

# 1 **Response letter**

2 We thank the editor and reviewers for their constructive and helpful comments and suggestions. In the following sections, we  
3 reply to each comment individually, and explain the changes we have made to the revised manuscript. Note that we also  
4 slightly corrected the revised manuscript for stylistic issues and minor mistakes (grammar mistakes, recalculation of N<sub>2</sub>O flux  
5 densities, etc.). These changes do not affect the conclusion of the manuscript and are shown in the marked-up manuscript  
6 version. We address all comments in detail below, but would like to highlight some major changes:

- 7 (1) In accordance with comments from the editor and reviewers, we adapted the manuscript to put our research in a wider  
8 scientific context, see response below. Furthermore, we changed the title to “Seasonal variability of nitrous oxide  
9 concentrations and emissions in a temperate estuary” to address a broader audience.
- 10 (2) We recalculated N<sub>2</sub>O flux densities and emissions using four different parametrizations for the gas transfer coefficient  
11 and different wind speeds.
- 12 (3) We adapted most of our figures according to the reviewers suggestions.

13 Reviewer comments are written in bold italics, our answers are kept in plain font.

## 14 **General remarks from the editor**

15 *Thank you for submitting your paper to Biogeosciences. Two referees have evaluated your paper and provided detailed*  
16 *feedback, in particular regarding the need to improve the presentation and to better put your results in a wider context*  
17 *(beyond a case study). In your detailed rebuttal, you indicate that you will be able to resolve most issues and I therefore*  
18 *believe that a revised paper might be qualified for publication in Biogeosciences. Your revised version will likely be*  
19 *evaluated again by one or both referees.*

20 We understand the need to put our results into a wider context. To accomplish this, we focussed more on the relation between  
21 DIN and N<sub>2</sub>O as suggested by reviewer 2. In line with several other researchers (Borges et al., 2015; Marzadri et al., 2017;  
22 Wells et al., 2018), we found a limited relation between both parameters and thus, we focused on understanding the drivers for  
23 this discrepancy. Since we identified organic matter availability as a main driver for N<sub>2</sub>O production in the Elbe Estuary, we  
24 concluded that in heavily managed estuaries with high agricultural loads, N<sub>2</sub>O emissions are clearly linked to eutrophication  
25 phenomena as already proposed by Wells et al. (2018). Therefore, we rewrote and restructured parts of our abstract and  
26 introduction towards a broader research question centered on N loads as drivers of N<sub>2</sub>O production in estuaries, and modified  
27 the last section of the discussion and conclusion to address the interplay of DIN and N<sub>2</sub>O in estuaries. Furthermore, we changed  
28 the title to “Seasonal variability of nitrous oxide concentrations and emissions in a temperate estuary” to address a broader  
29 audience. We hope that these changes are sufficient to meet the reviewers’ suggestions.

30 We changed the figures in line with suggestions from both reviewers.

31 **1. Review comment (RC1) – 08.03.2023**

32 **Line 17-18: what do you mean by “compensated the effect of decreasing dissolved inorganic nitrogen (DIN) loads”?**

33 Also considering the comments of reviewer 2, we revised the manuscript to highlight the relevance of our research to a broader  
 34 audience. Thus, we rewrote our abstract, focusing the relevance for a broader scientific community and highlighting the  
 35 connection between eutrophication and N<sub>2</sub>O emissions. We also rewrote this phrase.

Lines	Change
L24 - 29	Changed to: “A comparison with previous measurements in the Elbe Estuary revealed that N <sub>2</sub> O saturation did not decrease alongside with DIN concentrations after a significant improvement of water quality in the 1990s that allowed for phytoplankton growth to reestablish in the river and estuary. This effect of phytoplankton growth and the overarching control of organic matter on N <sub>2</sub> O production, highlights that eutrophication and agricultural nutrient input can increase N <sub>2</sub> O emissions in estuaries.”

36 **Line 25: How does 0.24±0.06 Gg N<sub>2</sub>O y<sup>-1</sup> emission compare to global estuarine N<sub>2</sub>O emission?**

37 We changed the emission calculation as suggested by both reviewers. We removed the emission estimate from the abstract  
 38 focusing more on the relation with DIN loads and seasonal varying drivers in the Elbe Estuary, which lead to year-round high  
 39 N<sub>2</sub>O emissions. We highlighted the relevance for a broader scientific community by focusing on the connection between N<sub>2</sub>O  
 40 emissions and DIN loads, as well as the linkage to eutrophication in estuaries with high agricultural loads. In the new section  
 41 4.5 of our discussion, we now compare N<sub>2</sub>O emission estimates and the resulting N<sub>2</sub>O:DIN relation across estuaries.

Lines	Change
	Removed 0.24 ± 0.06 Gg N <sub>2</sub> O yr <sup>-1</sup> from the abstract
L262-265	Comparison of N <sub>2</sub> O saturation with other estuaries
L421-434	Comparison of N <sub>2</sub> O emissions and N <sub>2</sub> O:DIN relation with other estuaries

42

43 **Lines 40-42: Denitrification could also occur in anoxic water column contributing to N<sub>2</sub>O production (Ji et al., 2018; Tang**  
 44 **et al., 2022).**

Lines	Change
L44-45	Added denitrification in the water column as possible production pathway.

45

46 **Line 44: specify Port Hamburg as the third largest port in Europe.**

Lines	Change
L55	Changed “biggest” to “largest”

47

48

49 **Line 70: how deep is the Elbe estuary? This gives an idea if sedimentary processes (e.g., N<sub>2</sub>O production) may affect N<sub>2</sub>O**  
 50 **concentration in the surface water column.**

Lines	Change
L82, L84-85	Added information about the depth of the Elbe Estuary in our study site description

51

52 **Figure 1: There are too many city names on the map, which is distractive. It may be clearer to label only the key cities like**  
 53 **Cuxhaven or island Scharhorn where the Elbe River enters the North Sea or Oortkaten.**

Lines	Change
L86	Changed Map (Fig. 1)

54

55 **Lines 85-87: Why transect sampling was performed after high tides? What's the effect of tides on N<sub>2</sub>O concentration? Tidal**  
 56 **cycles of N<sub>2</sub>O concentration have been observed in other estuaries (Goncalves et al., 2015; Barnes et al., 2006).**

57 We chose our sampling strategy (upstream against the outgoing tide) to prevent interference of tidal effects on our  
 58 measurements. Our aim was to obtain comparable data for each cruise at similar tidal phase, with comparable current and  
 59 mixing conditions. We started after high-tide and travelled against the outgoing tide to make sure that we did not move with  
 60 the same water masses while travelling upstream.

61 Tidal effects will very likely affect nitrous oxide concentrations in the Elbe estuary, but this is not the focus of the present  
 62 manuscript. We briefly addressed possible tidal effects in the revision.

