



1 **Global database of actual nitrogen loss rates in coastal and marine sediments**

2 Yongkai Chang¹, Ehui Tan^{1*}, Dengzhou Gao², Cheng Liu³, Zongxiao Zhang⁴,

3 Zhixiong Huang¹, Jianan Liu¹, Yu Han¹, Zifu Xu¹, Bin Chen⁵, Shuh-Ji Kao^{1*}

4 ¹ State Key Laboratory of Marine Resource Utilization in South China Sea, School of
5 Marine Science and Engineering, Hainan University, Haikou, China

6 ² Key Laboratory for Humid Subtropical Eco-Geographical Processes of the Ministry
7 of Education, School of Geographical Sciences, Fujian Normal University, Fuzhou,
8 China

9 ³ Shandong Key Laboratory of Eco-Environmental Science for the Yellow River Delta,
10 Shandong University of Aeronautics, Binzhou, China

11 ⁴ School of Environmental Science and Engineering, Southern University of Science
12 and Technology, Shenzhen, Guangdong, China

13 ⁵ State Key Laboratory of Marine Environmental Science, College of Ocean and Earth
14 Sciences, Xiamen University, Xiamen, China

15 ***Corresponding author:**

16 Ehui Tan (ehuitan@hainanu.edu.cn) and Shuh-Ji Kao (sjkao@hainanu.edu.cn)

17



18 **Abstract**

19 Denitrification and anaerobic ammonium oxidation (anammox) convert reactive
20 nitrogen to inert N_2 , and play vital roles in nitrogen removal in coastal and marine
21 ecosystems, weakening the adverse effects caused by terrestrial excessive nitrogen
22 inputs. Given the importance of denitrification and anammox in nitrogen cycle, lots of
23 studies have measured denitrification and anammox through intact core incubations
24 across different systems, and nitrogen loss processes are affected by a series of
25 environmental factors such as organic carbon, nitrate, dissolved oxygen and
26 temperature. However, a global synthesis of actual nitrogen loss rates is lacking and
27 how environmental factors regulate nitrogen loss remains unclear. Therefore, we have
28 compiled a database of nitrogen loss rates, including denitrification and anammox in
29 coastal and marine systems from published literatures. This database includes 473,
30 466, and 255 measurements for total nitrogen loss denitrification and anammox,
31 respectively. This work deepens our understanding of the spatial and temporal
32 distribution of denitrification, anammox and the relative contribution of anammox to
33 total nitrogen loss and their corresponding environmental controls. To our knowledge,
34 the constructed database for the first time offers a comprehensive overview of actual
35 nitrogen loss rates in coastal and marine ecosystems on a global scale. This database
36 can be utilized to compare nitrogen loss rates of different regions, identify the key
37 factors regulating these rates, and parameterize biogeochemical models in the future.
38 This database is available in Figshare repository :
39 <https://doi.org/10.6084/m9.figshare.27745770.v3> (Chang et al., 2024).



40 **KEYWORDS:** nitrogen cycle, denitrification, anammox, coastal and marine

41 ecosystems, isotope pairing technology, intact core incubations

42



43 **1 Introduction**

44 The production of anthropogenic reactive nitrogen has intensified remarkably since
45 the mid-20th century to meet the increasing global population (Kennedy, 2021). It is
46 estimated that nitrogen is entering Earth's ecosystems at more than twice its natural
47 rate, drastically disrupting the pristine nitrogen cycle (Canfield et al., 2010). Much of
48 the excess nitrogen, primarily in the form of nitrate, is conveyed downriver to coastal
49 and marine systems due to the low use efficiency of crops (Cui et al., 2013), resulting
50 in a series of environmental issues including harmful algal blooms, eutrophication,
51 and hypoxia (Dai et al., 2023). Consequently, it is critical to understand the
52 transformations, particularly the fates of reactive nitrogen, encountering the fact that
53 the nitrogen cycle has been intensively altered and is currently functioning beyond the
54 safe operating space for humanity (Richardson et al., 2023).

