

## Reply to reviewer #1

### *General comments:*

*The present paper examined the seasonal and annual variations of dissolved N<sub>2</sub>O in a time-series station located in the southwestern Baltic Sea. The results show the coupled variations between the N<sub>2</sub>O anomalies, the oxygen concentrations, and nutrients.*

*The paper presents a valuable new dataset of N<sub>2</sub>O and related biogeochemical parameters in a marine region subject to extensive human activities and so nutrients inputs, responsible of the deoxygenation in the Baltic Sea. After the revision, I consider that the manuscript is highly interesting and provide relevant information about processes occurring to the N<sub>2</sub>O in the Boknis Eck. The paper is well written and structured, with an appropriate description of the state of the art, objectives are clearly outlined and discussion precisely referenced. The main strength of the paper is the monthly sampling undertaken during twelve years.*

*However, there are several weaknesses in the paper. First, the authors make the discussion of the results based on data that are included in this study. There are references to data that exist but are not shown. But, in order to discuss about upwelling and hydrographic changes, about algal blooms and ammonium changes, the salinity, temperature, chlorophyll and ammonium data should be included and shown in this paper.*

Both reviewers requested to show additional temperature, salinity, chlorophyll *a* and ammonium data. These data were not explicitly shown in our manuscript because they have been published already in Lennartz et al.: Long-term trends at the Boknis Eck time series station (Baltic Sea), 1957–2013: does climate change counteract the decline in eutrophication? Biogeosciences, 11, 6323–6339, <https://doi.org/10.5194/bg-11-6323-2014>. However, in order to provide this obvious lacking information, we decided to include the seasonal and annual variations of temperature, salinity, chlorophyll *a* and ammonium in a new supplement to our manuscript.

*Secondly, the paper lack of a proper description of the water masses presents at BE and their temporal variability.*

Considering that all the measurements were conducted at one fixed location, it is difficult to investigate the water masses based on our data. However, the hydrological condition at BE are not complicated. As is written in Lines 86-87, there is no pronounced river input, and saline water from the North Sea plays a dominant role. By showing the seasonal and annual variations of temperature, salinity, dissolved oxygen and other parameters, readers gain a comprehensive idea about the hydrographic conditions at BE during 2005-2017.

We rewrite the lines 86-87 which read now:

There is no significant river runoff to Eckernförde Bay. Hence, the hydrographical conditions are mainly dominated by saline water input from the North Sea and less saline water from the Baltic Proper, which is typical for that region.

### *Specific comments:*

-Lines 129-130: *How did you shifted the data to the 15th? Include procedure and assumptions in the text.*

Since the time of sampling varied every month (usually 20-40 days interval), it would be easier for comparison and data analysis if all the samples were collected with a regular spacing. In this case, we ignored the slight time difference and assumed that all of the samples were collected on the same day every month.

We have replaced lines 129-130 with the following sentence: Sampling time varied for every month (usually 20-40 day interval), but for the statistical analysis, data was assumed to be regularly spaced as differences on weekly scales were minor.

*Line 170: Could you explicitly explain in the text how did you computed Sc, instead just give the reference? What is the equation for computing Sc?*

Sc was computed as:

$$Sc = v/D_{N2O}$$

$$D_{N2O} = 3.16 \times 10^{-6} e^{-18370/RT}$$

where  $v$  is the kinematic viscosity of seawater, which is calculated from the empirical equations given in Siedler and Peters (1986), and  $D_{N2O}$  is the diffusion coefficient of N<sub>2</sub>O in seawater.  $R$  is the universal gas constant and  $T$  is the water temperature in K. We used the  $D_{N2O}$  from Rhee (2000).

We have incorporated this part into section 2.4.

*Lines 176-184: The comparison of the range of concentrations found between Boknis and other time-series would be better move later in the text, since the reader at this point does not have enough information about the causes that differentiate it from other time-series. The authors should better discuss not only the different magnitudes of the N<sub>2</sub>O concentrations, but also the site-specific processes responsible of such differences.*

Thank you for the suggestion. The purpose of the comparison is just to give a general idea about the values of the few time-series N<sub>2</sub>O measurement published so far, because it might be different from the normal cruises which only last for days or months. A comparison of N<sub>2</sub>O concentrations between different time-series analysis is not the major topic of the manuscript, and a discussion about the site-specific processes requires detailed information on the environmental variability of the time-series stations, which does not fit in the scope of the manuscript. In this case, we keep this part in section 3.1.

*Lines 207...: In case there is additional information in the BTS, such as chlorophyll, during the study period, show the data in figures instead to refer to previous studies.*

Chlorophyll *a* data were added and are now shown in a new supplement.

*Lines 235: are there NH<sub>4</sub> data available at the study site during the study period? In that case, it would be better to show them for the discussion instead to appeal to a reference NH<sub>4</sub><sup>+</sup> data were added and are now shown in a new supplement.*

*Lines 238-239: "Denitrification is inhibited by the presence of O<sub>2</sub> and thus nitrification is presumably responsible for the high N<sub>2</sub>O concentrations in winter/early spring." This statement is not correct at all. The production of N<sub>2</sub>O by denitrification can occur at suboxic and hypoxic environments. Please, modify this sentence.*

Thank you for pointing out the problem. We modified the text to “Denitrification is inhibited by the presence of high concentrations of dissolved O<sub>2</sub> (> 20 µmol L<sup>-1</sup>) and...”

*Line 239-240: The authors should normalize the N<sub>2</sub>O and pH to a constant temperature. Otherwise, temperature changes can be the responsible of this relationship because of thermodynamics changes and not necessarily due to nitrification. In fact, it is not as clear the positive correlation between the N<sub>2</sub>O and pH in figure 5, since for pH higher than 7.6, there is no apparent trend between N<sub>2</sub>O and pH. The relationship between pH and N<sub>2</sub>O obtained during incubations experiments described by Rees et al. (2016) cannot be directly compare to this study, since the experimental conditions and approaches are completely different. The authors should rewrite the entire paragraph.*

We need to delete this part from the manuscript because after double-checking the data, we realize that some of the pH values were not calibrated properly. After re-calibration, the relationship between N<sub>2</sub>O and pH no longer exists. We are very sorry about the mistake.

*Lines 263-269: Again, the temperature salinity and Chla information at Boknis are mentioned in the text, but data are missed. If data for these parameters exist, the authors should include in the manuscript. It would reinforce some of the statement that now could look only speculative.*

Temperature, salinity and Chlorophyll *a* data were added and are now shown in a new supplement.

*Lines 287-288: “Although the observed temperatures and salinities during October 2016–April 2017 were comparable to other years,..”. Please, show temperature and salinity.*

Data is now shown in the supplement (see replies above).

*Lines 295-296: “Considering the classical view of N<sub>2</sub>O consumption via denitrification under hypoxic and anoxic conditions”. This is contrary to the statement done at lines 238-239. Consider to rewrite the first one.*

Thank you pointing this out. We have revised the first one as suggested.

*Lines 304-306; 308-309: The authors should make use of temperature, salinity or density to show changes in water masses.*

Temperature and salinity data can be found in the supplement. Mixed layer variations can be seen in Fig. 4. As is mentioned above, it is difficult to show the changes in water masses. We suggested that the low-N<sub>2</sub>O water was a result of advection because vertical exchange can be excluded. However, we do not have any evidence since we did not measure dissolved N<sub>2</sub>O from adjacent waters.

*Lines 313: Instead of “presence” it would be more correct “concentration/level”*  
We have revised it as suggested.

*Lines 320: “We did not observe an exceptional spring algae bloom in 2017”. Please, consider to include Chla or POM to support this statement.*

Chlorophyll a data is now shown in the supplement (see replies above). Unfortunately, POM data are not available. Secchi depth, a proxy of water transparency, is slightly lower in March 2017 than the average value. This could be used to support the statement.

We modified the text to “Secchi depth, a proxy of water transparency, was 3.8 m in March 2017, which is only slightly lower compared to the monthly average value for March ( $4.5 \pm 1.8$  m). There was no exceptional spring algae bloom and thus we infer that assimilative uptake of nutrients by phytoplankton was not responsible for the low nutrients concentrations.”

*Lines 319: Why can not be shown the Chla data?*

Chlorophyll *a* data is now shown in the supplement.

*Lines 331-335: The author should also discuss the potential dependence of rates on temperature and its impact on the seasonal variations of N<sub>2</sub>O production/consumption through the text.*

Unfortunately, we did not measure N<sub>2</sub>O production/consumption rates at BE. A discussion about the potential temperature dependence of rates is, thus, too speculative. Besides, there is no significant temperature anomaly during the low-N<sub>2</sub>O-event. In this case, we suggest that the impact of temperature on the low-N<sub>2</sub>O event could be excluded.

*Lines 356-357: Please, consider to support this statement with the salinity data 371-373. Please, show the density (or temperature and salinity) record to track the upwelling event in autumn 2017. Lines 377-378: Please, show the chlorophyll data. Lines 385-386: Please, show the ammonium data.*

The data are now shown in the supplement.

*Lines 394-399: This is a very speculative paragraph as it is written. Could you give any evidence for these potential explanations of the homogeneous distribution of N<sub>2</sub>O?*

Although the oxic/anoxic interface, where enhanced N<sub>2</sub>O production occurs, lasts for several months, the high N<sub>2</sub>O concentrations were usually observed only in late autumn (Fig. 4). We agree this paragraph is speculative. There are just some “potential explanations” for why there is no enhanced N<sub>2</sub>O in early autumn. Unfortunately, we do not have any further evidence to support the conjecture.

*-Section 3.5: The author should evaluate in the results the impact of the dissolved gas analysis uncertainty in the air-sea flux computation and the uncertainty introduced in the net seasonal and annual air-sea NO fluxes.*

The uncertainty of flux density, which is mainly derived from K<sub>N<sub>2</sub>O</sub>, with a minor contribution from the error of trace gas analysis, was estimated to be 20% (Wanninkhof, 2014). The average flux density at BE was  $3.5 \pm 12.4$   $\mu\text{mol m}^{-2} \text{ d}^{-1}$ . With a large uncertainty in the flux density, it is difficult to compute meaningful seasonal/annual fluxes. In this case, we only discussed the variation of flux density in section 3.5.

We have added the uncertainty of flux density in section 3.5.

*Lines 416-424: The authors show that N<sub>2</sub>O concentration change seasonally, but the saturation stay almost constant. So, how can the author affirm that emissions are controlled by temperature?*

There is a seasonality in surface N<sub>2</sub>O concentrations but not for the N<sub>2</sub>O saturation. During the transformation of concentrations into saturations, the effect of temperature on the saturation is more essential than the effect of salinity. In summer when surface N<sub>2</sub>O concentrations are low, N<sub>2</sub>O saturations are increased by the relative high temperature (because the equilibrium concentration is decreased). In winter the N<sub>2</sub>O concentrations are high, but N<sub>2</sub>O saturations are decreased because of the high N<sub>2</sub>O solubility at low temperature condition. Temperature is “buffering” the variation of saturation and thus affecting N<sub>2</sub>O emissions, and this is our point of “a modulating role”.

To clarify this point we rewrite the paragraph:

We found a weak seasonal cycle for surface N<sub>2</sub>O concentrations, with high N<sub>2</sub>O concentrations occurring in winter/early spring and low concentrations occurring in summer/autumn, but no such cycle for N<sub>2</sub>O saturation. The seasonality in concentration but not in saturation could be largely attributed to the effect of temperature on N<sub>2</sub>O solubility: In summer when surface N<sub>2</sub>O concentrations are low, N<sub>2</sub>O saturations are increased by the relative high temperature; and vice versa in winter. Although salinity also affects N<sub>2</sub>O solubility, its contribution is negligible compared to temperature. Temperature alleviated the fluctuation of surface N<sub>2</sub>O saturation and thus affected the sea-to-air N<sub>2</sub>O fluxes. We conclude that temperature plays a modulating role for N<sub>2</sub>O emissions.

