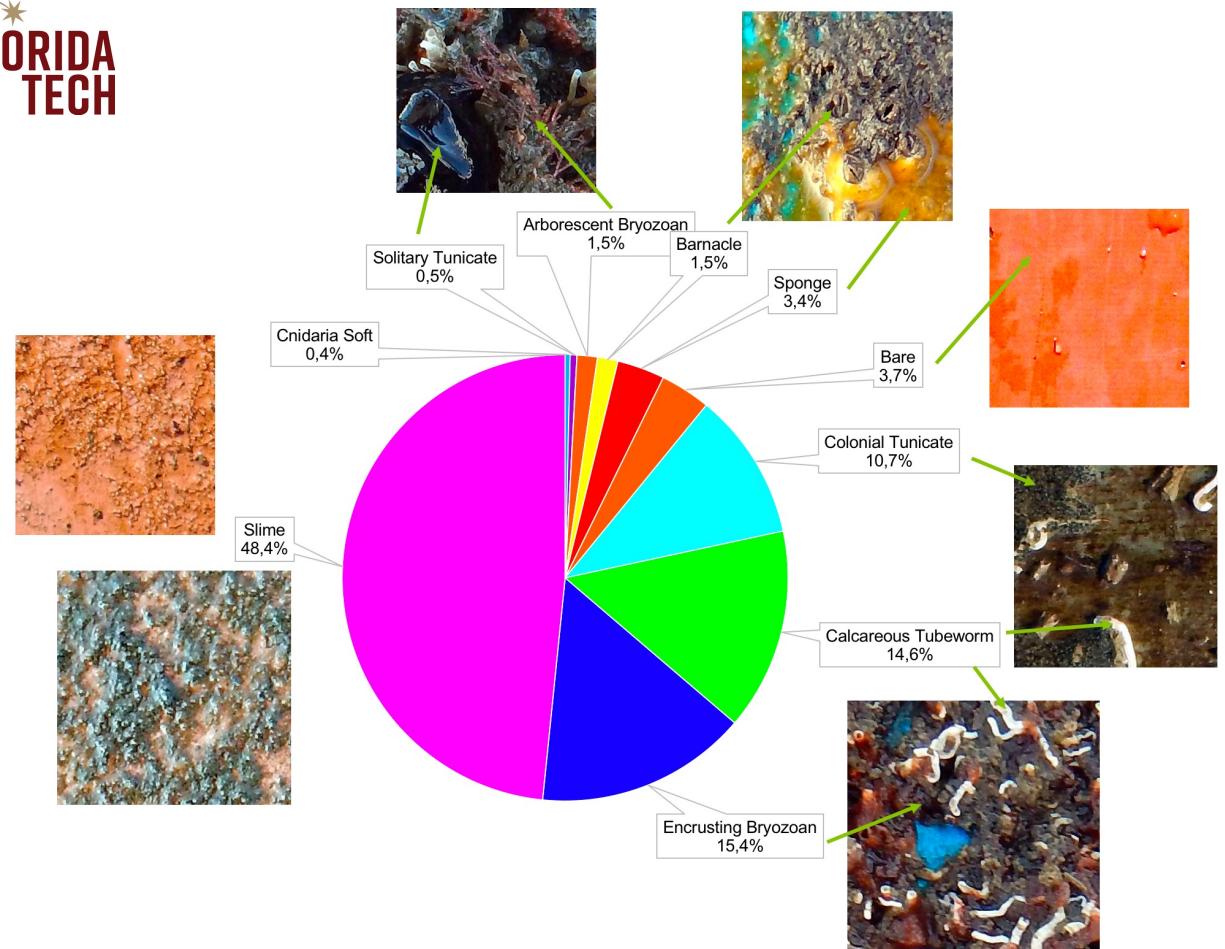




Data base from FIT : 486 panel images  
Point assignment according to CPcE (24300 data points)

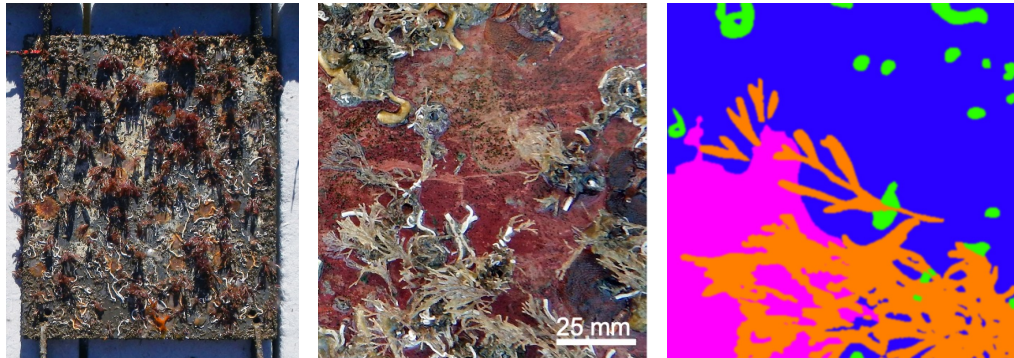
## Challenges for automated image analysis

