

# Sea shells record large biases from the marine bomb-<sup>14</sup>C in NW European seawater between the late 1960s and 2019

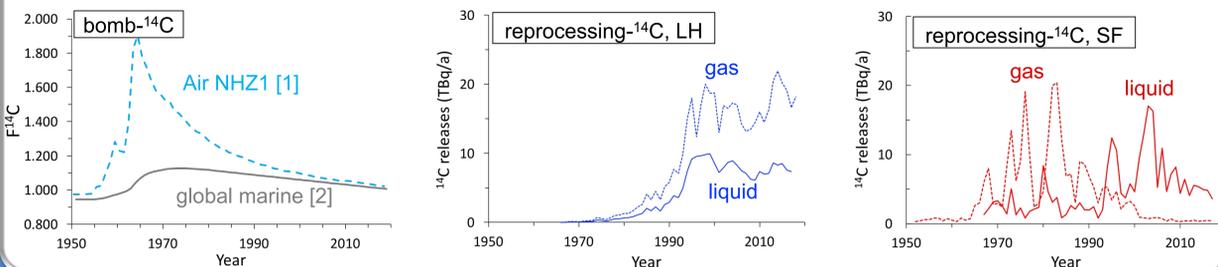
Maxi Castrillejo<sup>1</sup> (maxic@phys.ethz.ch), Christopher A. Richardson<sup>2</sup>, Rob Witbaard<sup>3</sup>, Rob Dekker<sup>3</sup>, Caroline Welte<sup>1,4</sup>, Lukas Wacker<sup>1</sup>, Christiane Yeman<sup>1</sup>, Núria Casacuberta<sup>1</sup>, Hans-Arno Synal<sup>1</sup> and Marcus Christl<sup>1</sup>



<sup>1</sup>Laboratory of Ion Beam Physics (LIP), ETH-Zurich, Switzerland. <sup>2</sup>School of Ocean Sciences, Bangor University, UK. <sup>3</sup>NIOZ and Utrecht University, The Netherlands. <sup>4</sup>Biogeosciences, ETH-Zurich, Switzerland.

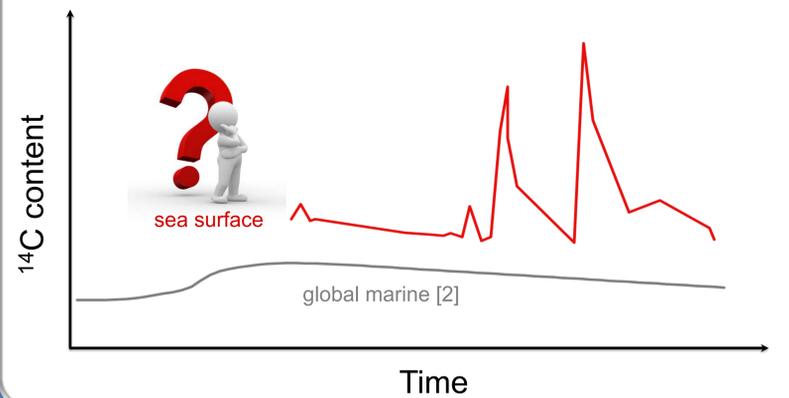
## Abstract

- Accurate knowledge of **radiocarbon (<sup>14</sup>C) content** in the sea surface mixed layer is **valuable for a broad range of applications** in oceanography.
- Globally**, the above-ground nuclear weapon testings (**bomb-<sup>14</sup>C**) in the 1950/60ies **has been the primary source** of increased atmospheric [1] and marine <sup>14</sup>C [2]. **Regionally**, however, other sources can be important. In Europe, the nuclear fuel **reprocessing plants** of La Hague (LH) and Sellafield (SF) **have discharged significant amounts of <sup>14</sup>C** since the 1950s that have been, in large part, documented or reconstructed.
- Previous studies** conducted closeby or downstream of reprocessing plants **found <sup>14</sup>C above bomb levels** in seawater, biota and sediments [e.g. 3-5]. Yet, a comprehensive view on **the temporal evolution of <sup>14</sup>C in sea surface was still missing**. Here, **bivalve shells of known collection date and age are used to reconstruct the <sup>14</sup>C contamination since the late 1960s to 2019**.
- The results show **significant regional deviations from the global and regional marine bomb-<sup>14</sup>C pulse** caused by liquid reprocessing releases that **can have important implications in broad topics including <sup>14</sup>C-dating, tracer oceanography and nuclear forensics**.

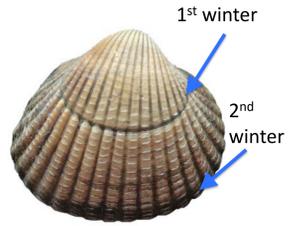


## Research question

What is the impact of nuclear fuel reprocessing on the temporal evolution of <sup>14</sup>C in the sea surface mixed layer?