Lines	Change
L94-95	Explained our chosen sampling strategy
L416-420	Addressed the possible effects of tides, diel variations and currents on N <sub>2</sub> O emissions

63

64 **Line 116 in Equation 1: is N<sub>2</sub>O<sub>cw</sub> the partial pressure of N<sub>2</sub>O in water? Otherwise, the saturation should be calculated as**  
 65 **the N<sub>2</sub>O<sub>cw</sub>/N<sub>2</sub>O<sub>eq</sub>\*100 where N<sub>2</sub>O<sub>eq</sub> is the equilibrated N<sub>2</sub>O concentration with atmosphere. Similarly in Equation 3.**  
 66 **N<sub>2</sub>O<sub>air</sub> should be N<sub>2</sub>O<sub>eq</sub>.**

67 For our calculations, we used the average atmospheric N<sub>2</sub>O concentrations measured on each specific day of the cruise to  
 68 calculate expected atmospheric equilibrium concentrations considering the solubility function of Weiss and Price (1980) and  
 69 atmospheric pressure.

Lines	Change
L125-126	Changed: “and in the air (N <sub>2</sub> O <sub>air</sub> )” to “atmospheric equilibrium concentrations (N <sub>2</sub> O <sub>eq</sub> )”
L127	Changed Eq. 1: “N <sub>2</sub> O <sub>air</sub> ” to “N <sub>2</sub> O <sub>eq</sub> ”

70

71

72 ***Line 143: Why nitrate concentration increased at 700 km? Are there tributaries or point sources?***

73 We regard this a result of nitrification, rather than a point source. We now refer to potential point sources in the revised version  
74 (see below). Dähnke et al. (2008) identified the Elbe estuary along its salinity gradient as a significant source of nitrate with  
75 high nitrate production in the maximum turbidity zone (MTZ). Sanders et al. (2018) measured highest nitrification in the  
76 Hamburg port region, which was not covered in the research done by Dähnke et al. (2008). Both studies highlighted the  
77 importance of nitrification along the Elbe estuary. Further, our results in section 4.2 showed ongoing nitrification fueled by  
78 marine organic matter along the mesohaline estuary and indicated that nitrate is produced by coupled remineralization with  
79 nitrification from both the riverine and marine site of the estuary.

80 We believe the offset between region of highest nitrification rates (Port of Hamburg) and nitrate peak in the estuary (stream  
81 kilometer 680-700) is a result of different spatiotemporal scales, as suggested by Sanders et al. (2018). They found that the  
82 position of the nitrate maximum and nitrate gain over the entire estuary depended on the processing rates and was thus coupled  
83 to discharge conditions. However, we decided not to address the offset in our manuscript. Hopefully, the added information  
84 regarding the point sources will help to clarify the text, but addressing the nitrate dynamics in more detail is beyond the scope  
85 of the paper.

Lines	Change
L78-79	Added information about point sources to study site description

86

87 ***Lines 148-149: Why ammonium and nitrite concentration increased near Hamburg Port? Is it due to internal organic***  
88 ***matter remineralization or point sources or sedimentary flux?***

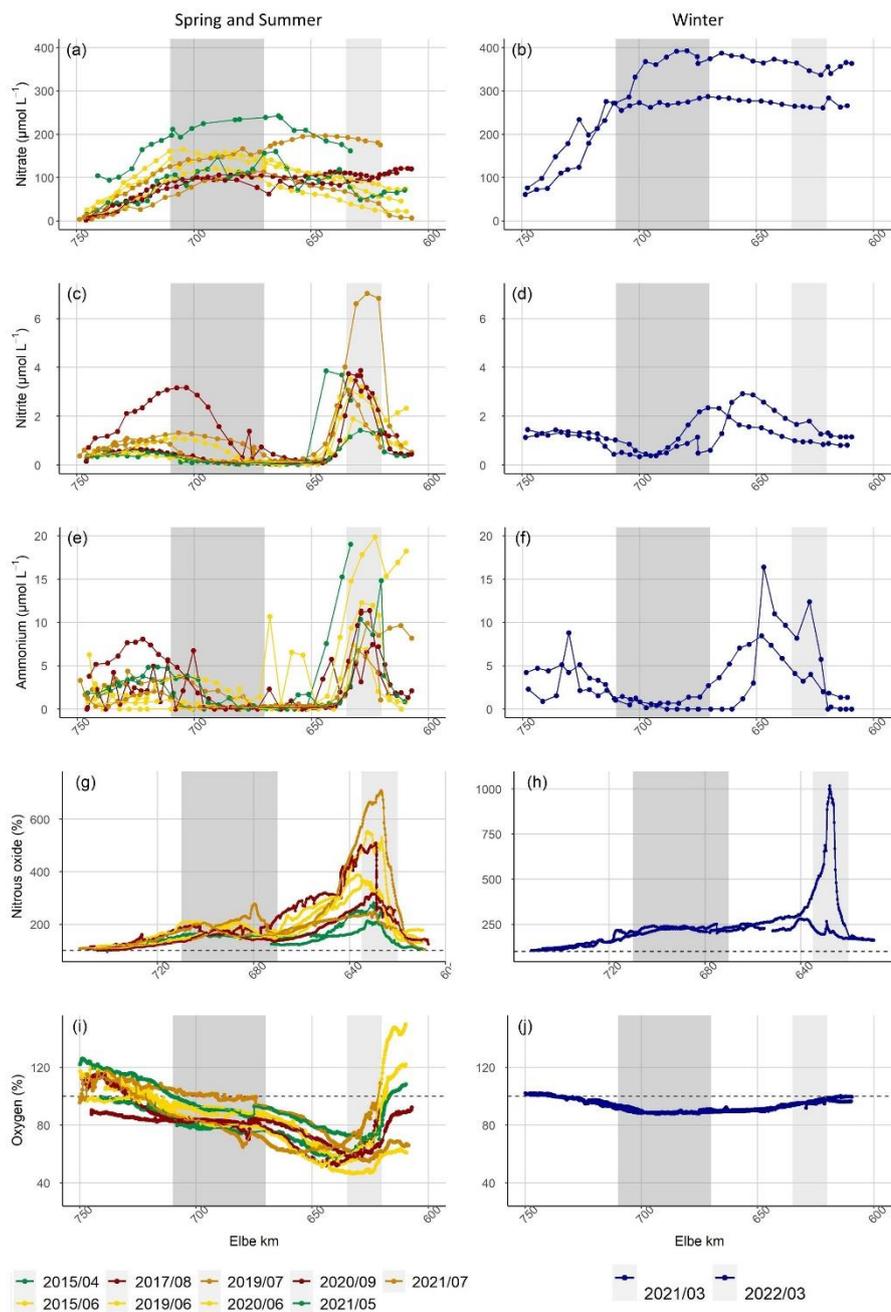
89 We restructured section 4.3 addressing possible nitrogen turnover processes and benthic-pelagic coupling in more detail.  
90 Therefore, we added a short paragraph to clarify succession of nitrogen turnover in the Port of Hamburg.

Lines	Change
L344-348 L352-356	Summarized state of research regarding nitrogen turnover in the Port of Hamburg

91

92 ***Figure 2: It is hard to tell the difference among each cruise with so many colored lines. How about presenting data from***  
93 ***the same season using the same color to illustrate the seasonality as a supplementary figure?***

94 We tried to implement the suggestion of the reviewer. However, we felt that the figure did not help to illustrate seasonality  
95 and therefore we decided not to include it into the supplements (see figure below).