- Complex composition of fouling organisms
- Mixture of species at certain spots
- Vicinity of randomly chosen CPcE points can be heterogeneous
- Imbalanced panel depending species distribution

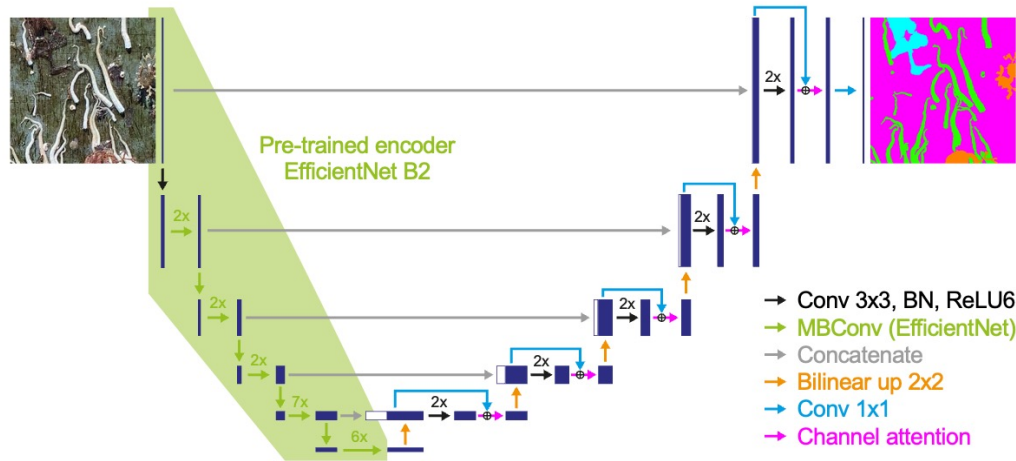




# Unet architecture for semantic segmentation



- Not only object detection and counting but semantic segmentation
- Labeling of entire panels is time demanding and induces a species imbalance
- Separated Panel into patches of  $\approx 5\%$  of the total panel size
- $\approx 300$  randomly chosen fully manually segmented images (13 fully manually segmented panels)



**Supplementary Figure 5: Final model architecture.** A pre-trained EfficientNet B2 encoder is deployed for the contraction path of the U-Net. Arrows denote operations and rectangles represent the resulting tensor. Crossing of arrows represents element-wise addition.