55 Denitrification and anammox are two key nitrogen loss processes in aquatic
56 environments, playing important roles in mitigating the adverse effects of excessive
57 nitrogen inputs (Chen et al., 2021; Tan et al., 2022). Denitrification is the sequential
58 reduction of nitrate, nitrite, nitric oxide, and nitrous oxide (N₂O) to dinitrogen gas
59 (N₂), which is the most energetically favorable respiratory pathway in the absence of
60 oxygen (Devol, 2015), serving as the predominant mechanism for nitrogen loss in
61 coastal ecosystems (Damashek & Francis, 2018; Deng et al., 2024). Anaerobic
62 ammonium oxidation (Anammox), an alternate nitrogen loss pathway, utilizes nitrite
63 and ammonium to generate N₂ with no greenhouse gas N₂O production under
64 anaerobic conditions (Graaf et al., 1995), and is a chemoautotrophic process with no



65 direct demand for organic carbon (Strous et al., 1999). Therefore, anammox is an
66 environment-friendly and energy-saving process compared to denitrification.

67 The ^{15}N isotope pairing technique (IPT) has been applied to a variety of sediments to
68 quantify nitrogen loss rates in these settings (Nielsen, 1992; Robertson et al., 2019).

69 Slurry incubation and intact core incubations in combination with IPT are two widely
70 used methods for studying benthic nitrogen transformation pathways (Song et al.,

71 2016b). Slurry incubations have been used to estimate the potential rates due to
72 advantage of simple operation in incubations (Thamdrup & Dalsgaard, 2002), and a

73 large number of studies have used this method to study sediment nitrogen loss.
74 However, slurry incubations could not reflect the genuine benthic nitrogen

75 transformation rates, as the natural gradients of substrates and redox in sediments
76 were disrupted during incubations (Trimmer et al., 2006). The application of intact

77 core incubations can overcome this drawback and will enable us to fully clarify and
78 understand the nitrogen cycle in field aquatic ecosystems.

79 Over the past thirty years, the introduction of isotope pairing technology has enabled
80 numerous studies to measure anammox and denitrification using intact core

81 incubations across a range of coastal and marine environments. These environments
82 include intertidal wetlands (Adame et al., 2019; Liu et al., 2020), estuaries and coasts

83 (Chen et al., 2021; Cheung et al., 2024; Deek et al., 2013; Hellemann et al., 2017),
84 lagoons (Bernard et al., 2015; Magri et al., 2020) and oceans (Deutsch et al., 2010; Na

85 et al., 2018). Despite decades of research, the majority of studies on denitrification
86 and anammox have been limited to local or regional scales. Various environmental



87 factors, such as the availability of organic carbon (Yin et al., 2015) and nitrate
88 (Asmala et al., 2017), dissolved oxygen (Bonaglia et al., 2013; Song et al., 2021), and
89 temperature (Tan et al., 2022) influence these processes in coastal marine ecosystems.
90 However, to date, the global patterns and drivers of sediment nitrogen loss rates
91 remain poorly understood in coastal and marine systems.

92 In view of the critical role of nitrogen removal processes and the current lack of a
93 comprehensive database on actual nitrogen loss in coastal and marine systems, we
94 have integrated actual nitrogen loss rates, including denitrification and anammox,
95 from published studies, and constructed a dataset on nitrogen removal rates in these
96 systems. This study provides a global-scale overview of the biogeography and
97 potential controlling factors of denitrification and anammox in coastal and marine
98 ecosystems. It also highlights the potential applications of this database such as using
99 machine learning to predict the distribution of denitrification and anammox and
100 offering a crucial dataset for the parameterization and development of biogeochemical
101 models.

102 **2 Methods**

103 **2.1 Data compilation**

104 Nitrogen loss rates, including denitrification and anammox measured through intact
105 core incubations in coastal and marine ecosystems, were extracted from the literature
106 published between 1996 and 2024. Table 1 summarized the locations, observation
107 numbers, core incubation methods and references of nitrogen loss rates measurements.