## Materials and methods



- Study areas nearby or along the path of reprocessing releases: Balgzand (1969-2018) and Traeth Melynog (1975-2019), locations 1 and 2, respectively, in the maps shown below.
- The european common cockle (*C. edule*) bivalve is used as <sup>14</sup>C recorder of surface seawater.
- Cockles were **captured alive** making elaborate age determination unnecessary.
- Individuals with 2 ± 1 years of age were selected** mainly to accurately place data in time. Age was determined by counting of winter lines.

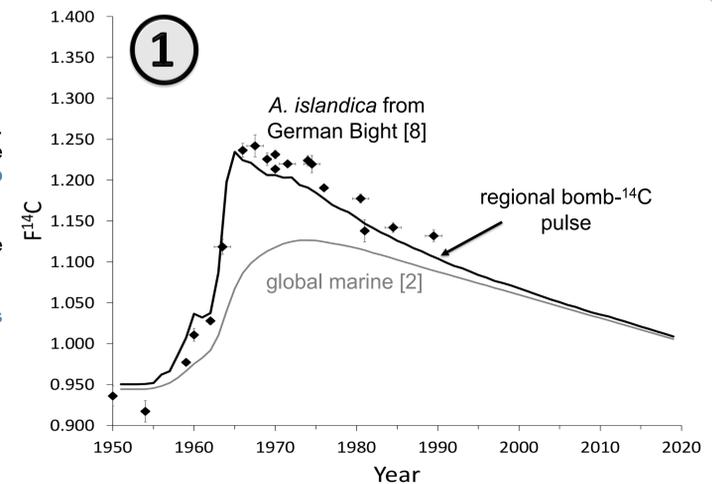
- Bivalve shells were** stored after boiling and removal of the flesh, **cleaned** with deionized water **and ground** to powder size.
- Each sample, of about 1 mg, was prepared following [6].
- <sup>14</sup>C measurement was **done by gas ion source – MICADAS AMS at LIP**.



## Results

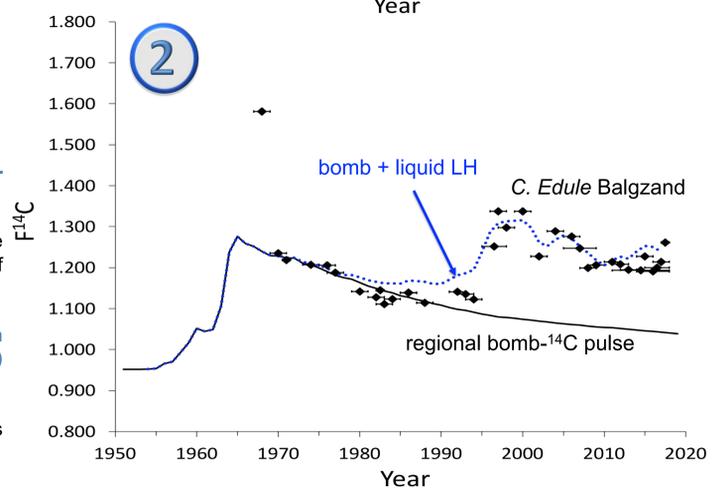
1

- The marine bomb pulse in the surface mixed layer expresses regional differences [7]. Thus, **we used published shell <sup>14</sup>C data of *A. islandica*** [8] that was captured before the large reprocessing releases (see figures in Abstract) **to define a regional bomb <sup>14</sup>C pulse**.
- We found that <sup>14</sup>C displayed by *A. islandica* can be approached by combining 25% of the atmospheric 'Northern Hemisphere 1' [1] and 75% of the Marine09 curve [2].
- The calculated **regional bomb-<sup>14</sup>C pulse** (black line) **was used to identify the excess <sup>14</sup>C** due to other anthropogenic sources, notably the discharge of liquid reprocessing-<sup>14</sup>C.