*Lines 476-484: Unless the author do not include salinity and temperature, they should not used them to conclude the hydrographic conditions at Boknis Eck. Further studies about the hydrography at the BE would complete the picture together with the biogeochemical data at the BE time-series station.*

We agree that changing hydrographic conditions will affect N<sub>2</sub>O cycling at Boknis Eck as well. This, however, will require modeling studies which need to take into account the ongoing environmental changes of temperature, deoxygenation, changing frequency of North Sea water inflow etc. We think that a detailed discussion of potential future projections of the environmental variability of Boknis Eck/Eckernförde Bay is beyond the scope of the manuscript.

## Reply to reviewer #2

*Quantifying the concentrations and dynamics of dissolved N<sub>2</sub>O in seawater is important for understanding the climate change, but conducting measurements of sufficient duration to determine trends over seasonal, interannual, and decadal time frames for any marine ecosystem remains a challenging task. A long-term Time-Series Station like Boknis Eck in the Eckernförde Bay can provide invaluable information for documenting the role of oceans in relation to N<sub>2</sub>O, hence this type of study is significant for our scientific understanding. The paper is well-written and clear. I only have few comments/suggestions below:*

*Lines 95-96: more information is needed on seawater sample collection.*

The sampling procedure is described in detail in lines 95-100 on page 3. Thus we do not see a need to revise the text.

*Lines 108-109: The N<sub>2</sub>O concentration of standard gases should be provided.*

We have added this information in the method section.

*Line 154-155: : : :dry mole fractions of atmospheric N<sub>2</sub>O at the time of the sampling. This description is not the fact since atmospheric N<sub>2</sub>O at the time of the sampling has not been measured and the monthly average of N<sub>2</sub>O data measured at Mace Head was used to N<sub>2</sub>Oeq.*

Thank you for pointing this out. We modified the sentence. It reads now: “Since the atmospheric N<sub>2</sub>O mole fractions were not measured at the BE Time-Series Station, atmospheric dry mole fractions of N<sub>2</sub>O were derived from the monthly average of N<sub>2</sub>O data at Mace Head, Ireland (AGAGE, <http://agage.mit.edu/>), instead.”

*Lines 185-193: NH<sub>4</sub><sup>+</sup> concentrations should be provided in the text as well as Figure 2.*

Both reviewers requested to show additional temperature, salinity, chlorophyll *a* and ammonium data. These data were not explicitly shown in our manuscript because they have been published already in Lennartz et al.: Long-term trends at the Boknis Eck time series station (Baltic Sea), 1957–2013: does climate change counteract the decline in eutrophication? Biogeosciences, 11, 6323–6339, <https://doi.org/10.5194/bg-11-6323-2014>. However, in order to provide this obvious lacking information, we decided to include the seasonal and annual variations of temperature, salinity, chlorophyll *a* and ammonium in a new supplement to our manuscript.

*Lines 221-222: The expression caused misunderstanding.*

We would like to change it to “The seasonal variations of NO<sub>2</sub><sup>-</sup> and NO<sub>3</sub><sup>-</sup> were significantly correlated with each other ([NO<sub>3</sub><sup>-</sup>]=11.59[NO<sub>2</sub><sup>-</sup>]-0.51, R<sup>2</sup>=0.80, n=72, p<0.0001) and high concentrations were observed for both in winter.”

*Lines 239-243: Does N<sub>2</sub>O correlate with NH<sub>4</sub><sup>+</sup>?*

There is no straightforward relationship between N<sub>2</sub>O and NH<sub>4</sub><sup>+</sup>. By the way, we realized that there were some problems with the calibration of pH data and, therefore, this part will be deleted.

*Line 462: The year of 2015 should be 2005?*

Thank you for pointing out the mistake. It should be 2005, which is the beginning of the N<sub>2</sub>O measurement.

*Line 476: There is no temperature data provided at all in this study but with a conclusion 'Temperature plays a modulating role for the N<sub>2</sub>O emission at the BE Time-Series Station'. I suggest to provide t data in Figure 2 and more t data provided in related discussion in the text*

We would like to show temperature data in the supplement. Also will rewrite the relevant text in section 3.5 to explain how temperature modulates N<sub>2</sub>O emissions in. See our reply to reviewer #1 as well.

*Figure 2: It would be better for the authors to provide the vertical profiles of t, s, density, NH<sub>4</sub><sup>+</sup> and Chl a.*

The data is now shown in the supplement.

*Figure 6: The vertical profiles of hydrological parameters, such as t, s and density are needed to help understand the possible influence of physical processes on N<sub>2</sub>O distribution as discussed between lines 301 and 309.*

Vertical profiles of temperature, salinity, NH<sub>4</sub><sup>+</sup> and Chl a are shown in supplement. Mixed layer variations can be seen in Fig. 4.

*Figure 8: title is needed for x- and y-axis at figure b and d*

*Figure 9: title is needed for x- and y-axis*

We have added new titles in the figures.

According to the comments from the reviewers, the following changes are made in the manuscript:

1. Show seasonal and annual variations of temperature, salinity, chlorophyll *a* and ammonium in a new supplement to our manuscript.
2. Rewrite the lines 86-88.
3. Add more details about standard gases in line 109.
4. Rewrite the lines 129-131.
5. Rewrite the lines 156-158.
6. Add more details in *Sc* computation in Section 2.4 (lines 165-171).
7. Rewrite the lines 221-223.
8. Rewrite the lines 238-239.
9. Delete the discussion about the relationship between  $\text{N}_2\text{O}$  and pH in section 3.2, and the corresponding figure was removed as well.
10. Modify the sentence in line 309.
11. Rewrite the lines 316-320.
12. Rewrite the lines 414-422.
13. Add the uncertainty of flux density computation in lines 428-429.
14. Add new titles in Fig. 7 and 8.

Besides the changes mentioned above, we also revised:

1. The computation of  $\Delta\text{N}_2\text{O}$  and AOU were removed from section 2.4 because the corresponding discussion was no longer in Results and Discussion.
2. Few typos in the manuscript were corrected.

1 **A multi-year observation of nitrous oxide at the Boknis Eck**  
2 **Time-Series Station in the Eckernförde Bay (southwestern**  
3 **Baltic Sea)**

4 Xiao Ma<sup>1</sup>, Sinikka T. Lennartz<sup>1,2</sup>, and Hermann W. Bange<sup>1</sup>

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9 **Abstract.** Nitrous oxide ( $N_2O$ ) is a potent greenhouse gas and it is involved in stratospheric ozone  
10 depletion. Its oceanic production is mainly influenced by dissolved nutrient and oxygen ( $O_2$ )  
11 concentrations in the water column. Here we examined the seasonal and annual variations of  
12 dissolved  $N_2O$  at the Boknis Eck (BE) Time-Series Station located in Eckernförde Bay  
13 (southwestern Baltic Sea). Monthly measurements of  $N_2O$  started in July 2005. We found a  
14 pronounced seasonal pattern for  $N_2O$  with high concentrations (supersaturations) in winter/early  
15 spring and low concentrations (undersaturations) in autumn when hypoxic/anoxic conditions  
16 prevail. Unusually low  $N_2O$  concentrations were observed during October 2016–April 2017,  
17 which was presumably a result of prolonged anoxia and the subsequent nutrient deficiency.  
18 Unusually high  $N_2O$  concentrations were found in November 2017 and this event was linked to  
19 the occurrence of upwelling which interrupted  $N_2O$  consumption via denitrification and  
20 potentially promoted ammonium oxidation (nitrification) at the oxic/anoxic interface. Nutrient  
21 concentrations (such as nitrate, nitrite and phosphate) at BE are decreasing since 1980s, but  
22 oxygen concentrations in the water column are still decreasing. Our results indicate a close  
23 coupling of  $N_2O$  anomalies to  $O_2$  concentration, nutrients and stratification. Given the long-term  
24 trends of declining nutrient and oxygen concentrations at BE, a decrease in  $N_2O$  concentration,  
25 and thus emissions, seems likely due to an increasing number of events with low  $N_2O$   
26 concentrations.

27 **1. Introduction**

28 Long-term observation with regular measurement intervals can be an effective way to monitor  
29 seasonal and interannual variabilities as well as to decipher short- and long-term trends of an  
30 ecosystem, which are required to make projections of the future ecosystem development (see e.g.  
31 Ducklow et al., 2009). Recently, multi-year time-series measurements of nitrous oxide ( $N_2O$ ), a  
32 potent greenhouse gas and a major threat to ozone depletion (IPCC, 2013; Ravishankara et al.,  
33 2009), have been reported from the coastal upwelling areas off central Chile (Farías et al., 2015)  
34 and off Goa (Naqvi et al., 2010), in the North Pacific Subtropical Gyre (Wilson et al., 2017), and  
35 in Saanich Inlet (Capelle et al., 2018).

36 N<sub>2</sub>O production in the ocean is generally dominated by microbial nitrification ( $\text{NH}_4^+ \rightarrow \text{NO}_2^- \rightarrow \text{NO}_3^-$ ) and denitrification ( $\text{NO}_3^- \rightarrow \text{NO}_2^- \rightarrow \text{N}_2\text{O} \rightarrow \text{N}_2$ ). During bacterial/archaeal nitrification, 37 N<sub>2</sub>O is produced as a by-product with enhanced N<sub>2</sub>O production under low oxygen (O<sub>2</sub>) 38 conditions (e.g. Goreau et al., 1980; Löscher et al., 2012). N<sub>2</sub>O is produced as an intermediate 39 during bacterial denitrification (Codispoti et al., 2005). N<sub>2</sub>O could be further consumed via 40 denitrification to dinitrogen, however, this process is inhibited with the presence of O<sub>2</sub> because 41 of the low O<sub>2</sub> tolerance of the enzyme involved (Bonin et al. 1989). This incomplete pathway is 42 called partial denitrification and can lead to N<sub>2</sub>O accumulation (e.g. Naqvi et al., 2000; Farías et 43 al., 2009).

45 The oceans including coastal areas contribute ~25% of the natural and anthropogenic N<sub>2</sub>O 46 emissions (IPCC, 2013), with disproportionately high emissions from coastal and estuarine areas 47 (Bange, 2006). N<sub>2</sub>O emissions from coastal areas strongly depend on nitrogen inputs (Seitzinger 48 and Kroeze, 1998; Zhang et al., 2010). The increasing input of nitrogen (i.e. eutrophication) has 49 become a worldwide problem in coastal waters leading to enhanced productivity and severe O<sub>2</sub> 50 depletion caused by enhanced degradation of organic matter (Breitburg et al., 2018; Rabalais et 51 al., 2014). The decline in O<sub>2</sub> concentration (i.e. deoxygenation), either in coastal waters or the 52 open ocean, might result in favorable conditions for N<sub>2</sub>O production (Codispoti et al., 2001; 53 Nevison et al., 2003). The results of a model study by Kroeze and Seitzinger (1998) indicated a 54 significant increase of N<sub>2</sub>O in European coastal waters for 2050. Moreover, it has been suggested 55 that N<sub>2</sub>O production and emissions are very likely to increase in the near future, especially in the 56 shallow suboxic/anoxic coastal systems (Naqvi et al., 2000; Bange, 2006). However, model 57 projections show a net decrease in future global oceanic N<sub>2</sub>O emission during the 21<sup>st</sup> century 58 (Martinez-Rey et al., 2015; Landolfi et al., 2017; Battaglia and Joos, 2018).