108 The intact core incubations in this study include both traditional core incubations
109 (Bonaglia et al., 2017; Cheung et al., 2024) and continuous flow experiments
110 combined with core incubations (Liu et al., 2020; McTigue et al., 2016). The
111 peer-reviewed articles compiled in this study were sourced from the Web of Science
112 database as of June 2024. The search terms were “denitrification” or “anammox” or
113 “nitrogen loss” or “nitrogen removal”. Only data where denitrification and/or
114 anammox rates were measured using intact core incubations combined with ^{15}N
115 isotope pairing techniques were included, while those measured via slurry incubation
116 were excluded. The intact core incubation experiments were primarily conducted in
117 dark conditions and near-*in situ* or *in situ* ambient temperatures. In cases where
118 nitrogen loss rates were measured under both light and dark conditions, only those
119 measured in the dark were included. Measurements under light conditions have been
120 detailed in studies reported by Bartoli et al. (2021), Chen et al. (2021),
121 Risgaard-Petersen et al. (2004), Rysgaard et al. (1996b), and Welsh et al. (2000).
122 Some studies have investigated the changes in nitrogen loss processes under varying
123 oxygen concentrations (Bonaglia et al., 2013; Neubacher et al., 2011; Song et al.,
124 2021), however, only nitrogen loss rates measured under ambient oxygen
125 concentrations were extracted for this database. Additionally, studies examining the
126 effects of meiofauna or antibiotics on nitrogen removal (Bonaglia et al., 2014b; Wan
127 et al., 2023) were not included, only rates measured without meiofauna or antibiotic
128 additions were considered. At least one environmental variable was recorded for each
129 selected study, and means and sample sizes had to be reported for nitrogen removal



130 rates. Articles that only reported nitrogen loss rates without any environmental
131 variables were excluded. Data on total nitrogen loss rates (the sum of denitrification
132 and anammox), denitrification rates, anammox rates, and related environmental
133 variables were collected from tables, text, and/or supplementary materials, and in
134 some cases, extracted from graphs using Origin 2020 software. The unit conversions
135 were performed where necessary. In addition, longitude and latitude were extracted
136 from figures from published articles if not shown in the main text.

137

138 The database includes observation details (year of sampling, month of sampling,
139 latitude, and longitude), sediment parameters, and water physicochemical factors,
140 such as sediment organic carbon, the ratios of carbon to nitrogen (C/N ratios), oxygen
141 penetration depth, and water salinity, depth, temperature, DO, ammonium and nitrate
142 concentrations. Note that some environmental variables were not reported in the
143 original studies. NM represents parameters that were not measured, and empty or NA
144 indicates data not available or reported. In total, the database comprises 473, 466, 255,
145 and 255 measurements of total nitrogen loss rates, denitrification rates, anammox
146 rates, and the relative contribution of anammox to total nitrogen loss, respectively.
147 Authors and interested readers are welcomed to contact us to indicate an error or
148 update the data in the database.

149 For quality control, extreme nitrogen loss rate values were excluded from the database
150 following Chauvenet's criterion (Glover et al., 2011), a method typically applied to
151 normally distributed data to identify outliers whose deviation from the mean has a



