

JERICO-RI: Progress toward an automated detection of phytoplankton in European coastal areas



Puillat I.* (LOPS-Ifremer, 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: <u>ipuillat@ifremer.fr</u> <u>jerico@ifremer.fr</u> & <u>www.jerico-ri.eu</u>

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 fluorometry (fluoroprobe bbe Moldaenke): discrimination of 4 pigmentary groups (Haptophytes, brown algae, phycocyanin and Phycoerythrin) containing micro-algae during a *Phaeocystis globosa*



Up to 13 groups discriminated

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

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



EEC-SNS JERICO-NEXT 2017 cruises

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







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

Distribution of the number of particles.ml⁻¹ according to the phytoplankton functional types



✓ 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.

FerryBox: Fluorescence (Fluores), Quantum I Difference in the distribution of the phytoplankton functional types with a higher yield (QY), Salinity (SAL), Temperature (SST) 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.

✓ Automatedfcm: 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.



Acknowledgments: The JERICO-NEXT project (www.jerico-ri.eu) has received funding from the European Union's Horizon 2020 research and innovation programme under grand agreement No 654410.