59 The Baltic Sea is a nearly enclosed, marginal sea with a very limited access to the open ocean via 60 the North Sea. The restricted water exchange with the North Sea and extensive human activities, 61 such agriculture, industrial production and sewage discharge in the catchment area led to high 62 inputs of nutrients to the Baltic Sea. As a result, the areas affected by anoxia have been 63 expanding in the deep basins of the central Baltic Sea (Carstensen et al., 2014). In order to 64 control this situation, the Helsinki Commission (HELCOM) was established in 1974 and a series 65 of measures have been taken to prevent anthropogenic nutrient input into the Baltic Sea. 66 Consequently, the nutrient inputs (by riverine loads, direct point-sources and, for nitrogen, 67 atmospheric deposition) to the Baltic Sea are declining (HELCOM, 2018a). However, the 68 number of low O<sub>2</sub> (i.e. hypoxic/anoxic) events in coastal waters of the Baltic Sea is increasing 69 and deoxygenation is still going on (Conley et al., 2011; Lennartz et al., 2014). The 70 deoxygenation in the Baltic Sea can affect the production/consumption of N<sub>2</sub>O. Our group has 71 been monitoring dissolved N<sub>2</sub>O concentrations at the Boknis Eck Time-Series Station, located in 72 Eckernförde Bay (southwestern Baltic Sea), for more than a decade. In this study, we present 73 monthly measurements of N<sub>2</sub>O and biogeochemical parameters such as nutrients and O<sub>2</sub> from 74 July 2005 to December 2017. The major objectives of our study were: 1) to decipher the seasonal

75 pattern of N<sub>2</sub>O distribution in the water column, 2) to identify short-term and long-term trends of  
76 the N<sub>2</sub>O concentrations, 3) to explore the potential role of nutrients and O<sub>2</sub> for N<sub>2</sub>O  
77 production/consumption, and 4) to quantify the sea-to-air N<sub>2</sub>O flux density at the time-series  
78 station.

## 79 **2. Material and methods**

### 80 **2.1 Study site**

81 Sampling at the Boknis Eck (BE) Time-Series Station ([www.bokniseck.de](http://www.bokniseck.de)) started on 30 April  
82 1957 and, therefore, it is one of the oldest continuously operated time-series stations in the world.  
83 The BE station is located at the entrance of the Eckernförde Bay (54°31' N, 10°02' E, Fig. 1) in  
84 the southwestern Baltic Sea. The water depth of the sampling site is 28 m. Various physical,  
85 chemical and biological parameters are measured on a monthly basis (Lennartz et al., 2014).  
86 **There is no significant river runoff to Eckernförde Bay. Hence, the hydrographical conditions are**  
87 **mainly dominated by saline water input from the North Sea and less saline water from the Baltic**  
88 **Proper, which is typical for that region.** Seasonal stratification usually starts to develop in April  
89 and lasts until October, during which hypoxia or even anoxia (characterized by the presence of  
90 hydrogen sulphide, H<sub>2</sub>S) sporadically occurs, as a result of restricted vertical water exchange and  
91 bacterial decomposition of organic matter in the bottom water (Hansen et al., 1999; Lennartz et  
92 al., 2014). Thus, BE is a natural laboratory to study the influence of O<sub>2</sub> variations and  
93 anthropogenic nutrient loads on N<sub>2</sub>O production/consumption.

### 94 **2.2 Sample collection and measurement**

95 Monthly sampling of N<sub>2</sub>O at the BE Time-Series Station started in July 2005. Triplicate samples  
96 were collected from six depths (1, 5, 10, 15, 20 and 25 m). Seawater was drawn from 5 L Niskin  
97 bottles into 20 mL brown glass vials after overflow. The vials were sealed with rubber stoppers  
98 and aluminum caps. The bubble-free samples were poisoned with 50 µL of a saturated mercury  
99 chloride (HgCl<sub>2</sub>) solution and then stored in a cool, dark place until measurement. The general  
100 storage time before measurements of the N<sub>2</sub>O concentrations was less than three months.

101 The static headspace-equilibrium method was adopted to measure the dissolved N<sub>2</sub>O  
102 concentrations in the vials. 10 mL helium (99.9999 %, AirLiquide, Düsseldorf, Germany)  
103 headspace was created in each vial with a gas-tight glass syringe (VICI Precision Sampling,  
104 Baton Rouge, LA). Samples were vibrated with Vortex (G-560E, Scientific Industries Inc., New  
105 York, USA) for 20 seconds and then left for at least two hours until equilibrium. 9.5 mL  
106 subsample of the headspace was subsequently injected into a GC-ECD (gas chromatograph  
107 equipped with the electron capture detector) system (Hewlett-Packard 5890 Series II, Agilent  
108 Technologies, Santa Clara, CA, USA), which was calibrated with two standard gas mixtures  
109 (N<sub>2</sub>O in synthetic air, 320 ppb and 1000 ppb, Deuste-Steininger GmbH, Mühlhausen, Germany  
110 and Westfalen AG, Münster, Germany) prior to the measurement. The average precision of the

111 measurements, calculated as the median standard deviation from triplicate measurements, was  
112 0.4 nM. Triplicates with a standard deviation of >10% were omitted. More details about the N<sub>2</sub>O  
113 measurement can be found in Kock et al. (2016). Dissolved oxygen (O<sub>2</sub>) concentrations were  
114 measured by Winkler titrations (Grasshoff et al., 1999). Nutrient concentrations were measured  
115 by the Segmented Continuous Flow Analysis (SCFA, Grasshoff et al., 1999). A more detailed  
116 summary of the parameters measured and methods applied can be found in Lennartz et al. (2014).

## 117 **2.3 Times series analysis**

118 A time-series can be decomposed into three main components, i.e. trend, cycle and residual  
119 component (Schlittgen and Streitberg, 2001). We used the Mann–Kendall test and wavelet  
120 analysis to detect the trend and periodical cycles in the time-series data, respectively. As for the  
121 residual component, we highlight unusual high/low N<sub>2</sub>O concentrations during 2005-2017 and  
122 discuss the potential causes for these events.

### 123 **2.3.1 Wavelet analysis**

124 In order to decipher periodical cycles of the parameters collected at the BE Time-Series Station,  
125 a wavelet analysis method was adopted. Wavelet analysis enables the detection of the period and  
126 the temporal occurrence of repeated cycles in time-series data. One of the requirements for  
127 wavelet analysis is a regular, continuous time-series. Since there is data missing (maximum 2  
128 months in a row) in the BE time-series, due to terrible weather or the ship's unavailability,  
129 missing data was interpolated from the previous and following months. **Sampling time varied for**  
130 **every month (usually 20-40 day interval), but for the statistical analysis, data was assumed to be**  
131 **regularly spaced as differences on weekly scales were minor.** Considering the band width in both  
132 frequency and time domain, a Morlet mother wavelet with a wave number of 6 was chosen  
133 (Torrence and Compo, 1998). The mother wavelet was then scaled between the frequency of a  
134 half-year cycle and the length of the time-series with a stepsize of 0.25. The wavelet analysis was  
135 conducted with the MatLab code by Torrence and Compo [2004]. More information about the  
136 method can be found on the website <http://paos.colorado.edu/research/wavelets/>.

### 137 **2.3.2 Mann–Kendall test**

138 Mann–Kendall test (MKT) is a non-parametric statistical test to assess the significance of  
139 monotonic trends for time-series measurements. It tests the null hypothesis that all variables are  
140 randomly distributed against the alternative hypothesis that a monotonic trend, either increase or  
141 decrease, exists in the time-series on a given significance level  $\alpha$  (here  $\alpha=0.05$ ). MKT is flexible  
142 for data with missing values and the results are not impacted by the magnitude of extreme values,  
143 which makes it a widely used test in hydrology and climatology (e.g. Xu et al., 2003; Yang et al.,  
144 2004). However, MKT is sensitive to serial correlation in the time-series. The presence of  
145 positive serial correlation would increase the probability of trend detection even though no such  
146 trend exists (Kulkarni and von Storch, 1995). In order to avoid this situation, data from 12

147 months were tested individually. It is assumed that there is no residual effect left from the same  
148 month last year, considering that the nitrogen species are rapidly biologically cycled. The Matlab  
149 function from Simone (2009) was used for the MKT.

## 150 **2.4 Calculation of saturation and sea-to-air flux density**

151  $N_2O$  saturations ( $S_{N2O}$ , %) were calculated as:

$$152 \quad S_{N2O} = 100 \times N_2O_{obs}/N_2O_{eq} \quad (1)$$

153 where  $N_2O_{obs}$  and  $N_2O_{eq}$  (in nM) are the observed and equilibrated  $N_2O$  concentrations in  
154 seawater, respectively.  $N_2O_{eq}$  was computed as a function of surface seawater temperature, in  
155 situ salinity (Weiss and Price, 1980) and the dry mole fractions of atmospheric  $N_2O$  at the time  
156 of the sampling. Since the atmospheric  $N_2O$  mole fractions were not measured at the BE Time-  
157 Series Station, atmospheric dry mole fractions of  $N_2O$  were derived from the monthly average of  
158  $N_2O$  data at Mace Head, Ireland (AGAGE, <http://agage.mit.edu/>), instead.

159  $N_2O$  flux density ( $F_{N2O}$ , in  $\mu\text{mol m}^{-2} \text{ d}^{-1}$ ) was calculated as:

$$160 \quad F_{N2O} = k_{N2O} \times (N_2O_{obs} - N_2O_{eq}) \quad (2)$$

161 where  $k_{N2O}$  (in  $\text{cm h}^{-1}$ ) is the gas transfer velocity calculated with the method given by  
162 Nightingale et al. (2000), as a function of the wind speed and the Schmidt number ( $Sc$ ). The wind  
163 speed data were obtained from the Kiel Lighthouse (see: [www.geomar.de/service/wetter/](http://www.geomar.de/service/wetter/)), which  
164 is approximately 20 km away from the BE Time-Series Station. The wind speed was normalized  
165 to 10 m ( $u_{10}$ ) to calculate  $k_{N2O}$  (Hsu et al., 1994).  $k_{N2O}$  was adjusted by multiplying with  $(Sc/600)^{0.5}$ ,  
166 and  $Sc$  was computed as:

$$167 \quad Sc = v/D_{N2O} \quad (3)$$

$$168 \quad D_{N2O} = 3.16 \times 10^{-6} e^{-18370/RT} \quad (4)$$

169 where  $v$  is the kinematic viscosity of seawater, which is calculated from the empirical equations  
170 given in Siedler and Peters (1986), and  $D_{N2O}$  is the diffusion coefficient of  $N_2O$  in seawater.  $R$  is  
171 the universal gas constant and  $T$  is the water temperature in K.

## 172 **3. Result and discussion**

### 173 **3.1 Overview**

174  $N_2O$  concentrations at the BE Time-Series Station showed significant temporal and depth-  
175 dependent variations from 2005 to 2017 (Fig. 2).  $N_2O$  concentrations fluctuated between 1.2 and  
176 37.8 nM, with an overall average of  $13.9 \pm 4.2$  nM. This value was higher than the results from  
177 the surface water of Station ALOHA (5.9–7.4 nmol  $\text{kg}^{-1}$ , average  $6.5 \pm 0.3$  nmol  $\text{kg}^{-1}$ , Wilson et

178 al., 2017), which is reasonable considering the weak anthropogenic impact in the North Pacific  
179 Subtropical Gyre. The  $\text{N}_2\text{O}$  concentrations at BE were much lower than those measured at the  
180 time-series station in the coastal upwelling area off Chile (2.9–492 nM, average  $39.4\pm29.2$  nM in  
181 the oxyclines and  $37.6\pm23.3$  nM in the bottom waters, Farías et al., 2015) and a quasi-time series  
182 station off Goa (Naqvi et al., 2010), where significant  $\text{N}_2\text{O}$  accumulations were observed in  
183 subsurface waters at both locations. Our measurements were comparable to the time-series  
184 station from Saanich Inlet (~0.5–37.4 nM, average 14.7 nM, Capelle et al., 2018), a seasonally  
185 anoxic fjord which has similar hydrographic conditions as BE.