152 probability lower than $1/(2n)$. More details about Chauvenet's criterion can be found
153 in Glover et al., (2011) and Buitenhuis et al. (2013). Very high rates of denitrification
154 were observed in the Tama Estuary, Japan (Usui et al., 2001), a constructed wetland in
155 Casino, NSW, Australia (Erler et al., 2008), a coastal lagoon in Sacca di Goro lagoon,
156 Italy (Magri et al., 2020) and the Tropical Coastal Wetlands, Australia (Adame et al.,
157 2019). For anammox, high rates were found only in a constructed wetland in Casino,
158 NSW, Australia (Erler et al., 2008). Similarly, high values for anammox's contribution
159 to total nitrogen loss were observed in the Changjiang River Estuary, China (Liu et al.,
160 2020), the Norwegian Trench, Skagerrak (Trimmer et al., 2013), and the Great Barrier
161 Reef lagoon (Erler et al., 2013), with contributions exceeding 70%. Observations with
162 nitrogen loss rates of 0 or NA were excluded from the outlier analysis. For example,
163 anammox rates of 0 were reported in the Changjiang River Estuary, China (Liu et al.,
164 2020), the North Sea (Neubacher et al., 2011; Rosales Villa et al., 2019), the Pearl
165 River Estuary, China (Tan et al., 2019), the Norwegian Trench, Skagerrak (Trimmer et
166 al., 2013), and the Gulf of Finland, Baltic Sea (Jäntti et al., 2011). After excluding
167 observations of 0 and NA (0, 8, 252, and 253 observations for total nitrogen loss rates,
168 denitrification rates, anammox rates, and anammox's contribution to total nitrogen
169 loss), the nitrogen loss rates were natural-log transformed for further analysis.

170 **2.2 Methods for measuring denitrification and anammox rates**

171 Before the discovery of anammox, denitrification was regarded as the sole significant
172 pathway responsible for nitrogen loss (Dalsgaard & Thamdrup, 2002). The ^{15}N



173 isotope pairing technique (IPT) was developed to quantify denitrification rates
174 (Nielsen, 1992). In this method, the overlying water of intact sediment cores is
175 enriched with $^{15}\text{NO}_3^-$, which is mixed with the naturally occurring $^{14}\text{NO}_3^-$. After a few
176 hours of incubation, the denitrification products, ^{15}N -labeled dinitrogen gas ($^{29}\text{N}_2$ and
177 $^{30}\text{N}_2$), are measured. Incubations to measure nitrogen loss rates have been mostly
178 conducted in dark conditions and near-*in situ* or *in situ* ambient temperatures. After
179 incubating for 1 h to over 96 h, the incubation is halted by injecting saturated HgCl_2 or
180 ZnCl_2 saturation solution or 37% formaldehyde. The samples are then preserved for
181 $^{15}\text{N}_2$ gas analyses through isotope ratio mass spectrometer (IRMS) or membrane inlet
182 mass spectrometry (MIMS). Key experimental details, such as incubation conditions,
183 temperature control, incubation time, termination, and calculation references, are
184 compiled in the database if provided in the original studies. For more detailed
185 experimental information, refer to the corresponding references.

186 The production rate of unlabeled $^{14}\text{NO}_3^-$ (IPT p_{14} , also referred to as the genuine
187 production of N_2) can be calculated based on the assumption of random isotope
188 pairing during the denitrification of the uniformly mixed NO_3^- species. The following
189 equation is commonly used to estimate the genuine N_2 production (Nielsen, 1992;
190 Steingruber et al., 2001).

$$191 \quad \text{IPT}p_{14} = \frac{p^{29}\text{N}_2}{2 \times p^{30}\text{N}_2} \times (p^{29}\text{N}_2 + 2 \times p^{30}\text{N}_2) \quad (1)$$

192 Where $p^{29}\text{N}_2$ and $p^{30}\text{N}_2$ represent the total production rates of $^{29}\text{N}_2$ and $^{30}\text{N}_2$,
193 respectively.