96

97 **Figure 1: Salinity along the Elbe estuary (a) in spring/summer and (b) in winter. Suspended particulate matter (SPM) concentration**  
 98 **in ( $\text{mg L}^{-1}$ ) along the Elbe estuary in (c) spring/summer and (d) in winter. Particulate carbon to nitrogen ratio (C/N) along the Elbe**  
 99 **estuary in (e) in spring/summer and (f) in winter. Particulate nitrogen (PN) content in (%) in (g) spring/summer and (h) winter.**  
 100 **Particulate carbon (PC) content in (%) in (i) spring/summer and (j) winter. All values are plotted against stream kilometers. The**  
 101 **Hamburg port region is shown with a gray background. C/N ratios were measured with an Elemental Analyzer (Eurovector EA**  
 102 **3000) calibrated against a certified acetanilide standard (IVA Analysetechnik, Germany). The standard deviation was 0.05% and**  
 103 **0.005% for carbon and nitrogen respectively. Please note that there are no data for the suspended particulate matter composition**  
 104 **in 2015.**

105 *Lines 211-218 and lines 232-234: Figure 4 a and b are both from June, summer. The linear positive relationship between*  
 106 *AOU and excess N<sub>2</sub>O suggests N<sub>2</sub>O production from nitrification (e.g., Nevison et al., 2003). The increase in the slope*  
 107 *should be interpreted as an increase in the N<sub>2</sub>O production yield or external N<sub>2</sub>O input (e.g., point source).*  
 108 We indeed identified nitrification as responsible production processes both in the mesohaline estuary (section 4.2) and the  
 109 Hamburg port region (section 4.3). In the revised manuscript, we clarified that N<sub>2</sub>O yield varied due to changes in production,  
 110 rather than point sources. As stated above, we also addressed the role of point sources in the study site description.

Lines	Change
L78-79	Added information about point sources to study site description
L236	Presented all plots of AOU vs N <sub>2</sub> O <sub>xs</sub> in a revised version of Fig. 3
L237-239	Changed figure caption to match new Fig. 3
L229-235	Changed to: “Plots of excess N <sub>2</sub> O (N <sub>2</sub> O <sub>xs</sub> ) and apparent oxygen utilization (AOU) revealed excess N <sub>2</sub> O along the entire estuary (Fig. 3). During all cruises, elevated riverine N <sub>2</sub> O <sub>xs</sub> entered the estuary (stream kilometer < 620). A linear positive relationship between N <sub>2</sub> O <sub>xs</sub> and AOU suggested nitrification as main production pathway in large sections of the estuary (Nevison et al., 2003; Walter et al., 2004). However, in summer, a change of slope in the Port of Hamburg as well as in the mesohaline section of the estuary suggested either increased in-situ N <sub>2</sub> O production or external N <sub>2</sub> O input. In winter, we found an increasing slope in the Hamburg Port region and in the oligohaline part of the Elbe Estuary (Fig. 3h, k).“
L265-267	Rewritten: “The relation of N <sub>2</sub> O <sub>xs</sub> and AOU (Fig. 3), with changing slopes in the Port of Hamburg and mesohaline estuary, was determined by either initial riverine N <sub>2</sub> O production, or in-situ production along the estuary”
L286-287	Rewritten: “The N <sub>2</sub> O peak in the transition between oligohaline and mesohaline estuary was accompanied by a sudden change in the slope of the AOU vs N <sub>2</sub> O <sub>xs</sub> plots, (Fig. 3), pointing towards N <sub>2</sub> O production in the oxic water column”

111  
 112 *Figure 4: It would be interesting to systematically/statistically assess the relations between excess N<sub>2</sub>O and environmental*  
 113 *factors like salinity (non-conservative behavior of N<sub>2</sub>O) or dissolved inorganic nitrogen (infer N<sub>2</sub>O production pathways),*  
 114 *PN, PC, and SPM. There seems to be a good relation between N<sub>2</sub>O and ammonium/nitrite concentration shown in Figure*  
 115 *2.*

116 We understand that a systematically and statistical assessment of the relations would help the reader to follow our discussion.  
 117 We did assess the statistical relations in sections the previous version of our discussions, but for clarity, we added a section  
 118 regarding the statistical analysis in the results chapter in the revised manuscript.

119 During data interpretation, we tested diverse presentation and analysis methods and found that regressions were not necessarily  
 120 well suited to visualize, describe and analyze our data. Correlations were distorted by the spatial offset between the ammonium,  
 121 nitrite and N<sub>2</sub>O peaks, which we attribute to a succession of nitrogen bearing substances during turnover processes like

122 nitrification. Thus, we chose another way to visualize the data in Fig. 4 (L297) and Fig. S3-S13 of the supplementary material.  
123 Furthermore, we addressed the relations of N<sub>2</sub>O and various forms of nitrogen to identify N<sub>2</sub>O production processes and their  
124 controls in the discussion (e.g. L292-L294, L314-L315, and L356-357).

<b>Lines</b>	<b>Change</b>
L160-162	Added method section regarding statistical analysis
L240-259	Added result section regarding statistical analysis

125

126 ***Lines 242-243 and Figure 5: What about the variations of the N<sub>2</sub>O%, oxygen and total nitrogen concentration? The***  
127 ***riverine N concentration is decreasing, what about the changes in other point sources of N input along the estuary (e.g.,***  
128 ***from wastewater treatment plants) or concentration in the estuary?***

129 We agree with the reviewer that a more detailed analysis of a long-term trend of N<sub>2</sub>O concentrations and reasons for changes  
130 would be very interesting. However, for our study we focus on seasonal variations rather than a long-term trend analysis. With  
131 section 4.1, we aimed to compare our results with a broader spatial and temporal scale by including a short comparison to other  
132 estuaries as well as with previous measurements from the Elbe estuary. This gives a hint towards temporal trends, but seasonal  
133 variability and data coverage make the long-term data difficult to interpret.

134 Briefly, the biogeochemical processes occurring in the Elbe estuary have drastically changed over the last 50 years: (1) the  
135 reunification of Germany and the collapse of East-German industry had led to significant improvements of water quality (e.g.  
136 Guhr et al., 2000). (2) The decision to combat eutrophication in the North Sea in the 1980s and (3) improved waste water  
137 management resulted in a significant reduction of riverine nutrient loads (de Jong, 2007, p.2019; Van Beusekom et al., 2019;  
138 Bergemann and Gaumert, 2010). Dähnke et al. (2008) showed that this led to a change of dominating denitrification towards  
139 significant nitrification in the Elbe estuary. Thus, a profound long-term analysis would be in need of its own paper, for which  
140 we think our data coverage is insufficient, also considering the seasonal variability. As an example, measurements from only  
141 one cruise are available for the 1990s.

142

<b>Lines</b>	<b>Change</b>
L78-79	Added: “Point sources along the estuary provide only small part of the total nitrogen input to the Elbe Estuary (Hofmann et al., 2005; IKSE, 2018)”
L265	Added comparison to N <sub>2</sub> O saturation with other highly modified urban systems (Reading et al. 2020)
L279-280	Included reference to Dähnke et al. (2008) and change of dominating denitrification towards significant nitrification in the Elbe estuary
Fig. S2	Removed Fig. 4 from text and added it to supplementary material with more plot panels and adapted figure caption
L275-278	Added: “However, since the BIOGEST study in 1997 (Barnes and Upstill-Goddard, 2011), N <sub>2</sub> O remained relatively stable at ~ 200 % saturation despite a concurrent decrease in TN concentration from ~400 μmol L <sup>-1</sup> to around 200 μmol L <sup>-1</sup> (Fig. S2, Hanke and Knauth, 1990; Barnes and Upstill-Goddard, 2011; Brase et al., 2017; FGG, 2021).” instead of the figure to the text.