186  $\text{NO}_2^-$  concentrations fluctuated between below detection limit of 0.1  $\mu\text{M}$  and 1.6  $\mu\text{M}$ , with an  
187 average of  $0.2\pm0.3$   $\mu\text{M}$ .  $\text{NO}_3^-$  concentrations varied from below detection limit of 0.3  $\mu\text{M}$  to 17.9  
188  $\mu\text{M}$ , with an average of  $2.0\pm2.8$   $\mu\text{M}$ . The temporal and spatial distributions of nitrite ( $\text{NO}_2^-$ ) and  
189 nitrate ( $\text{NO}_3^-$ ) were similar during 2005–2017. A clear  $\text{O}_2$  seasonality can be seen with severe  $\text{O}_2$   
190 depletion in the bottom waters during summer and autumn. Anoxia with the presence of  $\text{H}_2\text{S}$   
191 were detected in September/October 2005, September 2007, September/October 2014, and  
192 September–November 2016. All of the extremely low  $\text{N}_2\text{O}$  concentrations (<5 nM) were  
193 observed in the bottom waters in autumn, coinciding with hypoxia/anoxia, while the high  $\text{N}_2\text{O}$   
194 concentrations (>20 nM) sporadically occurred at different depths either in spring or autumn.

### 195 3.2 Seasonal cycle

196 Significant cycles at different frequencies were detected via wavelet analysis at the BE Time-  
197 Series Station during 2005–2017 (Fig. 3). A half-year  $\text{NO}_2^-$  cycle sporadically occurred in 2007–  
198 2009, 2013 and 2015. There is a seasonal  $\text{NO}_2^-$  variability (at the frequency of 1 year) between  
199 2007 and 2016 (times before 2007 and after 2016 were outside the conic line), except during  
200 2010–2012, when high  $\text{NO}_2^-$  concentrations were not observed in winter (Fig. 2). A biennial  
201 cycle of  $\text{NO}_2^-$  could be observed as well during 2008–2015. The  $\text{NO}_3^-$  concentrations were  
202 dominated by an annual cycle and a minor half-year cycle. The biennial cycle only occurred in  
203 2008 and 2009. A remarkable seasonal variability of dissolved  $\text{O}_2$  prevailed all the time, which is  
204 also obvious from the times series data shown in Fig. 2. The annual  $\text{N}_2\text{O}$  cycle became gradually  
205 more and more evident until 2014, then declined and reoccurred less intensely in 2016. The  
206 periodical cycle was also present at other frequencies, indicated by the broadening of the red area  
207 before 2015 in Fig. 2d. For example, a biennial  $\text{N}_2\text{O}$  cycle occurred during 2013–2015.

208 The half-year cycles of  $\text{NO}_2^-$  and  $\text{NO}_3^-$  were probably associated with algae blooms which  
209 usually occur in each spring and autumn (Fig. S1 and S2). Since the time between the two  
210 blooms differed between years, the cycles were weak and thus not present in every year. Due to  
211 the fact that there was no half-year  $\text{O}_2$  cycle at all, nutrients apart from  $\text{O}_2$  might be the “drivers”  
212 of the sporadic half-year  $\text{N}_2\text{O}$  cycle in 2008 and 2015, because  $\text{N}_2\text{O}$  production depends on the  
213 concentration of the bioavailable nitrogen compounds (Codispoti et al., 2001).

214 Generally the wavelet analysis indicated a strong annual cycle for  $\text{NO}_2^-$ ,  $\text{NO}_3^-$ , dissolved  $\text{O}_2$  and  
215  $\text{N}_2\text{O}$  at the BE Time-Series Station, which enabled us to explore the seasonal pattern with annual  
216 mean data. Although extreme values were excluded as a result of averaging, the smoothed results  
217 generally reflect the seasonality of these parameters. Here, we focus on the annual cycle.

218 The annual mean vertical distribution of dissolved  $\text{O}_2$ ,  $\text{NO}_2^-$ ,  $\text{NO}_3^-$  and  $\text{N}_2\text{O}$  are shown in Fig. 4.  
219 Due to the development of stratification, the mixed layer was shallow in summer and deep in late  
220 autumn/winter.  $\text{O}_2$  depletion was observed in bottom waters from late spring until late autumn.  
221 The seasonal variations of  $\text{NO}_2^-$  and  $\text{NO}_3^-$  were significantly correlated with each other  
222 ( $[\text{NO}_3^-] = 11.59[\text{NO}_2^-] - 0.51$ ,  $R^2 = 0.80$ ,  $n = 72$ ,  $p < 0.0001$ ) and high concentrations were observed  
223 for both in winter. Minimum  $\text{N}_2\text{O}$  concentrations were found in the bottom waters during  
224 September and October, presumably as a result of consumption during denitrification under  
225 anoxic condition (Codispoti et al., 2005). High  $\text{N}_2\text{O}$  concentrations were observed in late spring  
226 and late autumn, respectively. In late spring  $\text{N}_2\text{O}$  accumulated in the bottom waters because the  
227 stratification prevented mixing of the water column. In late autumn, however,  $\text{N}_2\text{O}$  could be  
228 ventilated to the surface and thus emitted to the atmosphere due to the breakdown of the  
229 stratification. The high  $\text{N}_2\text{O}$  concentrations could be attributed to enhanced  $\text{N}_2\text{O}$  production via  
230 nitrification and/or denitrification within the oxic/anoxic interface (Goreau et al., 1980;  
231 Codispoti et al., 1992). Since there is no clear  $\text{O}_2$  concentration threshold,  $\text{N}_2\text{O}$  production from  
232 both nitrification and the onset of denitrification overlap at oxic/anoxic interface. To this end,  
233 direct  $\text{N}_2\text{O}$  production measurements (i.e. nitrification/denitrification rates) are required to  
234 decipher which process dominates the formation of the different  $\text{N}_2\text{O}$  maxima.

235 High  $\text{N}_2\text{O}$  concentrations prevailed all over the water column in winter/early spring.  $\text{NH}_4^+$  is  
236 released from the sediment into bottom waters due to the degradation of organic matter,  
237 especially after the autumn algae bloom (Fig. S1 and S2). The stratification usually completely  
238 breaks down at this time of the year and the water column becomes oxygenated. Denitrification  
239 is inhibited by the presence of high concentrations of dissolved  $\text{O}_2$  ( $> 20 \mu\text{mol L}^{-1}$ ) and thus  
240 nitrification is presumably responsible for the high  $\text{N}_2\text{O}$  concentrations in winter/early spring.

### 241 3.3 Trend analysis

242 The MKTs were conducted for the surface (1m) and bottom (25m)  $\text{N}_2\text{O}$  concentrations and  
243 saturations of the individual 12 months, respectively. Significant decreasing trends were detected  
244 for the concentrations in the bottom waters for February and August (Table 1a), and for the  
245 saturations in the surface for September and in the bottom for August and November (Table 1b).  
246 These results indicated that some systematical changes in  $\text{N}_2\text{O}$  took place at BE. For example,  
247 the significant decrease in  $\text{N}_2\text{O}$  concentration/saturation in August might be associated with the  
248 increasing temperature, which reinforces the stratification and accelerates  $\text{O}_2$  consumption in the  
249 bottom waters (Lennartz et al., 2014). As a result, hypoxia/anoxia starts earlier and thus enables  
250 the onset of denitrification to consume  $\text{N}_2\text{O}$ . During most of the months, trends in  $\text{N}_2\text{O}$   
251 concentration and saturation were not significant during 2005–2017.

252 A significant nutrient decline has been observed at the BE Time-Series Station since the mid-  
253 1980s, however, Lennartz et al. (2014) found that bottom O<sub>2</sub> concentrations were still decreasing  
254 over the past 60 years. The ongoing oxygen decline was attributed to the temperature-enhanced  
255 O<sub>2</sub> consumption in the bottom water (Meier et al., 2018) and a prolongation of the stratification  
256 period at the BE Time-Series Station (Lennartz et al., 2014). Please note that the trends in  
257 nutrients and O<sub>2</sub> concentrations were detected based on the data collection which lasted for  
258 approximately 30 and 60 years, respectively, while the N<sub>2</sub>O observations at BE Time-Series  
259 Station has lasted for only 12.5 years. Further MKT analysis for nutrients, temperature and  
260 oxygen for months with significant trends in N<sub>2</sub>O concentrations did not show any significant  
261 results ( $p>0.05$ ). The significant trends in N<sub>2</sub>O concentrations thus do not seem to be directly  
262 related to one of these parameters, and we cannot state a reason for the significant trends of N<sub>2</sub>O  
263 concentration in February and the N<sub>2</sub>O saturation in September and November at this point.  
264 Presumably, a longer monitoring period for N<sub>2</sub>O is required to detect corresponding trends in  
265 N<sub>2</sub>O and oxygen or nutrients.

## 266 **3.4 Extreme events**

### 267 **3.4.1 Low N<sub>2</sub>O concentrations during October 2016-April 2017**

268 Besides the low N<sub>2</sub>O concentrations occurring in autumn, we observed a band of pronounced  
269 low N<sub>2</sub>O concentrations which started in October 2016 and lasted until April 2017 (Fig. 5). In  
270 this period N<sub>2</sub>O concentrations varied between 5.5–13.9 nM, with an average of  $8.4\pm2.0$  nM.  
271 This is approximately 40% lower than the average N<sub>2</sub>O concentration during the entire  
272 measurement period 2005–2017. The average N<sub>2</sub>O saturation during 2005–2017 was  $111\pm30\%$ ,  
273 while from October 2016 to April 2017, the N<sub>2</sub>O saturations were as low as 43–93% (average  
274  $62\pm10\%$ ).

275 Undersaturated N<sub>2</sub>O waters have been previously reported from the Baltic Sea: Rönner (1983)  
276 observed a N<sub>2</sub>O surface saturation of 79% in the central Baltic Sea and attributed the  
277 undersaturation to upwelling of N<sub>2</sub>O-depleted waters. Bange et al. (1998) found a minimum N<sub>2</sub>O  
278 saturation of 91% in the southern Baltic Sea where the hydrographic conditions were  
279 significantly influenced by riverine runoff. Walter et al. (2006) reported a mean N<sub>2</sub>O saturation  
280 of  $79\pm11\%$  for shallow stations (<30 m) in the southwestern Baltic Sea in October 2003. The  
281 low-N<sub>2</sub>O event at BE was unusual because the concentrations were much lower than those  
282 reported values and it lasted for more than half a year.

283 Although the observed temperatures and salinities during October 2016–April 2017 were  
284 comparable to other years (Fig. S1), it is difficult to evaluate the role of physical mechanism in  
285 the low-N<sub>2</sub>O event because of insufficient data for water mass exchange at the BE Time-Series  
286 Station. Here we mainly focused on the chemical or biological processes. Anoxia events with the  
287 presence of H<sub>2</sub>S were observed in the bottom waters for three months in a row during  
288 September–November 2016. This is an unusual long period and is unprecedented at the BE

289 Time-Series Station. In December 2016 the stratification did not completely break down.  
290 Although the water column was generally oxygenated, bottom O<sub>2</sub> concentrations were the lowest  
291 observed during the past ten years. Considering the classical view of N<sub>2</sub>O consumption via  
292 denitrification under hypoxic and anoxic conditions, we inferred that denitrification accounted  
293 for low N<sub>2</sub>O concentrations in the bottom layer. However, the question still remains where the  
294 low-N<sub>2</sub>O-concentration water in the upper layers came from.