194 Thamdrup and Dalsgaard (2002) were the first to quantify anammox through



195 anaerobic slurry incubations in natural environments, discovering that anammox
196 could account for more than 60% of total N₂ production. This highlighted the
197 significant role of anammox in nitrogen removal. Following this, Risgaard-Petersen et
198 al. (2003) proposed a modification to the traditional IPT, allowing for more accurate
199 quantification of true N₂ production in environments where anammox and
200 denitrification coexist. This revision also enables the distinction between N₂ produced
201 by anammox and denitrification. The revised IPT (rIPT) follows the same procedure
202 as the classical IPT, with ¹⁵NO₃⁻ added to the overlying water of intact sediment cores,
203 though the calculation process is more complex. The following equations are
204 commonly used to estimate the actual N₂ production (rIPT_{p14}) and denitrification
205 (*p14DEN*) as well as anammox (*p14ANA*) (Risgaard-Petersen et al., 2003; Trimmer
206 & Nicholls, 2009; Trimmer et al., 2006). The total N₂ production rate is the sum of
207 denitrification and anammox rates.

$$208 \quad \text{rIPT}_{p14} = 2r_{14} \times (p^{29}\text{N}_2 + p^{30}\text{N}_2 \times (1 - r_{14})) \quad (2)$$

$$209 \quad p14\text{DEN} = 2r_{14} \times (r_{14} + 1) \times p^{30}\text{N}_2 \quad (3)$$

$$210 \quad p14\text{ANA} = 2r_{14} \times (p^{29}\text{N}_2 - 2 \times r_{14} \times p^{30}\text{N}_2) \quad (4)$$

211 In these equations, $p^{29}\text{N}_2$ and $p^{30}\text{N}_2$ are the total production rates of ²⁹N₂ and ³⁰N₂,
212 respectively, and r_{14} represents the ratio of ¹⁴NO₃⁻ and ¹⁵NO₃⁻ in the nitrate reduction
213 zone. There are 3 different methods to estimate r_{14} , with detailed explanations
214 available in Trimmer et al. (2006).

215 Subsequently, Hsu and Kao (2013) revised the rIPT method to incorporate both N₂O
216 production and anammox, enabling the determination of the absolute rate of each



217 nitrogen loss pathway, including denitrification, anammox, and N₂O production from
218 denitrification. Denitrification and anammox measurements based on the method of
219 Hsu and Kao (2013) are included in this database, whereas data on the true N₂O
220 production rate has not been included.

221 **3 Results and discussion**

222 **3.1 Overview of the database**

223 Overall, there are 473, 466, and 255 measurements for total nitrogen loss
224 denitrification and anammox, respectively (Fig. 1). Denitrification and anammox have
225 been measured simultaneously at 255 observations. The observations of nitrogen loss
226 rates are primarily distributed in the Eastern coast of the United States, the Baltic Sea,
227 the Eastern Coast of China, the Eastern Coast of Australia, and polar regions of the
228 Northern Hemisphere (Fig. 1a). Before 2000, nitrogen loss measurements were
229 predominantly focused on denitrification, while both denitrification and anammox
230 rates have been measured concurrently since 2000 (Fig. 1b). Notably, more
231 observations were recorded in 2011 and 2017. The studies in 2011 were mainly
232 conducted in the Changjiang estuary and its adjacent East China Sea (Song et al.,
233 2021), the Jinpu Bay, China (Yin et al., 2015), the North Sea (Bale et al., 2014), the
234 Northern Baltic Proper (Bonaglia et al., 2014a) and the hypoxic zone off the
235 Changjiang River estuary, China (Yang et al., 2022). In 2017, high observations were
236 found in the Northern East China Sea, China (Chang et al., 2021), the Changjiang
237 River Estuary, China (Liu et al., 2020; Liu et al., 2019; Tan et al., 2022), the Coast of



238 Victoria, Australia (Kessler et al., 2018) and the Jiulong River Estuary, China (Tan et
239 al., 2022).

240

241 **3.2 Distribution of denitrification**

242 In total, the vast majority of nitrogen loss rate measurements were conducted in the
243 Northern Hemisphere, and data in the Southern Hemisphere were limited (Fig. 2a, 2b,
244 2c). The low and middle latitudes of the Northern Hemisphere have a large body of
245 observations, especially in the 20-30°N, 30-40°N, and 50-60°N latitude bands.
246 Denitrification rates ranged from 0.04 to 750 $\mu\text{mol N m}^{-2} \text{ h}^{-1}$, with a median value of
247 $7.72 \pm 4.30 \mu\text{mol N m}^{-2} \text{ h}^{-1}$. There is a decreasing trend in the denitrification rates with
248 latitude in the Northern Hemisphere, though the observations in the high latitude are
249 still limited. The measurements of denitrification were mostly conducted in later
250 spring, summer, and early autumn, from April to September (Fig. 2d, 2e, 2f). On a
251 global scale, no clear seasonal pattern for denitrification rates was observed.