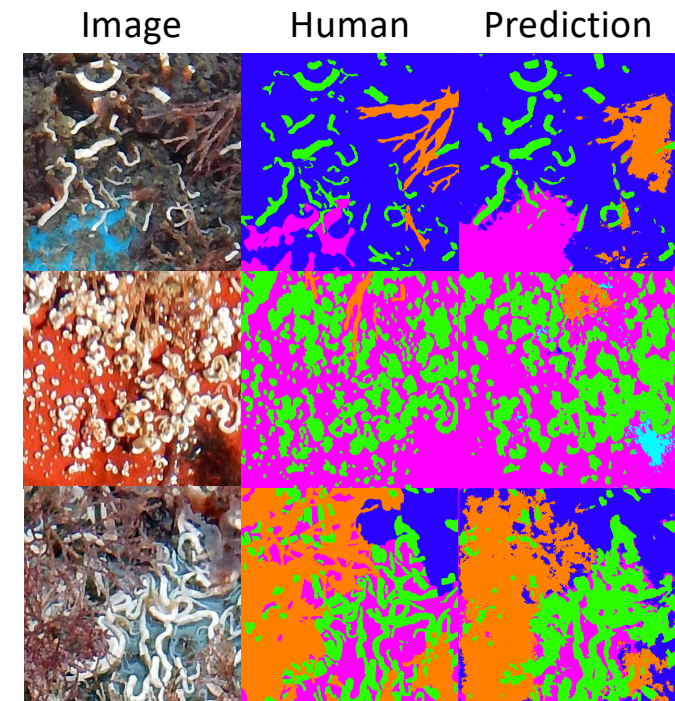
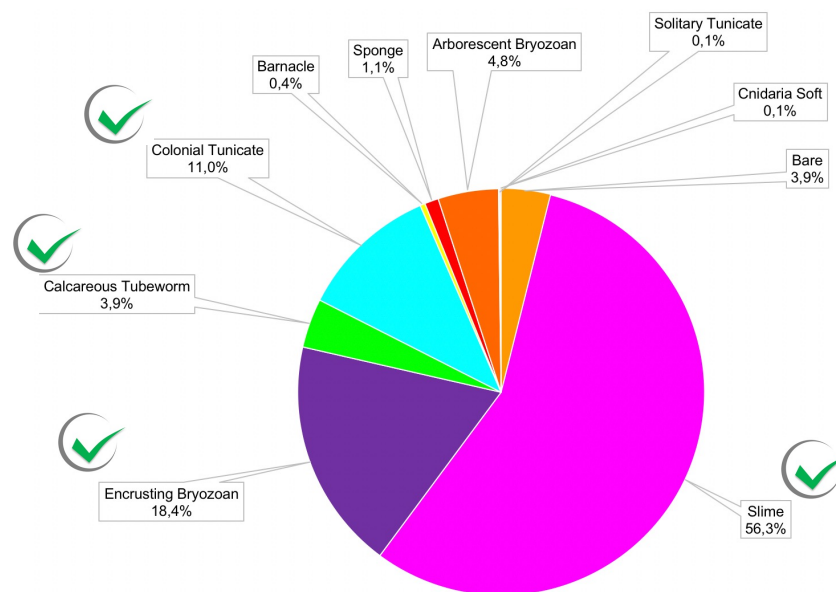
**Table 1: Performance of adapted U-Net model architectures.** Results for image tiles from validation set. Best performing configurations for a metric are highlighted.

Configuration	Accuracy	IoU	F1	Precision	Recall
U-Net (baseline)	0.959	0.614	0.746	0.758	0.739
+ Pretrained <u>EfficientNet</u> encoder	0.974	0.735	0.841	0.842	<b>0.841</b>
+ Residual decoder links	<b>0.976</b>	0.747	<b>0.849</b>	<b>0.864</b>	0.836
+ Channel attention & rebalanced decoder filters	<b>0.976</b>	<b>0.749</b>	<b>0.849</b>	0.861	0.840