295 In September 2016, low N<sub>2</sub>O concentrations were only observed in the bottom waters where the  
296 anoxia occurred. However, the situation was different in the following months. During  
297 October/November 2016, N<sub>2</sub>O concentrations were homogeneously distributed in the water  
298 column. Although the stratification gradually started to break down in late autumn, the density  
299 gradient was still strong enough to keep the bottom waters at anoxic conditions and prevented  
300 the low-N<sub>2</sub>O-concentration to reach the surface. Thus we inferred that the unusual low N<sub>2</sub>O  
301 concentrations in the upper layers (above 20 m) were probably resulting from advection of  
302 adjacent waters. Due to the fact that the upper layers were well-mixed and oxygenated, in situ  
303 N<sub>2</sub>O consumption in the water column could be neglected. We suggest therefore, that the N<sub>2</sub>O  
304 depleted waters were resulting from consumption of N<sub>2</sub>O in bottom waters elsewhere and then  
305 they were upwelled and transported to BE. Hence, N<sub>2</sub>O consumption via denitrification might  
306 have been, directly or indirectly, responsible for the low N<sub>2</sub>O concentrations during October–  
307 November 2016.

308 In December 2016, the bottom waters were ventilated with O<sub>2</sub>. Although N<sub>2</sub>O consumption by  
309 denitrification should have been inhibited by the high concentrations of O<sub>2</sub> (Codispoti et al.,  
310 2001), the N<sub>2</sub>O concentrations did not restore to their normal level under suboxic conditions.  
311 Since January 2017, the whole water column was well mixed and oxygenated. Usually a  
312 significant nutrient supply could be observed starting in November (Fig. 4) as a result of  
313 remineralization and vertical mixing, but the average NO<sub>2</sub><sup>−</sup> and NO<sub>3</sub><sup>−</sup> concentrations during  
314 November 2016–April 2017 were 0.2 and 1.4 μM, respectively, which was about 50% and 60%  
315 lower than in other years. Ammonium (NH<sub>4</sub><sup>+</sup>) and chlorophyll *a* concentrations during this  
316 period were comparable to that of other years (Fig. S1). Secchi depth, a proxy of water  
317 transparency, was 3.8 m in March 2017, which is only slightly lower compared to the monthly  
318 average value for March (4.5±1.8 m). There is no exceptional spring algae bloom and thus we  
319 infer that assimilative uptake of nutrients by phytoplankton was not responsible for the low  
320 nutrients concentrations. The nutrient deficiency might be attributed to enhanced nitrogen  
321 removal processes like denitrification or anammox (Voss et al., 2005; Hietanen et al., 2007;  
322 Hannig et al., 2007) during the prolonged period of anoxia in autumn 2016. During the low N<sub>2</sub>O  
323 event, we found that N<sub>2</sub>O concentrations were positively correlated with both NO<sub>2</sub><sup>−</sup>  
324 ( $[N_2O]=7.02[NO_2^-]+7.36$ ,  $R^2=0.29$ ,  $n=24$ ,  $p<0.01$ ) and NO<sub>3</sub><sup>−</sup> ( $[N_2O]=0.80[NO_3^-]+7.36$ ,  $R^2=0.51$ ,  
325  $n=24$ ,  $p<0.0001$ ). These results indicate that the development and maintenance of the low-N<sub>2</sub>O-  
326 concentration was closely associated with nutrient deficiency. Especially after the breakdown of

327 the stratification, when denitrification was no longer a significant  $\text{N}_2\text{O}$  sink, nutrients might have  
328 become a limiting factor for  $\text{N}_2\text{O}$  production.

329 In general, the low- $\text{N}_2\text{O}$ -concentration event during October 2016–April 2017 can be divided  
330 into two parts: in the stratified waters during October–November 2016,  $\text{O}_2$  played a dominant  
331 role and  $\text{N}_2\text{O}$  was consumed via denitrification under anoxic conditions. In the well-mixed water  
332 column during December 2016–April 2017, nutrient deficiency seemed to have constrained  $\text{N}_2\text{O}$   
333 production via nitrification under suboxic/oxic conditions.

334 In recent years a novel biological  $\text{N}_2\text{O}$  consumption pathway, called  $\text{N}_2\text{O}$  fixation, which  
335 transforms  $\text{N}_2\text{O}$  into particulate organic nitrogen via its assimilation, has been reported (Farías et  
336 al., 2013). This process can take place under extreme environmental conditions even at very low  
337  $\text{N}_2\text{O}$  concentrations. Cornejo et al. (2015) reported that  $\text{N}_2\text{O}$  fixation might play a major role in  
338 the coastal zone off central Chile where seasonally occurring surface  $\text{N}_2\text{O}$  undersaturation was  
339 observed. The relatively high  $\text{N}_2$  fixation rates in the Baltic Sea (Sohm et al., 2011) highlight the  
340 potential role of  $\text{N}_2\text{O}$  fixation (Farías et al., 2013). However, we cannot quantify the role of  
341 biological  $\text{N}_2\text{O}$  fixation for the  $\text{N}_2\text{O}$  depletion in the Baltic Sea due to the absence of  $\text{N}_2\text{O}$   
342 assimilation measurements.

### 343 **3.4.2 High $\text{N}_2\text{O}$ concentrations in November 2017**

344 High  $\text{N}_2\text{O}$  concentrations were observed at the BE Time-Series Station in November 2017. The  
345 average value reached  $35.4 \pm 1.5 \text{ nM}$ , which was the highest concentration measured during the  
346 entire sampling period from 2005 to 2017. Dissolved  $\text{N}_2\text{O}$  was homogeneously distributed in the  
347 water column, but this event did not last long. In December, dissolved  $\text{N}_2\text{O}$  returned to normal  
348 levels and the average concentration in the water column was comparable to that of other years.  
349 Average  $\text{N}_2\text{O}$  saturation in November 2017 was  $322 \pm 10\%$ , which was also the highest for the  
350 past 12.5 years. This value was much higher than the maximum surface  $\text{N}_2\text{O}$  saturation reported  
351 by Rönner (1983) in the central Baltic Sea, but was comparable to the results observed in the  
352 southern Baltic Sea (312%, Bange et al., 1998). Bange et al. (1998) linked the enhanced  $\text{N}_2\text{O}$   
353 concentrations to riverine runoff because those samples were collected in an estuarine area,  
354 however, the riverine influence around the BE Time-Series Station is negligible. As a result, the  
355 impact of fresh water input can be excluded.

356 Dissolved  $\text{O}_2$  seemed to play a dominant role in the high  $\text{N}_2\text{O}$  concentrations. Enhanced  $\text{N}_2\text{O}$   
357 production usually occurred at the oxic/anoxic interface, which was closely linked to the  
358 development of water column stratification. In general the breakdown of the stratification is  
359 faster than its establishment at the BE Time-Series Station. As a result, it took about half a year  
360 for bottom  $\text{O}_2$  saturation to gradually decrease from ~80% to almost 0% (i.e. anoxia), but only  
361 two months to restore normal saturation level in 2010 (Fig. 6). In late autumn, surface water  
362 penetrated into the deep layers via vertical mixing and eroded the oxic/anoxic interface. The  
363 entire water column quickly became oxygenated and the enhanced  $\text{N}_2\text{O}$  production was stopped.

364 Hypoxia/anoxia at BE is usually observed in the bottom waters in autumn, but in September  
365 hypoxic water ( $O_2$  saturation < 20 %, which was close to the criterion for hypoxia, see  
366 Naqvi et al., 2010) was found in the subsurface layer (10 m) as well. Surface  $O_2$  saturation was  
367 only ~50%, which was the lowest during the sampling period 2005–2017. The density gradient  
368 of the water column in September 2017 was much lower than in other years. These results  
369 indicate the occurrence of an upwelling event at BE Time-Series Station in autumn 2017, which  
370 might be a result of the saline water inflow from the North Sea considering the change of salinity  
371 in the water column (Fig. S1). Strong vertical mixing has interrupted the hypoxia/anoxia and  
372 bottom  $O_2$  saturation reached ~60% in October 2017. The presence of  $O_2$  prevented  $N_2O$   
373 consumption via denitrification, as a result, we did not observe a significant  $N_2O$  decline during  
374 that period (Fig. 5).

375 Considering the fact that a significant autumn algae bloom was observed in autumn 2017 (as  
376 indicated by high chlorophyll a concentrations, see Fig. S1), severe  $O_2$  depletion in the bottom  
377 water could be expected. Although the bottom  $O_2$  saturation was only slightly lower in  
378 November than in October, we speculate that even lower  $O_2$  saturation (but not anoxia) might  
379 have occurred between October and November. The “W-shaped”  $O_2$  saturation curve (see Fig. 6)  
380 suggests that the stratification did not completely break down in October and that there might  
381 have been a reestablishment of the oxic/anoxic interface providing favorable conditions for  
382 enhanced  $N_2O$  production. Due to the degradation of organic nitrogen,  $NH_4^+$  is released from the  
383 sediment into bottom waters (Dale et al., 2011), especially in autumn when  $O_2$  is low (Fig. S2).  
384  $NH_4^+$  concentrations in November 2017 were lower than in other years (Fig. S1), and  $NO_2^-$   
385 concentrations were higher (Fig. 5), indicating that nitrification occurred in bottom waters. To  
386 this end, we suggest that the reestablishment of the oxic/anoxic interface promoted ammonium  
387 oxidation (the first step of nitrification). In this case,  $N_2O$  could have temporary accumulated  
388 because its consumption via denitrification was blocked. Meanwhile, the relatively low density  
389 gradient (i.e. low stratification) allowed upward mixing of the excess  $N_2O$  to the surface.  
390 However, we inferred that that this phenomenon would only last for a few days due to the rapid  
391 breakdown of stratification at the BE Time-Series Station.

392 Due to the development of the pronounced stratification, the oxic/anoxic interface prevailed in  
393 summer/early autumn as well, but we did not observe  $N_2O$  accumulation during these months.  
394 One of the potential explanations is that enhanced  $N_2O$  production only took place within  
395 particular depths where strong  $O_2$  gradient existed, but our vertical sampling resolution was too  
396 low to capture this event. Also enhanced  $N_2O$  production might be covered by the weak mixing  
397 which brought low- $N_2O$  water from the bottom to the surface.

398 The upwelling event played different roles in autumn 2016 and 2017. First, upwelling took place  
399 somewhere else but at BE because of the strong density and  $O_2$  gradient in the water column  
400 during autumn 2016. Second, bottom water remained anoxic in autumn 2016, while the  
401 compensated water for upwelling in 2017 penetrated through stratification and brought  $O_2$  into  
402 bottom water (Fig. 6), which caused enhanced  $N_2O$  production. Similarly, autumn upwelling was

403 detected in 2011 and 2012 when we found relatively low O<sub>2</sub> concentrations in subsurface layers  
404 (10 m) (Fig. 2), but we did not observe an increase in bottom O<sub>2</sub> concentrations and N<sub>2</sub>O  
405 concentrations remained low during that time. These upwelling events seem to be driven by  
406 saline water inflow considering the prominent increase in salinity, but the mechanism dominates  
407 O<sub>2</sub> input into bottom water before the stratification break down remains unclear.

### 408 3.5 Flux density

409 During 2005–2017, surface N<sub>2</sub>O saturations at the BE Time-Series Station varied from 56 % to  
410 314 % (69–194 % excluding the extreme values discussed in Sect. 3.4), with an average of  
411 111±30 % (111±20 % without the extreme values). Generally the water column at BE was  
412 slightly oversaturated with N<sub>2</sub>O. Our results are in good agreement with the estimated mean  
413 surface N<sub>2</sub>O saturation for the European shelf (113%, Bange, 2006).

