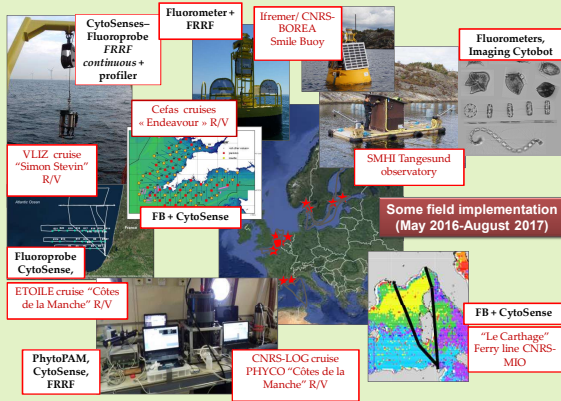


Puillat I.\* (LOPS-Ifrermer, FR); Artigas L. F. & Louchart A (CNRS-LOG, FR); Creach V. (CEFAS, UK); Debusschere E. (VLIZ, B), Rijkeboer M. (RWS, NL); Marrec P. & Thyssen M. (CNRS-MIO, FR); Karlson B. (SMHI, SW) and JERICO-NEXT partners.

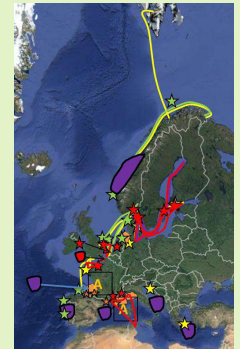
\*Corr. Author: [ipuillat@ifremer.fr](mailto:ipuillat@ifremer.fr)  
[jerico@ifremer.fr](http://jerico@ifremer.fr) & [www.jerico-ri.eu](http://www.jerico-ri.eu)

## Automated platform for the observation of phytoplankton diversity in relation to ecosystem services

A large set of instruments to qualify and expertise: SYKE, SMHI, HZG, RWS, VLIZ, CEFAS, CNRS (LOG, BOREA, OSU-V, MIO) & Ifremer (LER/BL)



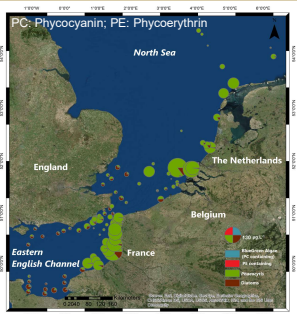
- Pulse-shape recording flow cytometer (Cytosense)
- Imaging in-flow (imaging in-flow Cytobot)
- FlowCAM
- FastCAM
- Spectral fluorometer (AOA, Fluoroprobe, Multiexciter)
- Absorption meter (PSI-CAM)
- Pulse Amplitude Modulated Fluorometers (PAM)
- Fast Repetition Rate Fluorometer (FRRF)
- Underwater Vision Profiler (UVP5)
- (Semi-)Automated data analysis



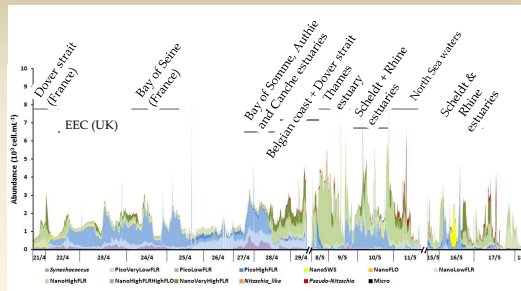
### Outcomes:

A set of recommendations of the most suitable and relevant combination of methods according to the environment considered, their limits and ways of implementing them as complementary sensors in combined platforms. A JERICO-NEXT deliverable to come by one year.

## Phytoplankton groups discriminated by automated pulse-shape recording flow cytometry (Cytobuoy b.v.)



Chlorophyll a equivalent of the multispectral fluorimetry (fluoroprobe bbe Moldaenke): discrimination of 4 pigmentary groups (Haptophytes, brown algae, phycocyanin and Phycoerythrin) containing micro-algae during a *Phaeocystis globosa* bloom.



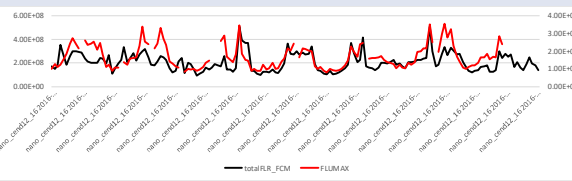
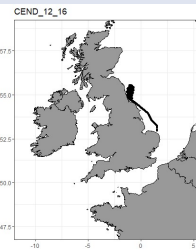
### Up to 13 groups discriminated

Waters under brackish influence: Dominated by nanophytoplankton (and microphytoplankton).

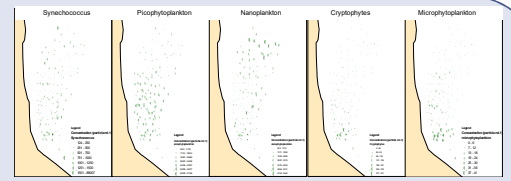
Offshore and other coastal waters: Dominated by pico-eukaryotes, *Synechococcus*-like (and NanoSWS coccolithophores-like)

JERICO-NEXT spring Channel-North Sea cruises (April-May 2017)

EEC-SNS JERICO-NEXT 2017 cruises



Same trend between the total fluorescence from the ferrybox and the total red fluorescence from the flow cytometer (FLR-cytosense) during the survey.



Distribution of the number of particles.ml<sup>-1</sup> according to the phytoplankton functional types

FerryBox: Fluorescence (Fluores), Quantum yield (QY), Salinity (SAL), Temperature (SST)

- ✓ More than 90 % of the particles = phytoplankton, ~ 93% of the total fluorescence.
- ✓ No relationship between salinity, temperature and the number of particles per functional types: *Synechococcus*, picophytoplankton, nanoplankton and microphytoplankton.
- ✓ Difference in the distribution of the phytoplankton functional types with a higher density of nanoplankton in the north of the area.

## Intercomparison exercise: 3 pulse shape recording automated low cytometers (fcm) (VLIZ, RWS and CNRS-LOC) during the JERICO-NEXT – LifeWatch spring cruise 2017 (8–12 May 2017).led by VLIZ

- ✓ Similar trends in measured total cell/colony abundance per ml.
- ✓ Automated fcm: high potential for large scale data collection at high temporal and spatial resolution.
- ✓ Harmonizing the sampling protocol will increase data comparison and greatly improve the insights into phytoplankton community dynamics.