# Manually segmented ground truth data

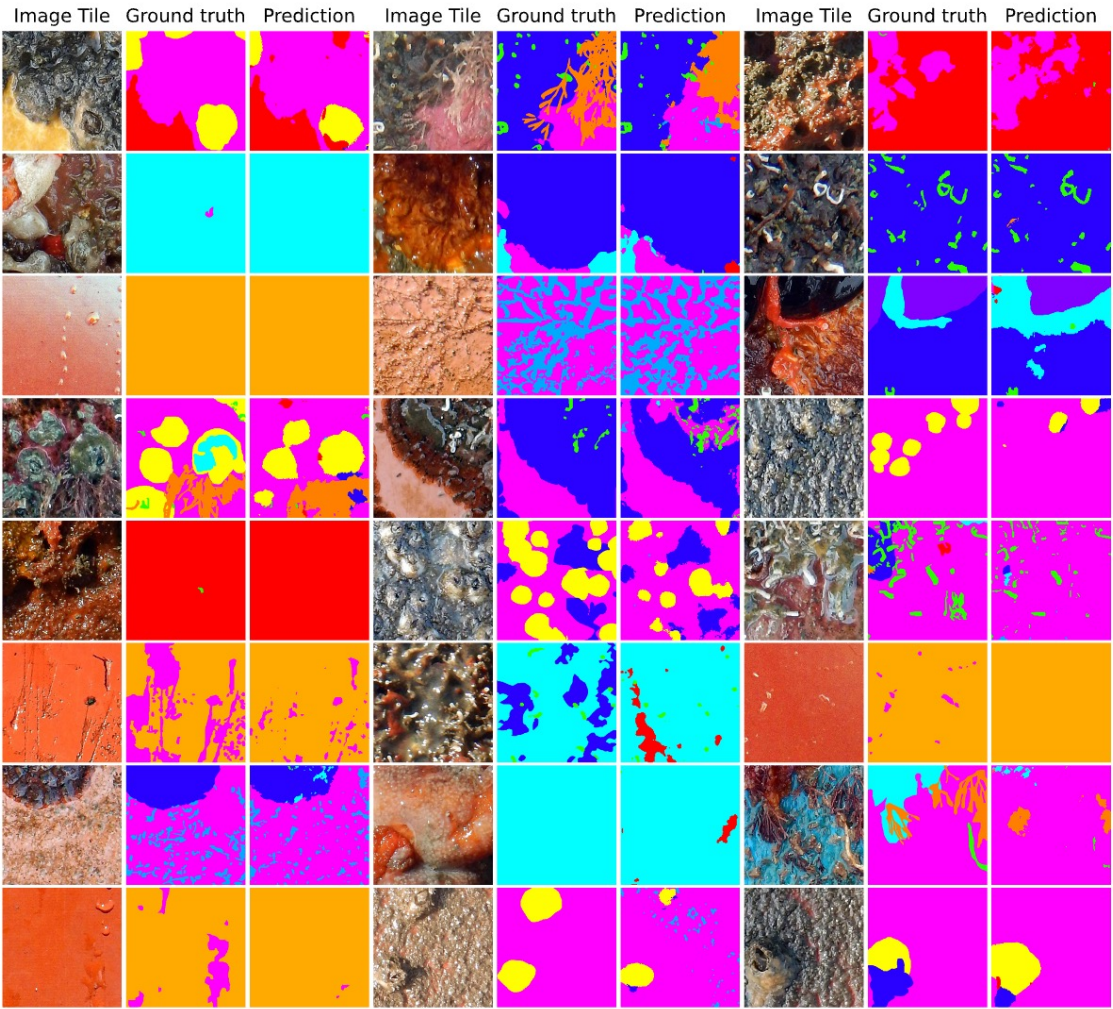
## Status in the middle of the project

- Intersection over union (IOU) of 0.3 for 10 classes
- $\approx 300$  training images BUT high class imbalance
- Architecture optimization and additional labeling of previously unlabeled regions/tiles of high entropy (e.g. bare versus slime as bare is underrepresented)



Supplementary Table 2: Segmentation performance of the enhanced U-Net. Metrics are reported as the class-wise mean of image tiles from the validation set. Best values ( $\geq 85\%$ ) are highlighted.