414 We found a weak seasonal cycle for surface N<sub>2</sub>O concentrations, with high N<sub>2</sub>O concentrations  
415 occurring in winter/early spring and low concentrations occurring in summer/autumn, but no  
416 such cycle for N<sub>2</sub>O saturation (Fig. 4; Fig. 7). The seasonality in concentration but not in  
417 saturation could be largely attributed to the effect of temperature on N<sub>2</sub>O solubility: In summer  
418 when surface N<sub>2</sub>O concentrations are low, N<sub>2</sub>O saturations are increased by the relative high  
419 temperature; and vice versa in winter. Although salinity also affects N<sub>2</sub>O solubility, its  
420 contribution is negligible compared to temperature. Temperature alleviated the fluctuation of  
421 surface N<sub>2</sub>O saturation and thus affected the sea-to-air N<sub>2</sub>O fluxes. We conclude that temperature  
422 plays a modulating role for N<sub>2</sub>O emissions.

423 The wind speed (u<sub>10</sub>) at the BE Time-Series Station ranged from 1.1 to 14.0 m s<sup>-1</sup>, with an  
424 average of 7.0±2.7 m s<sup>-1</sup>. N<sub>2</sub>O flux densities varied from -19.0 to 105.7 µmol m<sup>-2</sup> d<sup>-1</sup> (-14.1–30.3  
425 µmol m<sup>-2</sup> d<sup>-1</sup> without the extreme values), with an average of 3.5±12.4 µmol m<sup>-2</sup> d<sup>-1</sup> (3.3±6.5  
426 µmol m<sup>-2</sup> d<sup>-1</sup> without the extreme values). However, the true emissions might have been  
427 underestimated because our monthly sampling resolution is insufficient to capture short-term  
428 N<sub>2</sub>O accumulation events due to the fast breakdown of stratification in autumn. The uncertainty  
429 introduced in the flux density computation was estimated to be 20% (Wanninkhof, 2014). The  
430 flux densities at the BE Time-Series Station are comparable to those reported by Bange et al.  
431 (1998, 0.4–7.1 µmol m<sup>-2</sup> d<sup>-1</sup>) from the coastal waters of the southern Baltic Sea, but are slightly  
432 lower than the average N<sub>2</sub>O flux density reported by Rönner (1983, 8.9 µmol m<sup>-2</sup> d<sup>-1</sup>) from the  
433 central Baltic Sea. Please note that the results of Rönner (1983) were obtained only from the  
434 summer season and therefore are probably biased because of missing seasonality.

435 In December 2014, a strong saline water inflow from the North Sea was observed, which was the  
436 third strongest ever recorded (Mohrholz et al., 2015). Although the salinity in December 2014  
437 was comparable to other years, a remarkable increase in salinity was observed in the following  
438 several months. However, we did not detect a significant N<sub>2</sub>O anomaly or enhanced emission  
439 during that time. Similarly, Walter et al. (2006) investigated the impact of the North Sea water

440 inflow on N<sub>2</sub>O production in the southern and central Baltic Sea in 2003. The oxygenated water  
441 ventilated the deep Baltic Sea and shifted anoxic to oxic condition which led to enhanced N<sub>2</sub>O  
442 production, but the accumulated N<sub>2</sub>O was unlikely to reach the surface due to the presence of a  
443 permanent halocline (Walter et al., 2006).

444 Although we observed extremely high N<sub>2</sub>O flux density in November 2017, the low-N<sub>2</sub>O-  
445 concentration (<10 nM) events have become more and more frequent during the past ten years  
446 (Fig. 2). This phenomenon seldom occurred before 2011, but remarkable low N<sub>2</sub>O  
447 concentrations can be seen in 2011 and 2013, and to a less extent in 2012 and 2014. Similar  
448 events lasted for several months in 2015 and for even more than half a year during 2016–2017.  
449 The most striking was that the low-N<sub>2</sub>O-concentration water was not only detected in bottom  
450 waters, but also at surface which would significantly impact the air-sea N<sub>2</sub>O flux densities.  
451 Although the MKT result did not give a significant trend for the N<sub>2</sub>O flux densities, the data  
452 presented in Fig. 8 suggest a potential decline of N<sub>2</sub>O flux densities from the coastal Baltic Sea,  
453 challenging the conventional view that N<sub>2</sub>O emissions from coastal waters would most probably  
454 increase in the future, which was based on the hypothesis of increasing nutrient loads into coastal  
455 waters. Due to an effective reduction of nutrient inputs, the severe eutrophication condition in the  
456 Baltic Sea has been alleviated (HELCOM, 2018b), but ongoing deoxygenation points to the fact  
457 that it will take a longer time for coastal ecosystems to feedback to reduced nutrient inputs  
458 because other environmental changes such as warming may override decreasing eutrophication  
459 (Lennartz et al., 2014).

## 460 **4. Conclusions**

461 The seasonal and inter-annual N<sub>2</sub>O variations at the BE Time-Series Station from July 2005 to  
462 December 2017 were driven by the prevailing O<sub>2</sub> regime and nutrients availability. We found a  
463 pronounced seasonal cycle with low N<sub>2</sub>O concentrations (undersaturations) occurring in  
464 hypoxic/anoxic bottom waters in autumn and enhanced concentrations (supersaturations) all over  
465 the water column in winter/early spring. Significant decreasing trends for N<sub>2</sub>O concentrations  
466 were found for few months, while most of the year, no significant trend was detectable in the  
467 period of 2005–2017. During 2005–2017, no significant trends were present for O<sub>2</sub> and nutrients  
468 either, but these parameters all show significant decreasing trends on longer time scales (~60  
469 years) at BE. Our results show the strong coupling of N<sub>2</sub>O with O<sub>2</sub> and nutrient concentrations,  
470 and suggest similar changes on comparable time scales. Further monitoring of N<sub>2</sub>O at BE time  
471 series station is thus important to detect changes. Further studies on N<sub>2</sub>O  
472 production/consumption by nitrification and denitrification and analysis of the characteristic N<sub>2</sub>O  
473 isotope signature might be very helpful to decipher the potential roles of O<sub>2</sub>/nutrients for N<sub>2</sub>O  
474 cycling.

475 Temperature plays a modulating role for the N<sub>2</sub>O emission at the BE Time-Series Station.  
476 Although the hydrographic condition at BE is generally dominated by the inflow of saline North  
477 Sea water, this did not affect N<sub>2</sub>O production and its emissions to the atmosphere. It seems that

478 events with extremely low N<sub>2</sub>O concentrations and thus reduced N<sub>2</sub>O emissions became more  
479 frequent in recent years. Our results provide a new perspective onto potential future patterns of  
480 N<sub>2</sub>O distribution and emissions in coastal areas. Continuous measurement at the BE Time-Series  
481 Station with a focus on late autumn would be of great importance for monitoring and  
482 understanding the future changes of N<sub>2</sub>O concentrations and emissions in the southwestern Baltic  
483 Sea.

## 484 **Data availability**

485 Data are available from the Boknis Eck Database: [www.bokniseck.de](http://www.bokniseck.de)

## 486 **Author contribution**

487 X.M., S.T.L. and H.W.B. designed the study and participated in the fieldwork. N<sub>2</sub>O  
488 measurements and data processing were done by X.M. and S.T.L. X.M. wrote the manuscript  
489 with contributions from S.T.L. and H.W.B.

## 490 **Competing interests**

491 The authors declare that they have no conflict of interest.

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675 Table 1. The results of the Mann-Kendall test for the surface and bottom N<sub>2</sub>O concentrations and  
676 saturations of the 12 individual months.

677 Table 1a. MKT results for N<sub>2</sub>O concentrations

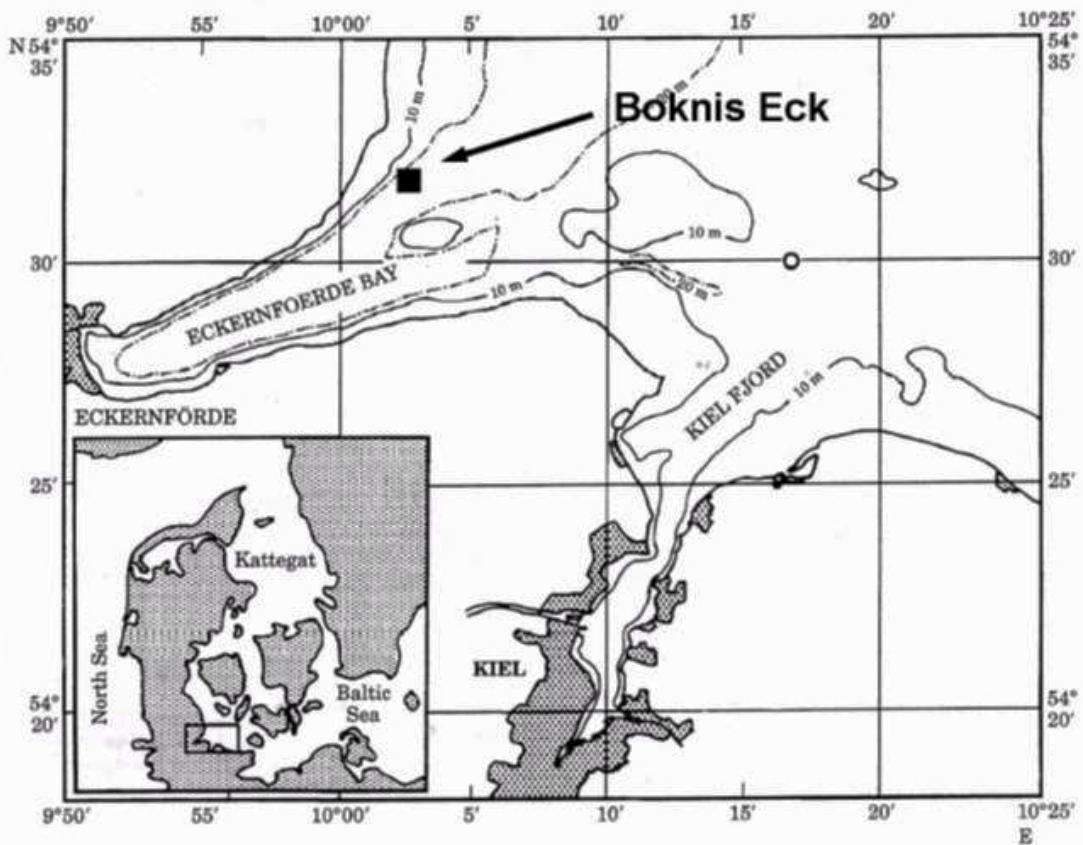
Month	January		February		March		April	
Depth/m	1	25	1	25	1	25	1	25
p	0.09	0.19	0.11	0.03(-)	0.19	0.63	0.09	0.30
Month	May		June		July		August	
Depth/m	1	25	1	25	1	25	1	25
p	0.63	0.24	0.15	0.95	0.16	0.16	0.20	0.03(-)
Month	September		October		November		December	
Depth/m	1	25	1	25	1	25	1	25
p	0.25	0.76	0.36	0.76	0.67	0.16	0.10	0.30

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Table 1b. MKT results for N<sub>2</sub>O saturations