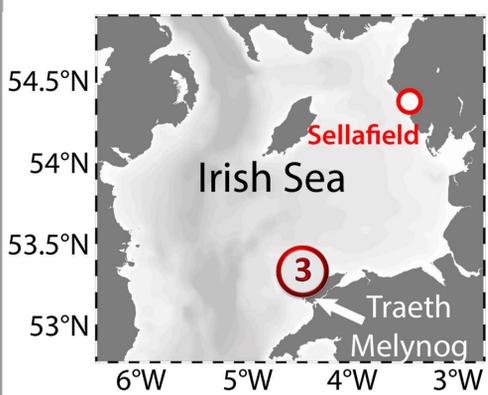
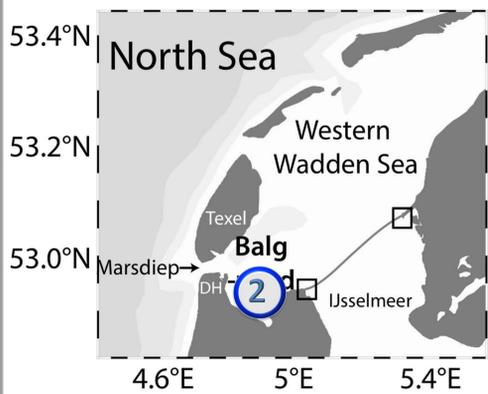
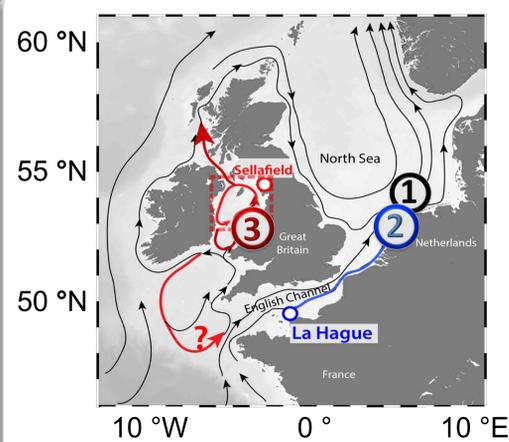
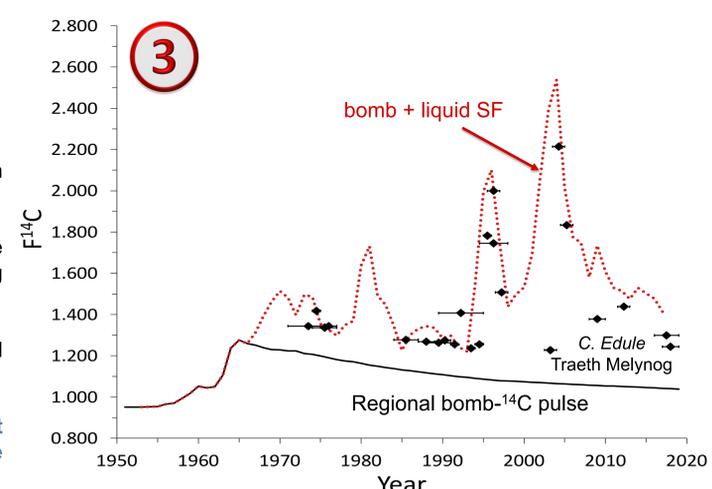
2

- In Balgzand, cockles generally presented  $F^{14}C$  values of 1.100 – 1.300.
- The **shell <sup>14</sup>C content was well above the regional bomb-<sup>14</sup>C pulse after the mid-1990s**.
- Shell <sup>14</sup>C are compared to simulated <sup>14</sup>C values** for the sea surface (blue line). The simulated <sup>14</sup>C reflects the regional bomb pulse with the addition of liquid releases of reprocessing-<sup>14</sup>C from the La Hague plant.
- By applying a multiple linear regression we found that **the combination of both sources explained over 85 % of the temporal evolution (variability and magnitude) of <sup>14</sup>C in the cockles**.
- Results also imply that **other anthropogenic sources** (nuclear power plants, gas reprocessing releases, etc.) **play a minor role** in sea surface levels of <sup>14</sup>C.



3

- In Traeth Melynog, cockles generally present  $F^{14}C$  values of 1.200 – 2.200.
- The **cockle <sup>14</sup>C content showed a greater temporal variability** than in Balgzand.
- In this case, the **shell data are compared to the red dotted line representing the simulated <sup>14</sup>C values** for Irish Sea surface waters. Here, the reprocessing component correspond to the liquid <sup>14</sup>C discharged from the Sellafield plant.
- A **similar regression model** was tested showing that the effect of the liquid reprocessing-<sup>14</sup>C released from Sellafield explained 63% of the variance.
- Overall, both datasets at Balgzand and Traeth Melynog evidenced the impact of reprocessing releases on the observed temporal evolution of <sup>14</sup>C in the surface mixed layer**.



References: 1. Hua *et al.*, Radiocarbon (2013), 2. Reimer *et al.*, Radiocarbon (2009), 3. Cookl *et al.*, J. Env. Rad. (1998) 4. Keogh *et al.*, Radiocarbon (2004), 5. Tierney *et al.*, J. Env. Rad. (2016), 6. Wacker *et al.*, Nucl. Ins. & Meth. Phys. Res. B (2013), 7. Scourse *et al.*, Radiocarbon (2012), 8. Weidman, PhD Dissertation (1995).

### Acknowledgements:

We thank all the scientists involved in the collection of shell material. This work was funded by the ETH Zurich Postdoctoral Fellowship Program (17-2 FEL-30), co-funded by the Marie Curie Actions for People COFUND Program. Additional internal funds were provided by LIP and LIP consortium partners (EAWAG, EMPA and PSI), NIOZ, NERC, BBSRC and Bangor University.