143

144 **Line 272: “this suggests”**

<b>Lines</b>	<b>Change</b>
L305	Changed to “suggests”

145

146 **Line 273: how is MTZ defined? What threshold of suspended particle material is used to define the MTZ?**

147 Generally, the occurrence of an MTZ is unique to each estuary and is generated by the balance between river-induced flushing  
 148 and upstream transport of marine SPM, as well as a function of estuarine geomorphology, gravitational circulation and tidal  
 149 flow, trapping the particles in the MTZ (Bianchi, 2007; Sommerfield and Wong, 2011; Winterwerp and Wang, 2013). Thus,  
 150 the MTZ is usually located in the onset of the salinity gradient of an estuary (Burchard et al., 2018).

151 The MTZ is – also in literature – often assessed based on relative changes in SPM or turbidity, and is mostly located between  
 152 stream km 670 and 710 (e.g. Bergemann, 2004) in the Elbe estuary. During our cruises, SPM and turbidity were not always  
 153 measured consistently, depending on instrument and personnel availability – during some cruises suspended particulate matter  
 154 concentrations were measured using filtration techniques, during some cruises we obtained the data from turbidity sensors,  
 155 which are not entirely intercomparable. Therefore, we did not define a threshold of suspended particulate matter to define the  
 156 MTZ, but used relative changes of SPM or turbidity for MTZ identification.

157 We added color bars indicating the relative change of SPM concentrations to Fig. 4 and supplement material.

<b>Lines</b>	<b>Change</b>
L298	Added color bars to Fig. 4
Fig. S3-S13	Added color bars to each figure

158

159 **Line 287 and 296-297: clarify the reference: Kappenberg and Fanger, 2007 (German?) and source of organic matter from**  
160 **the North Sea into the Elbe estuary.**

161 We included a peer-reviewed reference (Schoer, 1990) that shows an upstream transport of suspended matter into the Elbe  
162 estuary due to tidal transports in the Elbe estuary.

Lines	Change
L320	Included a new reference: (Schoer, 1990)

163

164 **Lines 311-313: How about showing the relations between ammonium, nitrite and N<sub>2</sub>O in figures?**

165 We decided to show the relation between ammonium, nitrite and N<sub>2</sub>O plotted against stream kilometers in Fig. 4 and  
166 Fig. S3-S13. We found that the spatial progression of nitrogen containing substances were more illustrative than scatter plots  
167 or correlations for each substance, cruise and production areas. These relations were distorted by the spatial offset between the  
168 occurring ammonium, nitrite and N<sub>2</sub>O peaks, which we explain by a succession of nitrogen bearing substances during turnover  
169 processes like nitrification (e.g. section 4.2). Therefore, we find our choice of presentation better suited. To address this issue,  
170 though, we added a section about the statistical analysis and relations of individual parameters in the results section. We further  
171 rephrased the statement in L311-313).

Lines	Change
L160-162	Added method section regarding statistical analysis
L240-259	Added result section regarding statistical analysis
L361-364	Changed to: “Overall, our data showed the succession of ammonium, nitrite and N <sub>2</sub> O production (Fig. 4 and supplementary material S3-S13) confirming simultaneous denitrification and nitrification responsible pathways for N <sub>2</sub> O production in the Port of Hamburg (Brase et al. 2017).”

172

173 **Line 315: What are R values? R is positive for nitrite concentration.**

174 R is the Pearson correlation coefficient. We added a short section about our statistical analysis in the *Methods* section of our  
175 manuscript. Nitrite concentrations correlated positive with N<sub>2</sub>O leading to a positive correlation coefficient.

Lines	Change
L160-162	Added method section about statistical analysis

176

177 **Line 320-321: Is nitrification responsible for the remaining oxygen consumption?**

178 Yes, nitrification is responsible for the remaining oxygen consumption. We have clarified this in the text.

Lines	Change
L369	Added: “whereas the remaining 25 % stem from nitrification (Schöl et al., 2014; Sanders et al., 2018)”

179

180 ***Line 326 and Figure S1: why C/N ratio was so high in 2021 March?***

181 We double-checked our measurements, and the data appear correct and sound. We have no easy explanation at hand, but  
182 speculate that a calcareous algae bloom in the North Sea might be a potential cause. However, we have no further evidence  
183 for this hypothesis. We will not address this further, as the C/N ratios are not a crucial parameter for our discussion. However,  
184 we would like to keep the data in the manuscript as they might be insightful for later research and other researchers.

185 ***Line 345-347: “Ammonium and N<sub>2</sub>O concentrations are high in the pore water of underlying sediments”. Reference or  
186 example of the concentration. What about the timing of deepening and dredging works in the Hamburg Port compared to  
187 the cruise periods?***

188 The Elbe estuary is constantly deepened and dredged along the entire transect to grant access for big container ships. However,  
189 we did not compare operation locations with the measured N<sub>2</sub>O concentrations except for our March cruises, as we did not see  
190 big differences in the spatial variation of the N<sub>2</sub>O profiles, which were not explainable by in-situ production, nor found strong  
191 correlations between N<sub>2</sub>O and suspended particulate matter concentrations.

Lines	Change
L82	Added: “The Elbe Estuary is dredged year-round”
L340-342	Added references and rephrased: “Ammonium concentrations in the sediment pore water are high (Zander et al., 2020, 2022) and N <sub>2</sub> O can be produced by nitrifier-denitrification in the sediments (Deek et al., 2013)”
L331-350	Moved from section 4.4 to 4.3
L389-410	Shortened discussion about possible effects of deepening and dredging in section 4.4

192

193 ***Lines 360-361: Has there been any N<sub>2</sub>O measurement from this wastewater treatment plant (WWTP) Köhlbrandhöft? The  
194 ammonium concentration in 2021/03 is not exceptionally high compared to previous cruises (e.g., 2020/06). What about  
195 the direct N<sub>2</sub>O output from the wastewater treatment plant?***

196 The operators of the WWTP measured N<sub>2</sub>O concentrations during our cruise, but did not detect elevated N<sub>2</sub>O concentrations.  
197 However, direct N<sub>2</sub>O output from the WWTP was not measured and the increased ammonium loads leaving the WWTP  
198 corresponded to the measured increase of N<sub>2</sub>O concentration. We assume that excess N<sub>2</sub>O is not produced within WWTP itself,  
199 but stems from elevated ammonium concentrations in the Elbe that are introduced with warmer waste water.

200 We likely did not see an extraordinary ammonium peak due to the distance of the WWTP outflow and our measurement  
201 transect, ~ 2 km. The outflow of the WWTP is located in the Southern Elbe, which joins the sample stretch at stream kilometer  
202 626. Ammonium is probably rapidly converted to N<sub>2</sub>O as the warmer and biological active waste water enters the Elbe estuary  
203 before it reaches the Northern Elbe (and our sampling site).