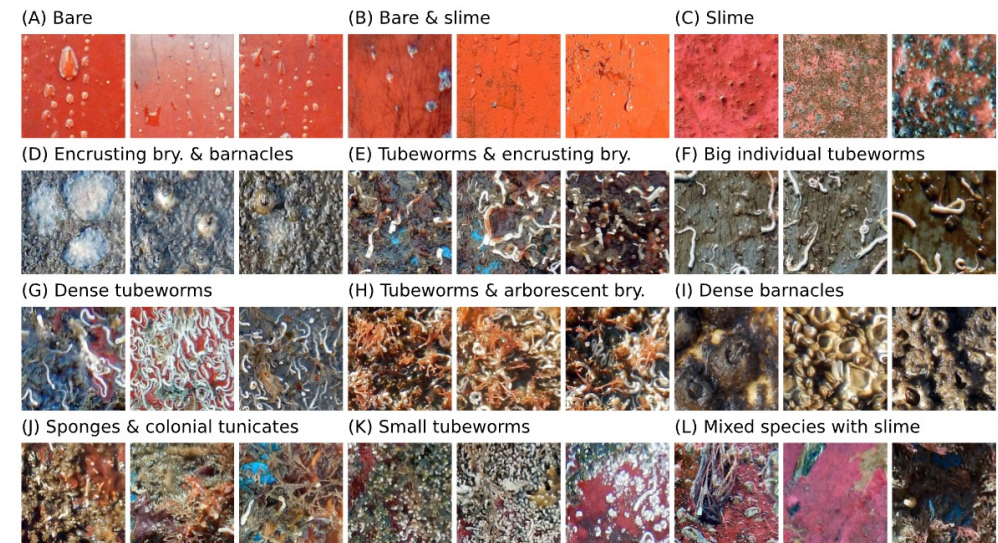
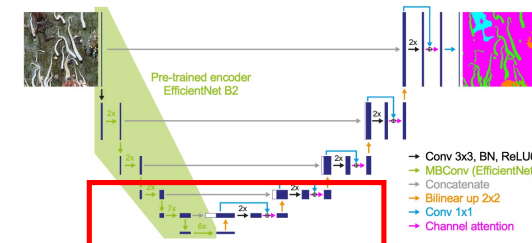
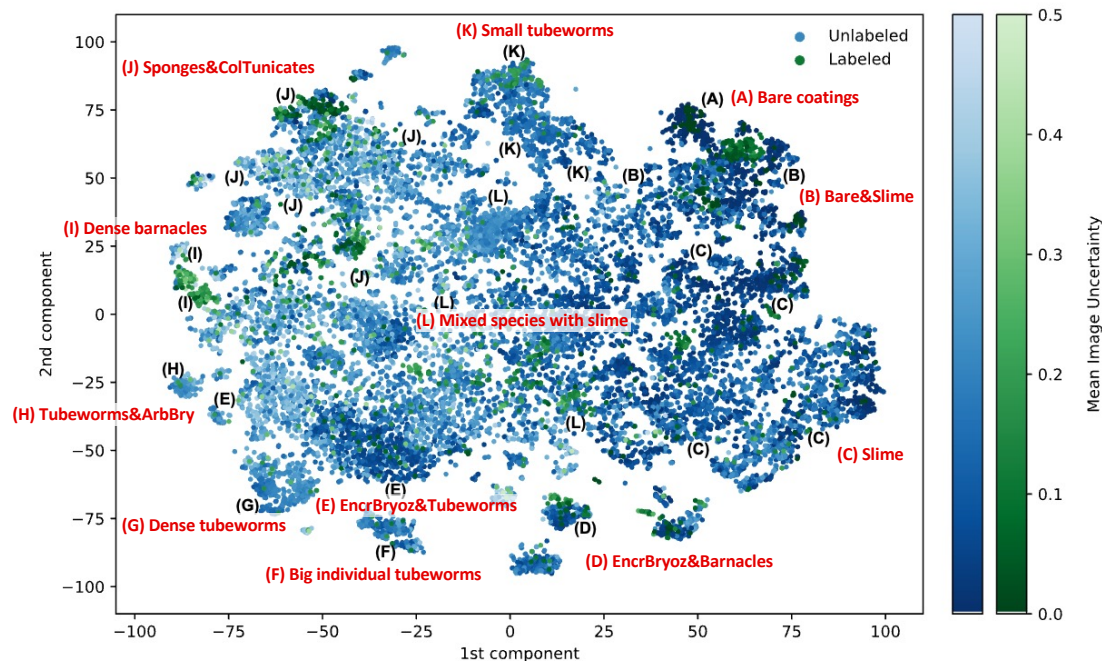
Class	Bare	Slime	Encrusting Bryozoan	Calcareous Tubeworm	Colonial Tunicate	Barnacle	Sponge	Arborescent Bryozoan	Solitary Tunicate	Cnidaria	Mean
Accuracy	<b>0.986</b>	<b>0.911</b>	<b>0.978</b>	<b>0.981</b>	<b>0.969</b>	<b>0.976</b>	<b>0.988</b>	<b>0.982</b>	<b>0.999</b>	<b>0.993</b>	0.976
IoU	<b>0.908</b>	0.807	0.772	0.644	0.667	0.666	0.879	0.505	<b>0.966</b>	0.672	0.749