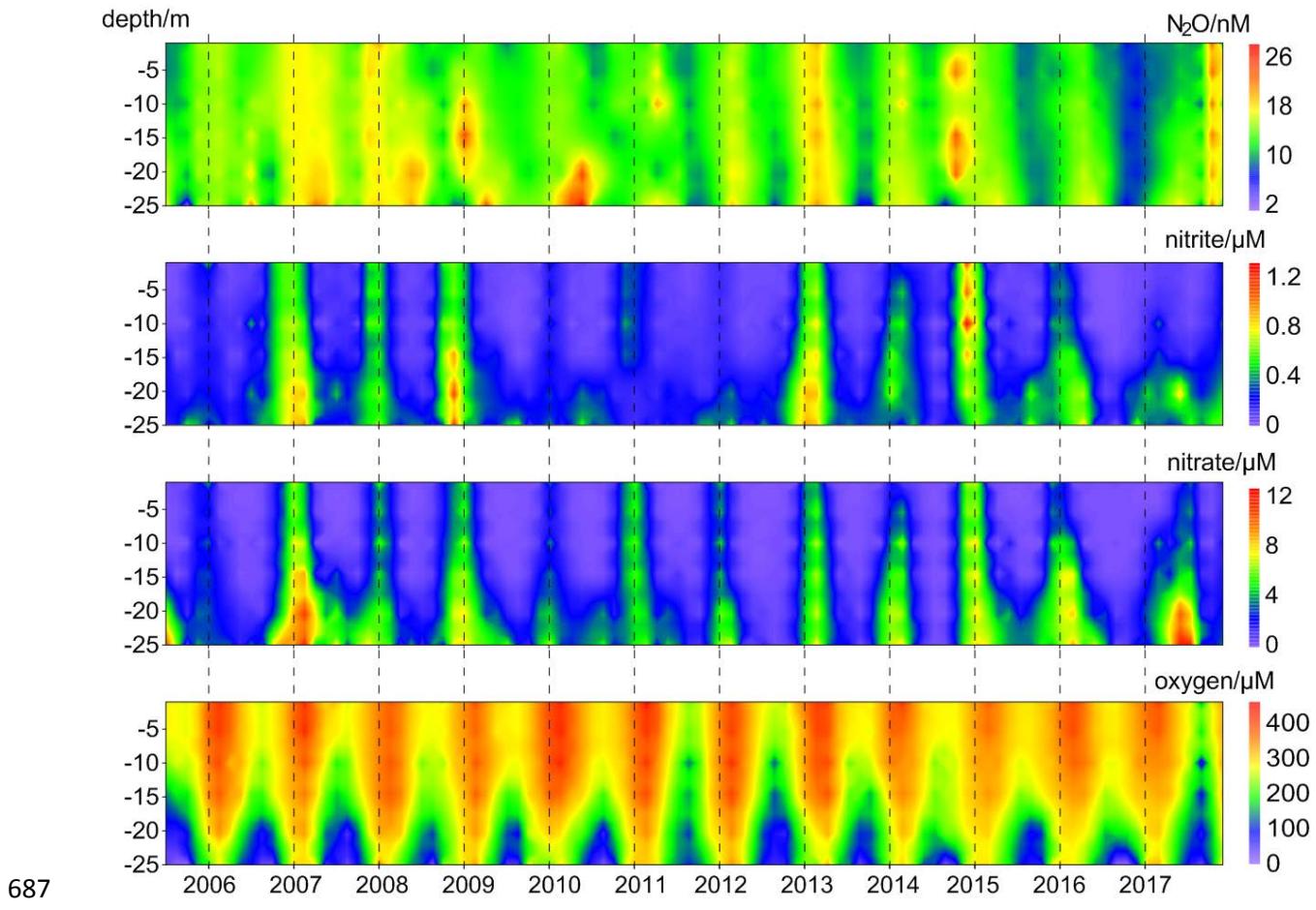
Month	January		February		March		April	
Depth/m	1	25	1	25	1	25	1	25
p	0.37	0.24	0.15	0.15	0.19	0.63	0.11	0.19
Month	May		June		July		August	
Depth/m	1	25	1	25	1	25	1	25
p	0.19	1	0.37	0.54	0.10	0.43	0.20	0.02(-)
Month	September		October		November		December	
Depth/m	1	25	1	25	1	25	1	25
p	0.04(-)	0.85	0.06	0.43	0.20	0.03(-)	0.16	0.36

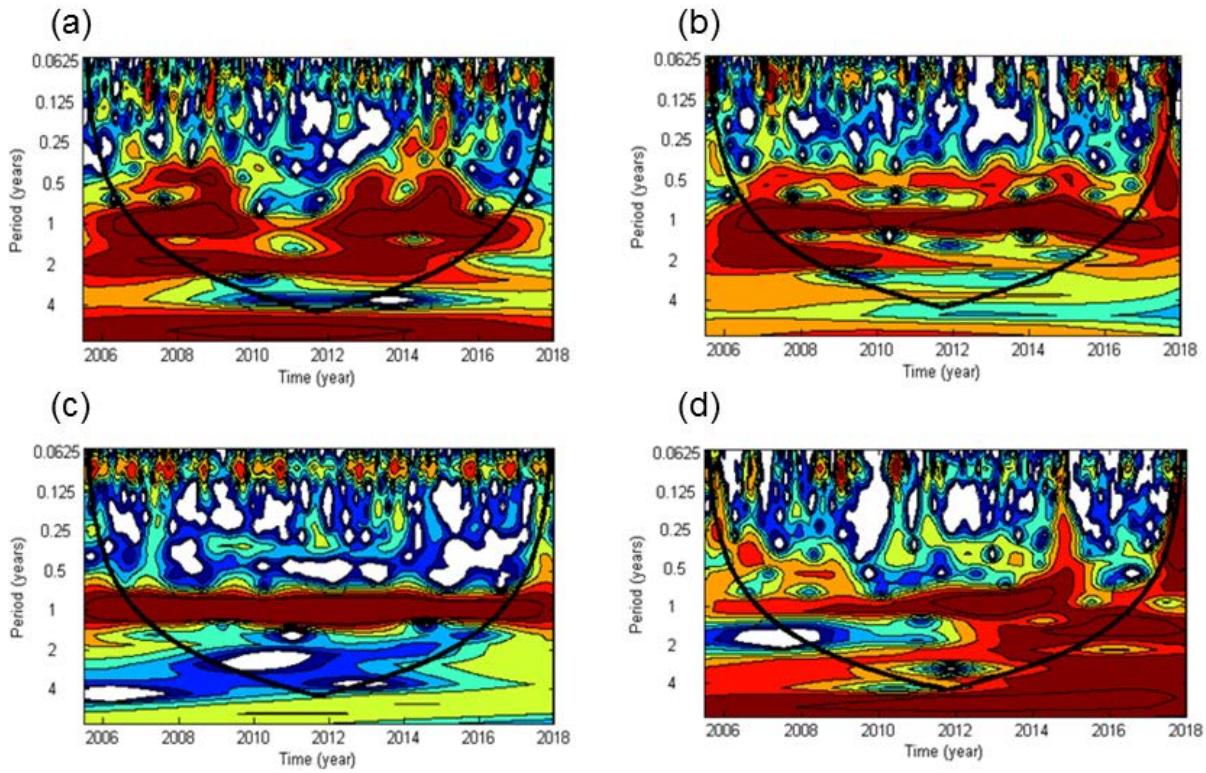
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681 p indicates the p-value of the test, which is the probability, under the null hypothesis, of obtaining a value of  
682 the test statistic as extreme or more extreme than the value computed from the sample.  
683 (-) indicates a rejection of the null hypothesis at  $\alpha$  significance level and a decreasing trend is detected.



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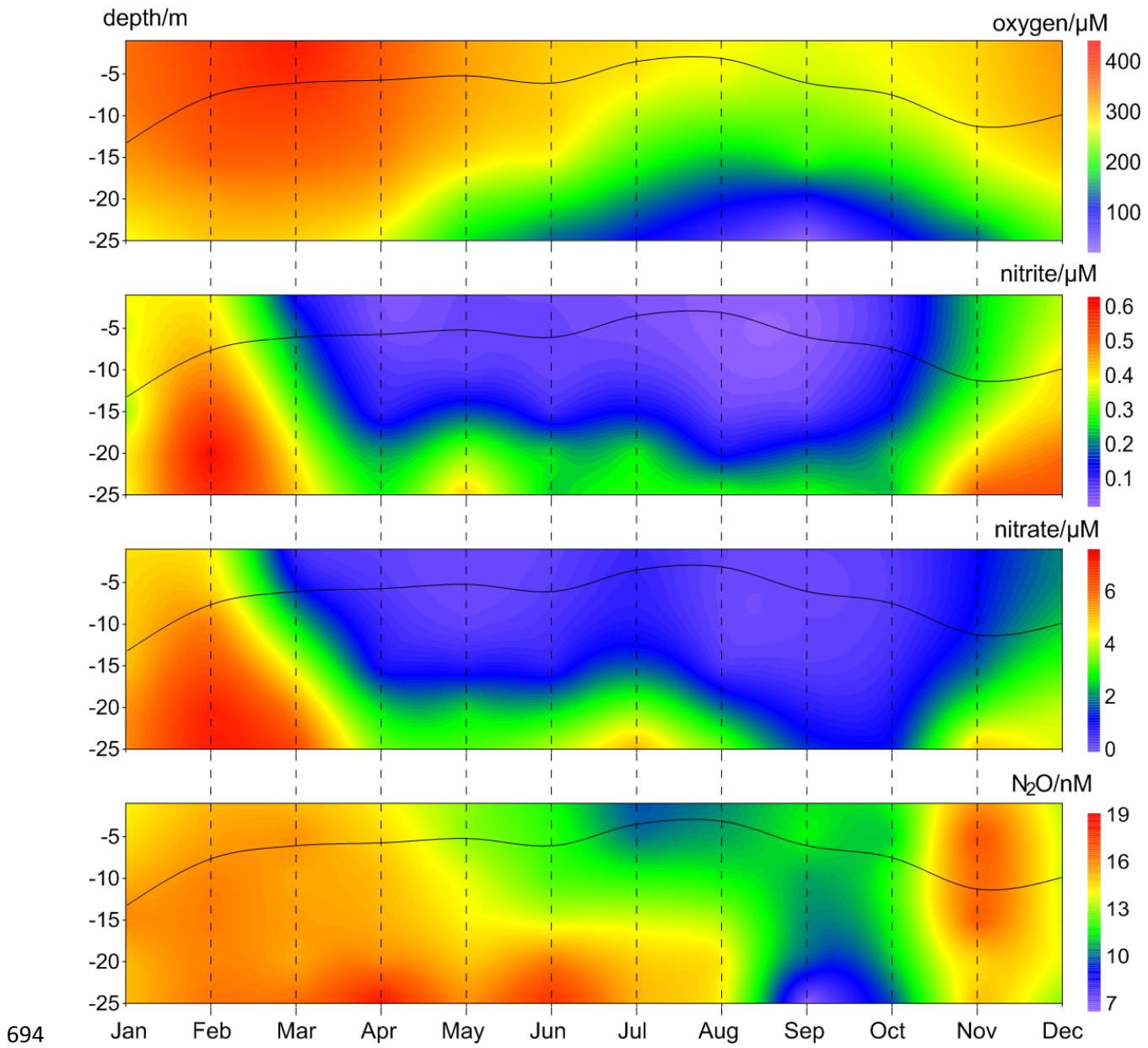
685 Fig. 1 Location of the Boknis Eck Time-Series Station in the Eckernförde Bay, southwestern Baltic Sea. (Map  
 686 from Hansen et al., 1999)