204 We were admittedly surprised by the extraordinary N<sub>2</sub>O concentrations in March 2021, and even more so when the results  
205 were not reproduced in the following year. Consequently, we concluded that an extraordinary event must have caused the N<sub>2</sub>O

206 peak. Since we could not detect any relation to the deepening and dredging work in the Port area and our measurements fitted  
 207 to extraordinary operation condition in the WWTP, we assumed this might be the source, especially as our hypothesis was  
 208 confirmed by the WWTP operators. Additional measurements confirmed that N<sub>2</sub>O concentration was not elevated near the  
 209 WWTP outlet under normal conditions.

Lines	Change
L399-410	Revised: “Another possible source of N <sub>2</sub> O is the WWTP outflow in the Southern Elbe that joins the main estuary at stream kilometer 626 (Fig. 1), matching the N <sub>2</sub> O peak at stream kilometer 627 (Fig. 2h). As explained above (section 4.3), the effect of this WWTP on N <sub>2</sub> O saturations under normal conditions should be negligible. This peak can be the result of an extraordinary event during our sampling. We indeed found that an extreme rain event occurred on March 11 <sup>th</sup> 2021 (HAMBURG WASSER, pers. Comm., Laurich 2022) with a statistical recurrence probability of one to five years ( <a href="https://sri.hamburgwasser.de/">https://sri.hamburgwasser.de/</a> , last access: 04.04.2023). This rare event caused aggravated operation conditions in the WWTP at the time of sampling. While the operators could still meet the limits for the effluent levels of nitrate and ammonium, higher than usual ammonium loads exited the treatment plant at this time. We assume that these elevated ammonium WWTP loads, were rapidly converted to N <sub>2</sub> O as the warmer and biologically active waste water entered the Elbe Estuary in March 2021. An important factor for aggravated conditions was a temperature drop in the WWTP caused by cold rain water, we hypothesize that a similar rain event in warmer months would not lead to comparable N <sub>2</sub> O peaks.”

210

211 ***Figure 7. Use month or season as the x axis instead of cruise number? Add description of the boxplot. Why not adding***  
 212 ***error bars for emissions?***

213 As we restructured section 4.5 of our discussion, we also changed the figure. However, we considered the comment of the  
 214 reviewer including description of the boxplots.

Lines	Change
L442	Deleted Fig. 7 included new Fig. 5
L443-447	Included description of boxplots in figure caption.

215

216 ***Table 3. How is annual N<sub>2</sub>O emission calculated? Since there is a seasonal variation in the N<sub>2</sub>O flux, monthly or seasonal***  
 217 ***N<sub>2</sub>O emission may be more representative. Because N<sub>2</sub>O flux was measured at a high spatial resolution, it may be useful to***  
 218 ***calculate the N<sub>2</sub>O flux across the whole estuary by integrating the flux and area section by section (e.g., River section,***  
 219 ***Hamburg port, Oligohaline section) instead of multiplying the average N<sub>2</sub>O flux by the whole area of Elbe estuary.***

220 We recalculated emissions as suggested by the reviewer: We separated the Elbe estuary into five regions: limnic (stream  
 221 kilometer 585 to 615), Port of Hamburg (stream kilometre 615 to 632), oligohaline (stream kilometre 632 to 704), mesohaline  
 222 (stream kilometre 704 – 727) and the polyhaline section (stream kilometre 727 to 750). Respective areas are found in the

223 supplementary material S6. For seasonality, we divided our cruises: winter (March), spring (April and May), summer (June  
 224 and July) and late summer/autumn (August and September). Following this, we calculated daily emissions for each section  
 225 and each season. To upscale to annual emissions, we applied our calculated emissions estimates to months without  
 226 measurements (winter: January to March and November to December, spring: April to May, summer: June to July and late  
 227 summer/autumn: August to October).

228 In line with reviewer 2, we also considered different wind speeds and parameterizations to calculate the gas transfer coefficient  
 229 for flux densities calculation and emission estimates. Thus, we rewrote the results section and included our emissions estimates  
 230 in section 3.3 “N<sub>2</sub>O flux densities and emissions”.

<b>Lines</b>	<b>Change</b>
L142-151	We included a detailed description of the calculation in the “Method” section
L204-220	We included detailed results in the “Results” section (results section was rewritten to include new emission calculations)
Table S2	We included flux-densities calculations using other parametrizations and wind speeds in the supplementary material
L262 and L414-434	We changed flux densities and N <sub>2</sub> O emission estimates in the revised manuscript so that they fit to the new calculations

231

232 ***Line 406: Why do you think there is no seasonality in N<sub>2</sub>O emission? N<sub>2</sub>O flux is different comparing spring, summer and***  
 233 ***winter shown in Table 3.***

234 For the revised manuscript, we calculated emissions as suggested by the reviewer and described above. We restructured our  
 235 last section of the discussion 4.5 in line with suggestions of the other reviewer focusing on the relevance of our research for a  
 236 broader audience, investigating the N<sub>2</sub>O:DIN relation discussing seasonal changing drivers for N<sub>2</sub>O production and emissions.  
 237 Thus, we also changed our abstract and conclusion.

<b>Lines</b>	<b>Change</b>
L413-447	Restructured and rewritten section 4.5
	Removed from abstract: “Surprisingly, estuarine N <sub>2</sub> O emissions were equally high in winter and summer”
	Removed from conclusion: “We saw no seasonality in N <sub>2</sub> O emissions, ...”

## 238 **2. Review comment (RC2) – 04.04.2023**

239 ***1. Converting dissolved concentrations to emissions: Like many studies, here the authors measured the dissolved***  
 240 ***concentration of the gas (N<sub>2</sub>O), and then converted this into water-air emissions based on a gas transfer velocity (k). Gas***  
 241 ***transfer velocities can be highly variable, especially in estuaries where the importance (and magnitude) of factors like wind,***

242 *flow velocity, and water depth can all vary a lot over space and time. This complexity is reflected in the wide range of*  
 243 *empirical k value parameterisations that have been developed for estuaries (see e.g., Rosentreter et al. (2021), also Hall and*  
 244 *Ulseth (2019) for a good review of the topic, albeit for freshwater systems). However, here the authors convert measured*  
 245 *concentrations to emissions using a single parameterisation (L116-125). This creates considerable uncertainty, which is*  
 246 *not reflected in the reported estuary emissions estimates. Emissions should be recalculated using 3-5 k parameterisations,*  
 247 *and the variability of these outputs reported in the results / figures. More information should also be supplied on the wind*  
 248 *speed data used in the parameterisations. It is important to understand how the values measured during the campaigns*  
 249 *compare to ‘average’ conditions around the estuary when considering the upscaled seasonal emissions values (e.g., are*  
 250 *emissions estimates likely to be on the low side because cruises were only done on low-wind days?).*

251 As suggested, we included calculations based on three other parameterizations. We calculated and discussed the effect of wind  
 252 speeds in relation to average conditions along the estuary and added the information concerning average wind speed.