F1	<b>0.952</b>	<b>0.893</b>	<b>0.872</b>	0.784	0.800	0.800	<b>0.936</b>	0.671	<b>0.983</b>	0.804	0.849
Precision	<b>0.932</b>	<b>0.885</b>	<b>0.861</b>	0.805	<b>0.862</b>	0.801	<b>0.926</b>	0.709	<b>0.974</b>	<b>0.858</b>	0.861
Recall	<b>0.973</b>	<b>0.901</b>	<b>0.882</b>	0.764	0.747	0.799	<b>0.946</b>	0.637	<b>0.991</b>	0.756	0.840





# Semantic segmentation of fouling in the field

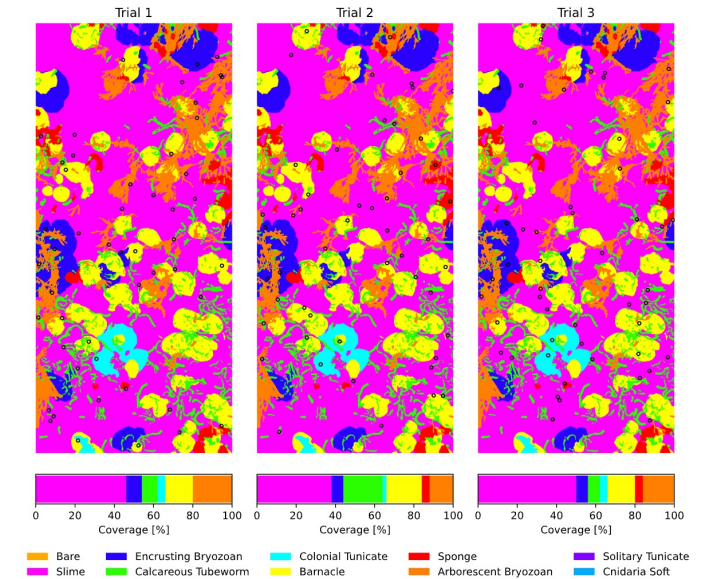
t-SNE (t-distributed stochastic neighbor embedding) of the high-level features extracted by the encoder path of U-Net