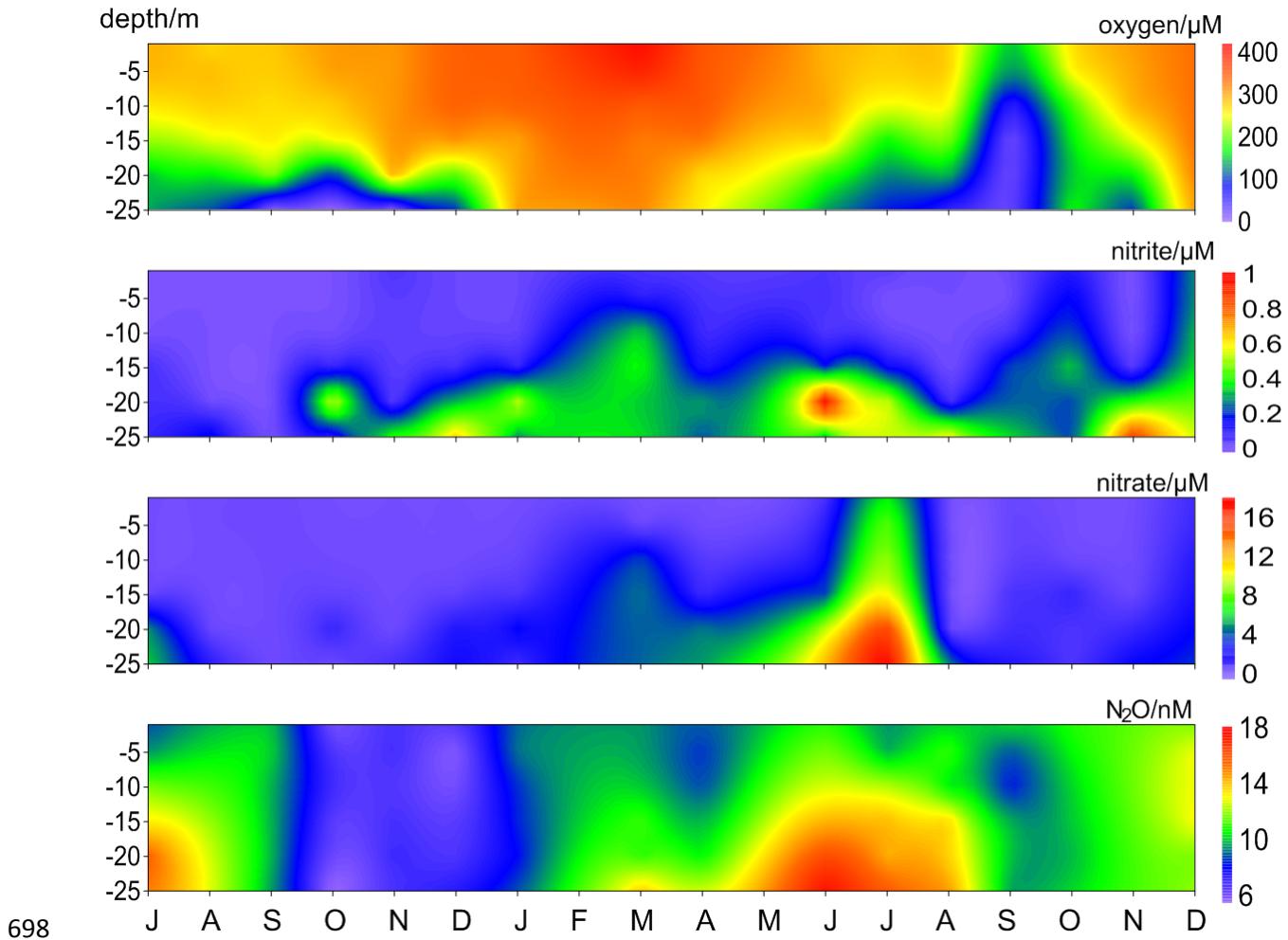




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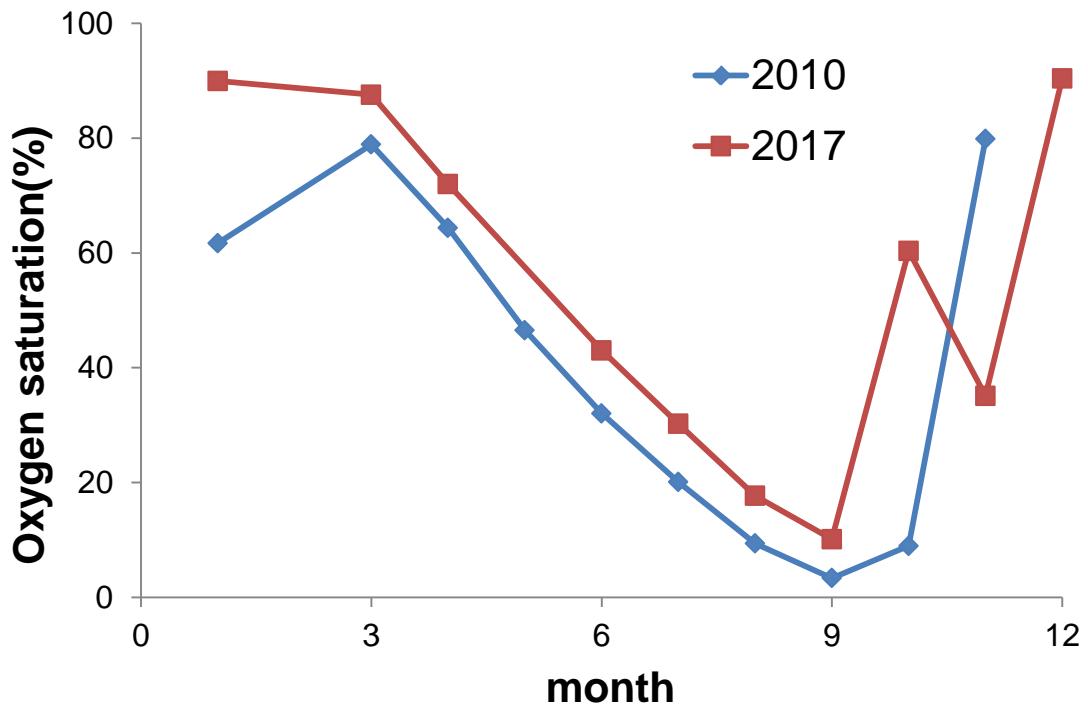
691 Fig. 3 Wavelet power spectra of  $\text{NO}_2^-$  (a),  $\text{NO}_3^-$  (b), dissolved  $\text{O}_2$  (c) and  $\text{N}_2\text{O}$  (d) from the BE Time-Series  
 692 Station. Red areas indicate high, blue indicate low power. The black conic line indicates the significant area  
 693 where boundary effects can be excluded.





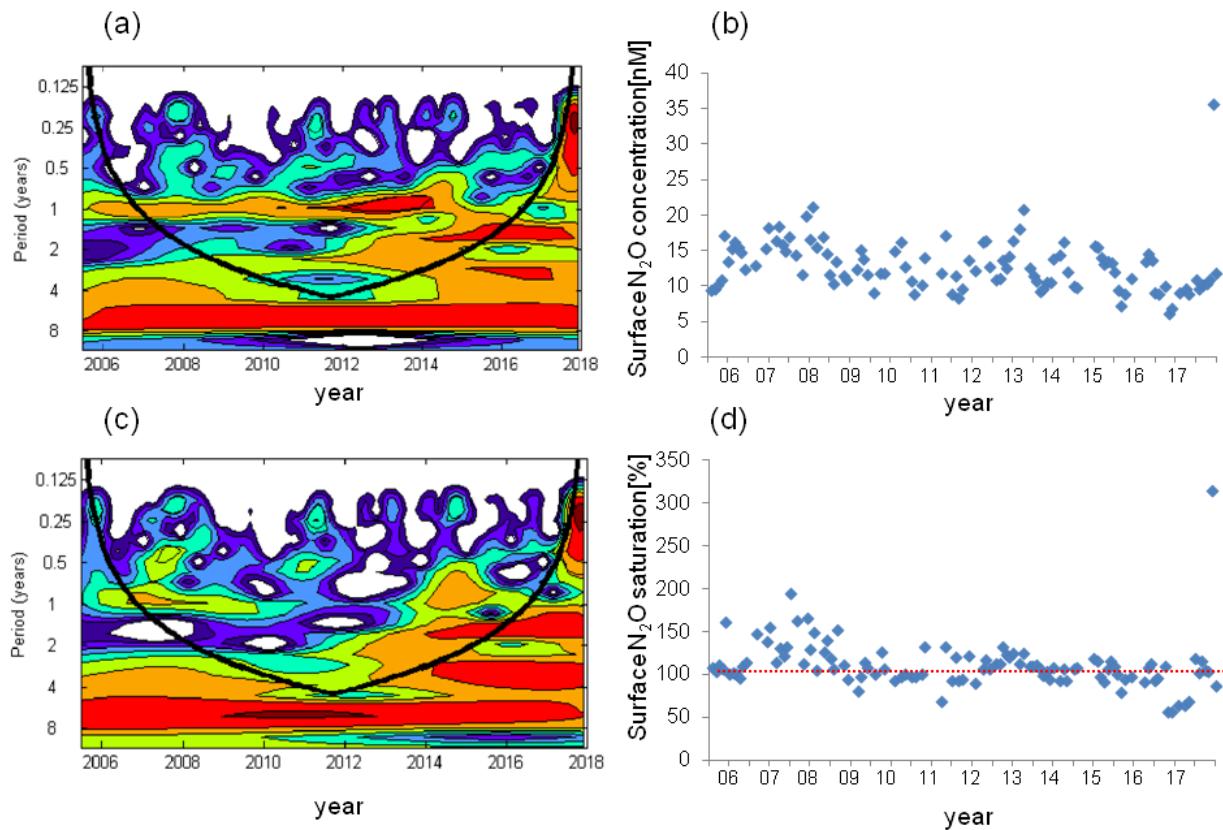
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Fig. 5 Vertical distribution of dissolved  $O_2$ ,  $NO_2^-$ ,  $NO_3^-$ , and  $N_2O$  from the BE Time-Series Station during July 2016–December 2017. Please note that the high  $N_2O$  concentrations in November 2017 were removed for better visualization.



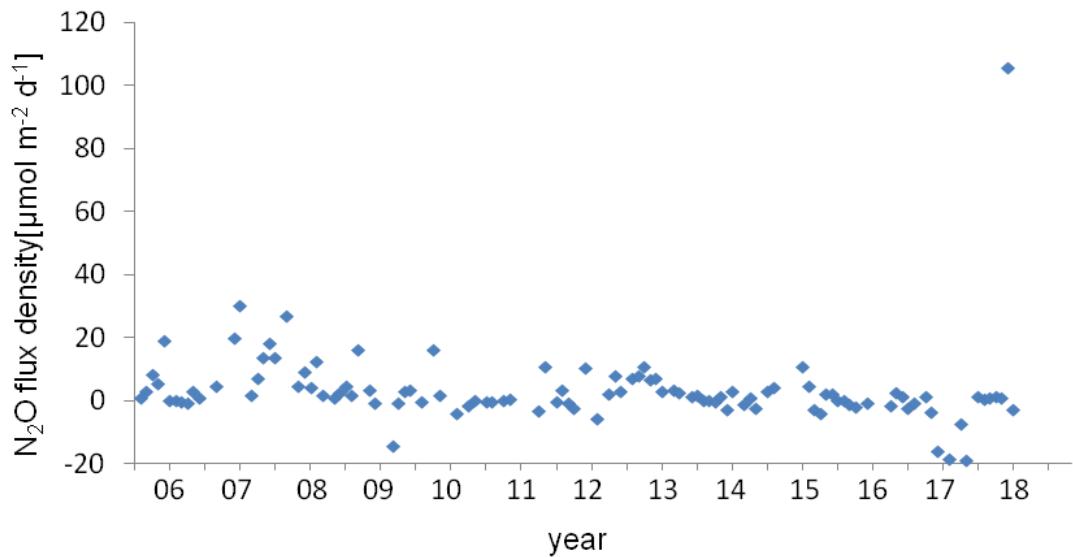
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703 Fig. 6 Variations of bottom O<sub>2</sub> saturation in 2010 (blue) and 2017 (red).



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705 Fig. 7 Wavelet analysis and the variation of surface  $\text{N}_2\text{O}$  concentrations (a, b) and surface  $\text{N}_2\text{O}$  saturations (c,  
706 d). The dashed red line in (d) indicates the saturation of 100%.



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708 Fig. 8 Variation of  $\text{N}_2\text{O}$  flux density at the BE Time Series-Station during 2005–2017. Negative values  
 709 indicated  $\text{N}_2\text{O}$  influx from the atmosphere and positive values indicated  $\text{N}_2\text{O}$  efflux to the atmosphere.