<b>Lines</b>	<b>Change</b>
L128-151	Calculated flux densities and emissions with four parametrizations and different wind speeds
L204-220	We included detailed results in the “Results” section for the new calculations of N <sub>2</sub> O flux densities and emissions
Table S2	We included flux-densities calculations using other parametrizations and wind speeds in the supplementary material
L262 and L414-434	We changed flux densities and N <sub>2</sub> O emission estimates in the revised manuscript so that they fit to the new calculations
L416-420	Addressed the uncertainties in the “Discussion” section

253

254 **2. Relationship between N<sub>2</sub>O and N inputs:** *As discussed in the paper intro here, aquatic N<sub>2</sub>O emissions are generally*  
 255 *predicted based on N loads to the system (i.e., leaching of N, inputs from WWTPs, etc). While here N<sub>2</sub>O emissions are*  
 256 *discussed and presented, the N inputs side of the equation is not clear to me. In the site description it says that annual N*  
 257 *load were ~80 Gg y<sup>-1</sup> (L67) – but does this mean the estuary receives this much N, or discharges this much N? And how*  
 258 *does this break down between sources (WWTPs v river discharge)? On L231 it says that N<sub>2</sub>O emissions were low relative*  
 259 *to other high N input estuaries. But how do N inputs into the Elbe stack up compare to these other estuaries? I particularly*  
 260 *wonder how the ‘point source’ N loads around the port might stack up with those in other urban estuaries where N<sub>2</sub>O*  
 261 *emissions have been measured, e.g., (Wells et al., 2018). Constraining the other side of the N<sub>2</sub>O emissions v N inputs*  
 262 *equations is critical for placing these findings into a more global context. Within the study, more information on N loads*  
 263 *will also be important for picking apart the seasonal emissions drivers. How much N enters the estuary at the port? Is this*  
 264 *input seasonally variable? Did it vary between the sampled years? Do these variations correspond with variations in*  
 265 *emissions (particularly the size of the winter N<sub>2</sub>O-excess excursion)?*

266 The N-loads of 80 Gg yr<sup>-1</sup> are calculated from concentrations data at the station “Seemanshoeft”, which is located at the  
 267 Hamburg Port (stream kilometer 628.9). In general, point sources play a subordinate role in the nitrogen input of the Elbe  
 268 estuary (Hofmann et al., 2005; IKSE, 2018) with dominating agricultural sources in the upper and middle Elbe River (Hofmann  
 269 et al., 2005; Johannsen et al., 2008). Further, we calculated annual varying DIN and total nitrogen (TN) loads for our  
 270 observation period and listed the results in the supplements as recent TN loads were lower varying between 43.1 kt-N yr<sup>-1</sup> and  
 271 70.2 kt-N yr<sup>-1</sup> from 2015 to 2021 (FGG, 2021).

272 We restructured our section 4.5 of the discussion: We now address the relation of N<sub>2</sub>O emissions and N inputs based on a  
 273 comparison of the amount of DIN released as N<sub>2</sub>O (for annual loads, seasonal loads and for each cruise separately). In the  
 274 revised version, we compare the relation of flux densities and N<sub>2</sub>O emission versus N input to a wider set of literature data and  
 275 across estuaries. We refer to the change of drivers of N<sub>2</sub>O emissions in winter (high riverine input and nitrification) versus  
 276 spring and summer (organic matter) more clearly. Finally, we highlight the link of N<sub>2</sub>O emissions to eutrophication to broaden  
 277 the scope of our study, which we also now address in the revised abstract, introduction and conclusion.

Lines	Change
L19-20	Rewritten abstract: “However, in spring and summer, N <sub>2</sub> O saturation and emission did not decrease alongside lower riverine nitrogen loads [...]”
L47-54	Adding new research aim to investigate the driving factors of N <sub>2</sub> O emissions along the estuary as well as looking into N <sub>2</sub> O and DIN relation.
L78-79	Added: “Point sources along the estuary provide only small part of the total nitrogen input to the Elbe Estuary (Hofmann et al., 2005; IKSE, 2018)”
L413-447	Rewritten and restructured section 4.5 focusing on N <sub>2</sub> O:DIN relation and comparing our results with other estuaries
L449-469	Rewritten conclusion
Table S4 and S5	Included annual and seasonal nitrogen loads for station Seemanshoeft

278

279 ***1. Introduction: It is not entirely clear how studying N<sub>2</sub>O in the Elbe estuary will advance understanding of aquatic N<sub>2</sub>O***  
 280 ***emissions / fill a needed research gap. A stronger transition between the penultimate and last paragraphs of the discussion***  
 281 ***is needed (how does the present study relate to the broader literature). Stating a testable hypothesis, rather than just site-***  
 282 ***specific study objectives, in the last paragraph may also help make the study more clearly relevant to the broader scientific***  
 283 ***community. Is this just a case study or will the data help us understand estuary N cycling and gaseous emissions in a more***  
 284 ***fundamental way?***

285 In the revised manuscript, we now elaborate a research question of interest for a wider audience by studying drivers for the  
 286 reported discrepancies in the N<sub>2</sub>O:DIN relation (Borges et al., 2015; Marzadri et al., 2017; Wells et al., 2018). Overall, the aim  
 287 of our research is to provide insight on drivers of N<sub>2</sub>O productions and emissions from heavily anthropogenic impacted  
 288 estuaries.

289

290

Lines	Change
L47-54	Elaborated new research question from interest for broad scientific community
L58-59	Added overall goal of our research

291

292 **2. Discussion:** *While I think overall the data interpretation makes sense, the discussion section currently reads as a bit*  
 293 *descriptive and could go further to place these findings in a broader context (rather than just the context of how we*  
 294 *understand the Elbe River Estuary). This could include in particular more discussion of N cycling in urban estuaries /*  
 295 *where there are point N pollution. Where else in the world would the observed seasonal patterns be expected to be found?*  
 296 *I also think there is missing some discussion of ‘alternative hypotheses’ – work through the logic of why denitrification is*  
 297 *not thought to be the primary driver of N<sub>2</sub>O in the estuary, and why benthic production (e.g., (Chen et al., 2022)) is also*  
 298 *ruled out. Also please carefully edit to ensure that you are not repeating results in this section.*

299 We now discuss alternative hypotheses in more detail as described below in the replies for the individual comments. We also  
 300 discuss the effects of benthic fluxes and production in more detail (see specific comment below). We have removed results  
 301 sections from the discussion section, and focused more on comparing our finding with research from other estuaries to address  
 302 a broader audience.

303 **3. Conclusion:** *This is currently very focused on untangling what exactly is happening within the Elbe River Estuary, but*  
 304 *the implications for broader understanding of aquatic N<sub>2</sub>O production and emissions are not clear.*

305 We modified our research question towards general controls of N<sub>2</sub>O production and emissions from estuaries with high  
 306 nitrogen loads. Consequently, we rewrote large parts of the discussion section 4.5. Thus, we also changed our conclusion and  
 307 abstract to highlight the new findings.