# Estimation of sampling error in CPCe annotation

**Supplementary Table 1: Error of random point annotation.** The class-wise random point annotation error was quantified by the mean absolute error (MAE), the mean absolute percentage error (MAPE), and the left-out probability (LOP). Metrics were obtained by uniform sampling of  $n = 50$  random points with segmentation masks from manual annotation and  $N = 1000$  repetitions per image. All values are given as percentage. Reported uncertainties refer to the standard error.

Class	Bare	Slime	Encrusting Bryozoan	Calcareous Tubeworm	Colonial Tunicate	Barnacle	Sponge	Arborescent Bryozoan	Solitary Tunicate	Cnidaria
MAE	$3.94 \pm 0.02$	$3.72 \pm 0.01$	$2.58 \pm 0.01$	$1.55 \pm 0.03$	$2.83 \pm 0.01$	$4.74 \pm 0.00$	$2.17 \pm 0.01$	$2.17 \pm 0.01$	$4.12 \pm 0.01$	$4.40 \pm 0.00$
MAPE	$5.7 \pm 0.0$	$48.0 \pm 0.4$	$72.9 \pm 0.4$	$113.6 \pm 0.5$	$67.3 \pm 0.4$	$20.1 \pm 0.0$	$60.0 \pm 0.5$	$88.1 \pm 0.5$	$33.6 \pm 0.2$	$24.4 \pm 0.0$
LOP	$0.0 \pm 0.0$	$17.4 \pm 0.2$	$27.9 \pm 0.3$	$53.7 \pm 0.3$	$30.0 \pm 0.2$	$0.0 \pm 0.0$	$23.4 \pm 0.2$	$38.4 \pm 0.3$	$6.2 \pm 0.1$	$0.0 \pm 0.0$

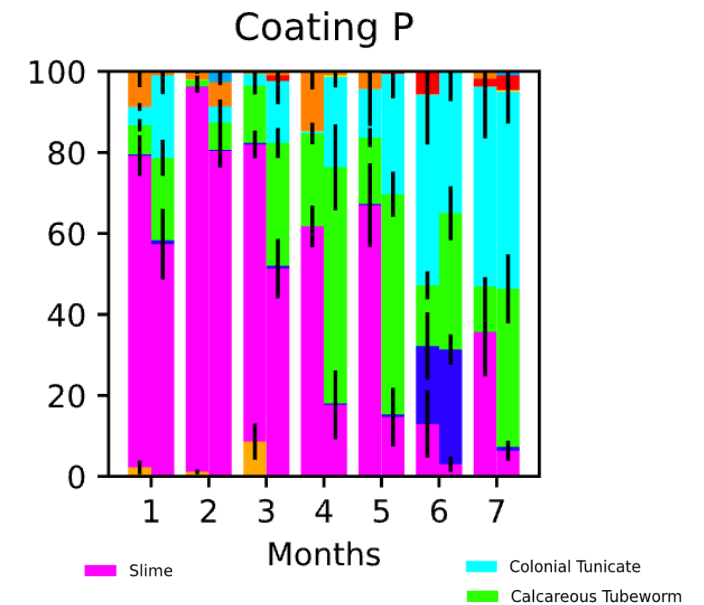
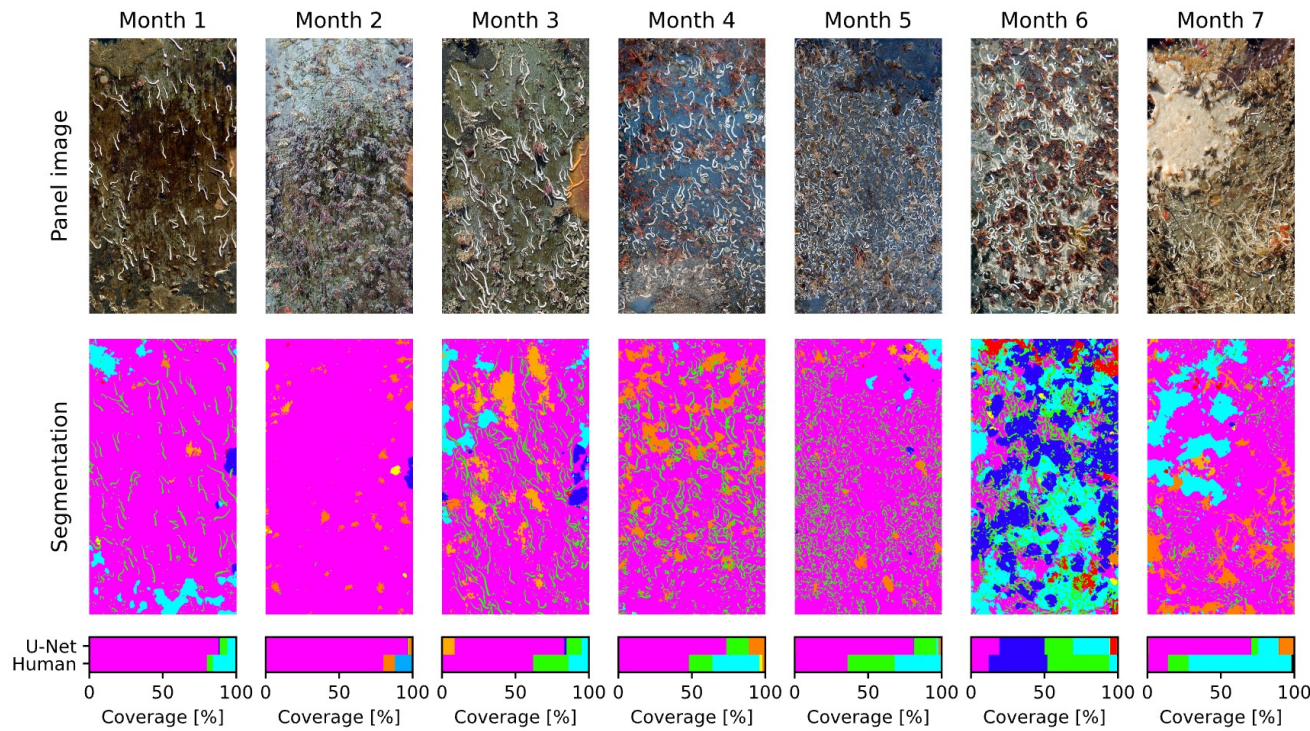


- Fully segmented panel image was used to "simulate" CPCe sampling accuracy
- While frequently occurring species that cover larger areas like bare substrate, barnacles, slime or tunicates has lower left-out probabilities, small objects like tubeworms have a high leaving out probability



# Fouling progression analysis

Criticizm on the method: Bias for top surface layer, but fouling organisms with surface contact relevant

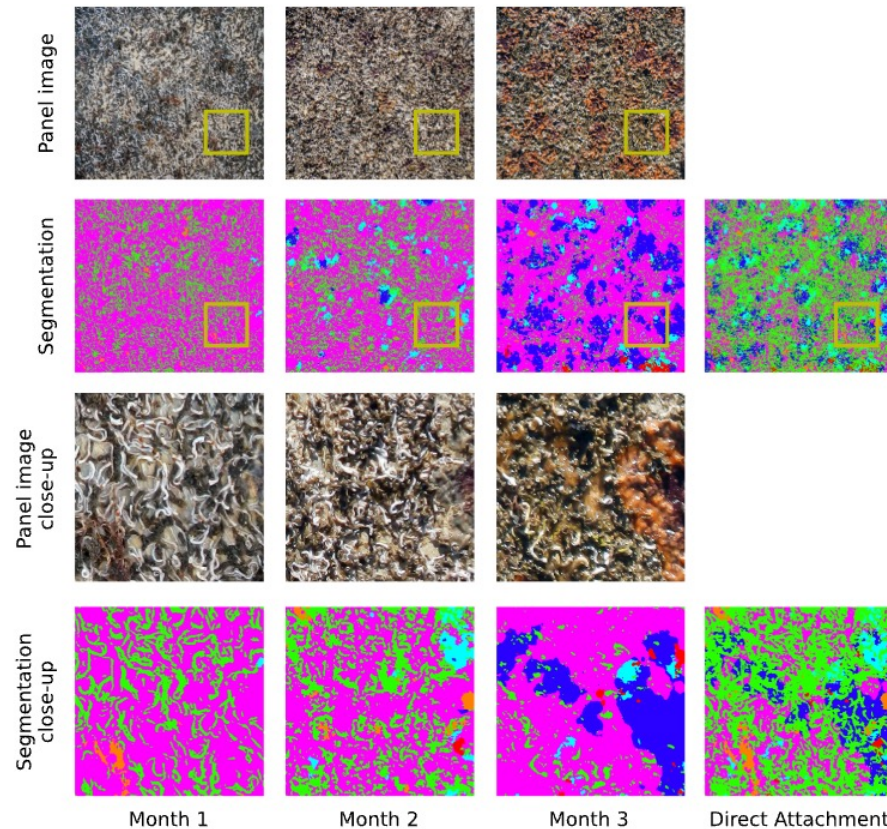


Coverage by organisms is changing over time

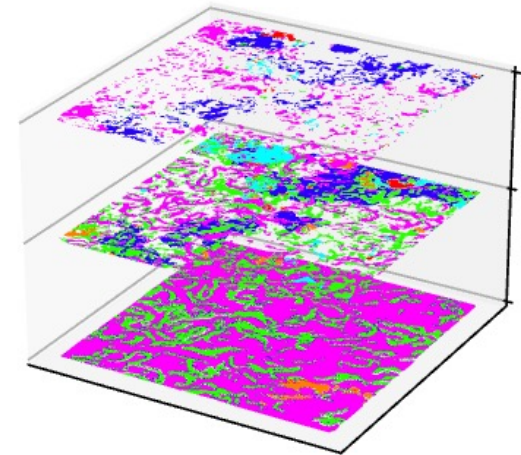
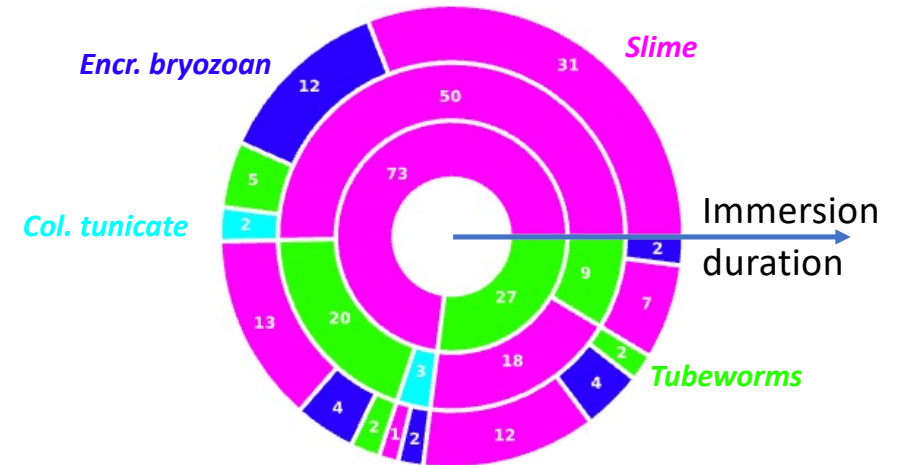
Bars: left U-Net, right human

Bare Encrusting Bryozoan Colonial Tunicate Sponge Solitary Tunicate Others  
 Slime Calcareous Tubeworm Barnacle Arborescent Bryozoan Cnidaria Soft

# Point resolved fouling progression analysis



■ Bare    ■ Encrusting Bryozoan    ■ Colonial Tunicate    ■ Sponge    ■ Solitary Tunicate    ■ Others  
■ Slime    ■ Calcareous Tubeworm    ■ Barnacle    ■ Arborescent Bryozoan    ■ Cnidaria Soft



- Very **early fouling** dominated e.g. by diatoms can be analyzed by image detection using a binary classifier
- Labeling and analysis of **fouling communities** on surfaces can be accomplished by semantic segmentation with deep learning

### Challenges

- Semantic segmentation has a **bias towards the top layer** of fouling. Analysis of time series can help.
- Many of the larger species (e.g. solitary tunicates, arborescent bryozoans) **collapse** if the panels are retrieved from water. Is the assessment accurate? (Similar question applies to CPCe method and to a lesser degree to for ASTM assessment). Does the analysis of underwater (diver) images help?
- If multiple species are grown **symbiotically as new entity**, how should such cases be analyzed/interpreted?
- Need to define how can the data be **reduced** into “heavy fouling”, “light fouling”, and “clean coatings”.

### Advantages

- Once established, it may serve as fast and reliable method for fouling assessment with limited human bias







International congress on marine corrosion and fouling

26.-31.7.26

Henry Ford Building, FU Berlin

Berlin Germany

Organizers:

T. Heusinger von Waldegge, D. Stübing

Fraunhofer IFAM, Bremen

with COIPM committe