Lines	Change
L27-29	Rewritten abstract: “This effect of phytoplankton growth and the overarching control of organic matter on N <sub>2</sub> O production highlights that eutrophication and agricultural nutrient input can increase N <sub>2</sub> O emissions in estuaries.”
L47-54	Elaborated new research question from interest for broad scientific community
L58-59	Added overall goal of our research
L413-447	Rewritten and restructured section 4.5 focusing on N <sub>2</sub> O:DIN relation and comparing our results with other estuaries
L449-470	Rewritten conclusion

308

309 **L17-19: This sentence is not clear (how does N<sub>2</sub>O ‘compensate’ for decreasing N loads?), please reword.**

310 This statement was also unclear to reviewer 1. Please see comment above for the changes in the revised version of our  
311 manuscript.

312

313 **L22-24: “In winter, high riverine N<sub>2</sub>O concentrations led to high N<sub>2</sub>O emissions from the estuary, whereas in summer,**  
314 **estuarine biological N<sub>2</sub>O production led to equally high N<sub>2</sub>O emissions.” This is I think getting at a crucial point (that**  
315 **although seasonal magnitude of N<sub>2</sub>O fluxes did not differ the drivers of these fluxes did), the meaning is not clear. What**  
316 **is the difference between winter ‘high N<sub>2</sub>O concentrations’ and summer ‘high N<sub>2</sub>O production’? Reword to be more precise**  
317 **about these differences.**

318 We revised the manuscript to highlight the relevance of our research for a broader audience. Thus, we rewrote our abstract,  
319 focusing on a comparison of our results with other research in heavily managed estuaries, highlighting that we found seasonal  
320 varying drivers of N<sub>2</sub>O emissions that did not scale with DIN loads and were directly linked to eutrophication phenomena.  
321 Thus, we also re-wrote this phrase.

Lines	Change
L18-21	Changed to: “We found that the estuary was a year-round source of N <sub>2</sub> O, with highest emissions in winter when dissolved inorganic nitrogen (DIN) loads and wind speeds are high. However, in spring and summer, N <sub>2</sub> O saturations and emissions did not decrease alongside lower riverine nitrogen loads, suggesting that estuarine in-situ N <sub>2</sub> O production is an important source of N <sub>2</sub> O.”

322

323 **L70: How often is ‘on a regular basis’? e.g., weekly, yearly, every three years?**

Lines	Change
L82-85	Added clarification

324

325 **L86: Suggest changing ‘steaming upstream’ to ‘travelling upstream’ (steaming sounds a bit antiquated)**

Lines	Change
L95	Changed “steaming” to “travelling”

326

327 **L101-104: More information on number of nutrient samples collected per survey, as well as method detection limits and**  
328 **precision, would be useful.**

329 We will included the detection limits and also added a short description addressing the range of samples numbers.

Lines	Change
L105	Clarified numbers of samples taken
L113-114	Added detection limits

330

331 **L109: How often was ‘regularly’? e.g., before each cruise?**

Lines	Change
L119-120	Clarified: “Twice a day, we analyzed two standard gas mixtures of N <sub>2</sub> O in synthetic air (500.5 ppb ± 5 % and 321.2 ppb ± 3 %) to validate our measurements.”

332

333 **L116: How often, and how, was dry air sampled during each cruise?**

Lines	Change
L126-127	Clarified: “Atmospheric N <sub>2</sub> O dry mole fractions were measured before and after each transect cruises using an air duct from the deck of the research vessel.”

334

335 **L122: The term ‘flux densities’ is not one I’m familiar with – more common to see something like ‘water-air fluxes’ or**  
336 **‘evasion’.**

337 In physics, fluxes per unit area are called flux densities (Nitrous oxide mass flux | environmentdata.org, 2023), whereas the  
338 term “fluxes” only describe amount of N<sub>2</sub>O moving between the sea-air interface and is unitless. The terms are indeed often  
339 used synonymously.

340 We would like to stick to the term “flux densities”, which has been used previously by other researchers (e.g. Brase et al.,  
341 2017; Bange et al., 2019; Morgan et al., 2019; Forster et al., 2009).

342

343 **L123-125: Please provide some clarification on the upscaling approach used to calculate whole-estuary emissions. From**  
344 **the description it sounds like the mean flux was multiplied by the estuary surface area? Or were these calculations area-**  
345 **weighted, and if so at what resolution?**

346 We changed our way of estimate N<sub>2</sub>O emissions, which we elaborated in detail for a comment of reviewer 1 above.

Lines	Change
L142-151	We included a detailed description of the calculation in the “Method” section
L213-220	We included detailed results in the “Results” section
Table S2	We included flux-densities calculations using other parametrizations and wind speeds in the supplementary material
L262 and L414-434	We changed flux densities and N <sub>2</sub> O emission estimates in the revised manuscript so that they fit to the new calculations

347

348

349 **L127-128: Citation?**

Lines	Change
L154	Added a citation

350

351 **L148: Low relative to what?**

352 “low” is less than 1  $\mu\text{mol L}^{-1}$  – we will specify this.

Lines	Change
L177	Added: “(< 1 $\mu\text{mol L}^{-1}$ )”

353

354 **L163-189: Separating the N<sub>2</sub>O data into different sections for the different units (molar concentrations, % saturation,**  
355 **water-air fluxes) is confusing as these are all inter-related. For instances, it is hard to make sense of the meaning of the**  
356 **molar concentrations without also considering whether these reflect changes in percent saturation (i.e., changes due to**  
357 **water temperature / salinity v source / production). I suggest integrating these lines of data (and thinking) to provide a**  
358 **clearer picture of estuary N<sub>2</sub>O patterns.**

359 In Figure 2, we changed N<sub>2</sub>O concentrations to N<sub>2</sub>O saturations. Thus, we removed our previous Fig. 3 from the text. We also  
360 changed the results sections accordingly.

Lines	Change
L182-190	Replaced Fig. 2i, j with N <sub>2</sub> O saturations and changed figure caption
L191-203	Changed section 3.2 to “Atmospheric N <sub>2</sub> O and N <sub>2</sub> O saturation”
L204-220	Changed section 3.3 to “N <sub>2</sub> O flux densities and N <sub>2</sub> O emissions”
	Deleted the description of N <sub>2</sub> O concentrations
	Removed previous Fig. 3 from the text

361

362 **L204: High relative to what?**

Lines	Change
L225	Specified: “(> 100 %)”

363

364

365 **L209-218: The AOU v N<sub>2</sub>O-excess relationship really highlights the importance, and seasonality, of the port for estuary**  
 366 **N<sub>2</sub>O emissions, with distinct peaks in the winter and consumption in the summer. Given that this underpins the discussion**  
 367 **around seasonal N<sub>2</sub>O source switching, I wonder if there is a way to include more than just these ‘representative’ plots in**  
 368 **the main text. For instance, a table with info on AOU v N<sub>2</sub>O-excess slopes, and min-max range for the port? I think if the**  
 369 **port data is excluded something like an ANCOVA could be used to compare shifts in slope relationships.**

370 In the Port region, N<sub>2</sub>O<sub>xs</sub> and AOU had no linear relation during most of our cruises (e.g. June 2015 and August 2017).  
 371 Therefore, a table with slopes and min-max ranges would miss crucial information. However, we will include figure S2 from  
 372 the supplement in the main text so that we do not only show representative plots but all cruises.

Lines	Change
L236-239	Included all AOU vs N <sub>2</sub> O <sub>xs</sub> plots in Fig. 3 and changed figure caption

373

374 **L256-260: Interesting relationship between NO<sub>2</sub>- and N<sub>2</sub>O. This could be connected to previous work, e.g., (Sharma et al.,**  
 375 **2022; Smith and Bohlke, 2019; Wertz et al., 2018)**

Lines	Change
L291-296	Added: “This co-occurrence of nitrite accumulation and increased N <sub>2</sub> O saturation has been interpreted as signs for N <sub>2</sub> O production via denitrification (e.g. Wertz et al., 2018; Sharma et al., 2022). However, denitrification does not seem likely in this oxic water column. Such a succession of nitrite and ammonium peaks is also typical for remineralization and nitrification, and the slight decrease of oxygen concentrations around the higher N <sub>2</sub> O saturation (Fig. 2g and i) suggests oxygen consumption, possibly caused by these two processes.”

376

377 **L314-316: This should be in the results section**

Lines	Change
L240-259	Added section 3.4 describing results from our statistical analysis and removed this paragraph

378

379 **L318-324: Interesting! I wonder if the algae themselves could also be contributing to the N<sub>2</sub>O production, e.g., (Fabisik et**  
 380 **al., 2023)**

Lines	Change
L372-373	Added: “Fabisik et al. (2023) showed that algae could additionally contribute to N <sub>2</sub> O production.

381

382

383 **L330-332: This makes sense, but is this the only possible explanation for high emissions around the port area? What about**  
 384 **wastewater inputs, enhanced benthic production, and/or enhanced groundwater connectivity due to dredging? Some**  
 385 **discussion of these points will make this conclusion stronger.**

386 We restructured our discussion focusing more on elaborating potential causes for the high N<sub>2</sub>O peak. We focused on (1) point  
 387 sources – mainly the wastewater treatment plant, (2) deepening and dredging operations in the Port of Hamburg and (3) in-situ  
 388 production, discussing both production in the water column and sediment. Therefore, we moved parts of section 4.4 to 4.3

<b>Lines</b>	<b>Change</b>
L330-356	Restructured the paragraph
L330-356 and L383-410	Moved parts from 4.4 to 4.3 and thus, adapted section 4.4

389

390 **L357-358: How extreme was this rain event, i.e., was it more extreme than any rainfalls over the other five years of**  
 391 **sampling? This will help verify the attribution, and also put the pulse into context. It would then be instructive to recalculate**  
 392 **the seasonal budget with and without this pulse.**

393 Considering the statistical recurrence probability, it is likely that similar events occurred during the last five years, but so far  
 394 not during a comparable sampling cruise. Moreover, we assume that temperature was equally important as water mass, because  
 395 cold rain water in the waste water treatment plant led to aggravated operation conditions. Thus, in warmer months the effect  
 396 might be different. We discuss this in more detail in the revised manuscript.

<b>Lines</b>	<b>Change</b>
L213-220	Calculated N <sub>2</sub> O emissions estimates with and without the pulse and reported the variability
L403-404	Added statistical recurrence probability
L408-410	Added: “An important factor for aggravated conditions was a temperature drop in the WWTP caused by cold rain water, we hypothesize that a similar rain event in warmer months would not lead to comparable N <sub>2</sub> O peaks”

397

398 **L392: If large riverine loads were the main driver, wouldn't there be a continuous decrease in concentration over distance?**  
 399 **But instead emissions peak in the port.**

400 We restructured the section 4.5 and thus removed this statement from the text.

<b>Lines</b>	<b>Change</b>
L413-447	Restructured section 4.5

401

402

403 **Table 2: Standard deviations for the air N<sub>2</sub>O concentrations would be helpful**

Lines	Change
L210-212	Added standard deviations to the Tab. 2

404

405 **Fig. 1: The most important pieces of info in this map (where sampling points are, where the port is, where the MTZ is)**  
406 **don't really stand out. Can you adjust colours, font size, etc to better highlight these key features? A scale bar for the main**  
407 **map would also be helpful.**

408 We changed the style of map to highlight key features and important locations as suggested by the reviewer.

Lines	Change
L86-88	Changed Fig. 1 and figure caption

409

410 **Fig. 2: I'm not sure that there is much value in showing N<sub>2</sub>O concentrations (in nM) here – the % saturation information**  
411 **in the subsequent figure is much more effective for showing fluctuations between seasons and over the salinity gradient,**  
412 **given the relatively low concentrations and the impact of both temperature and salinity on N<sub>2</sub>O solubility. It would also be**  
413 **helpful to have ‘summer’ and ‘winter’ headings at the top of the two columns to make the point of difference more**  
414 **immediately obvious.**

Lines	Change
L182	Included a heading to the plot
Fig. S1	Included a heading to the plot
L182-190	Changed Fig. 2i,j from N <sub>2</sub> O concentrations to saturations and changed figure caption
	Removed previous Fig. 3 from the text

415

416 **Fig. 3: A unified y axis scale would be helpful for picking out seasonal differences**

Lines	Change
	Removed previous Fig. 3 from the text

417

418 **Fig. 4: As above, unified axes scales would make differences between sampling dates much clearer.**

419 We decided against unified y-axis for all cruises. The March 2021 cruise differs so much that it is hard to see variabilities in  
420 the other cruises if we use the same y-axis. However, we used unified x-axes and y-axes scales for all other cruises.

Lines	Change
L236	Included all cruises in Fig. 3 with unified x-axes and unified y-axes except for March 2021
L237-239	Changed figure caption

421

422

423 **Fig. 5: Different y axes are needed for the different variables (N<sub>2</sub>O, O<sub>2</sub>, TN), if not different plot panels**

Lines	Change
Fig. S2	Moved this figure to the supplementary material
L275-278	Added instead of the figure: “However, since the BIOGEST study in 1997 (Barnes and Upstill-Goddard, 2011), N <sub>2</sub> O remained relatively stable at ~ 200 % saturation despite a concurrent decrease in TN concentration from ~400 μmol L <sup>-1</sup> to around 200 μmol L <sup>-1</sup> (Fig. S2, Hanke and Knauth, 1990; Barnes and Upstill-Goddard, 2011; Brase et al., 2017; FGG, 2021)”
Fig. S2	Included a plot panel for each variable and changed figure caption

424

425 **Fig. 6: I found this to be too many variables on the same plot to make much logical sense out of. I suggest separating into**  
 426 **two panels, one for all of the N species (y axis unit is μM N), and then another with two y axes, one for PN and one for**  
 427 **C/N.**

428 We adapted the plot as suggested by the reviewer.

Lines	Change
L298-304	Changed figure and figure caption
Fig. S3-S13	Changed figure and figure caption

429

430 **Fig. 7: It would be helpful to use a different pattern or colour scheme to distinguish the winter v summer cruises.**

431 We restructured and rewrote this part of the discussion and removed Fig. 7. The new Fig. 6 shows seasonal variations in N<sub>2</sub>O  
 432 saturation, nitrous oxide emissions, DIN loads and N<sub>2</sub>O:DIN ratios. We considered the comment of the reviewer regarding the  
 433 color scheme for the new figure.

Lines	Change
	Removed previous Fig. 7
L442-447	Added new Figure

434

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