252

253 **3.3 Distribution of anammox**

254 From a latitude perspective, the distribution of anammox rates closely mirrored that of
255 denitrification, with the majority of observations concentrated in the 20-30°N,
256 30-40°N, and 50-60°N latitude bands (Fig. 3a, 3b, 3c). However, compared to
257 denitrification, there were fewer anammox observations. Anammox rates spanned



258 from 0.01 to 48.94 $\mu\text{mol N m}^{-2} \text{ h}^{-1}$, with a median value of $1.00 \pm 0.39 \mu\text{mol N m}^{-2} \text{ h}^{-1}$.
259 Similar to denitrification, anammox rates also showed a decreasing trend with
260 increasing latitude in the Northern Hemisphere. Numerous anammox measurements
261 were conducted between April and September, consistent with the timing of
262 denitrification measurements (Fig. 3d, 3e, 3f). Additionally, February saw a high
263 number of anammox observations, and these observations were predominantly
264 conducted at the north East China Sea (Chang et al., 2021), the Changjiang Estuary
265 (Liu et al., 2019) and the Northeastern New Zealand continental shelf regions
266 (Cheung et al., 2024). On a global scale, there was no clear seasonal pattern for
267 anammox rates.

268

269 **3.4 Distribution of contribution of anammox to total N₂** 270 **production**

271 The relative importance of anammox to total N₂ production increased first and then
272 decreased, peaking in the 40-50°N latitudinal band in the Northern Hemisphere,
273 although data points in this band were limited (Fig. 4). The contribution of anammox
274 to total N₂ production varied from 0.22% to 67.33%, with a median value of 12.29%.
275 The highest value (67.33%) was recorded at a site on the North Atlantic continental
276 slope at a depth of 2000 m (Trimmer & Nicholls, 2009), where anammox accounted
277 for the majority of nitrogen removal. There were no significant monthly changes in
278 the relative importance of anammox to total nitrogen loss, except for March, when



279 anammox contributed a notably high percentage. High values in March were observed
280 in the Ulleung Basin, East Sea, and the continental shelf and slope, North Atlantic (Na
281 et al., 2018; Trimmer & Nicholls, 2009) where the stations were characterized by low
282 nitrate levels or deep water. These environmental conditions may inhibit
283 denitrification, thereby increasing the relative contribution of anammox to nitrogen
284 loss.

285

286 **3.5 Control factors on nitrogen loss rates**

287 The variations in denitrification rates, anammox rates, and the contribution of
288 anammox to total N_2 production (%) were compared against several environmental
289 variables, including sediment organic carbon, the ratios of carbon to nitrogen (C/N
290 ratios) and oxygen penetration depth, and water depth, temperature, salinity, dissolved
291 oxygen, ammonium, and nitrate concentrations. This comparison was conducted to
292 evaluate the controlling factors of nitrogen loss rates and the relative importance of
293 anammox to total nitrogen removal.

294 There was no significant relationship between denitrification rates and the contents of
295 sediment organic carbon ($p > 0.05$; Fig. 5a). Heterotrophic denitrification is primarily
296 carried out by facultative anaerobic heterotrophs (Devol, 2015), which use organic
297 carbon as an electron donor and energy source. Therefore, higher organic carbon
298 levels might be expected to promote denitrification (Damashek & Francis, 2018).
299 However, no such relationship was observed in this dataset. Denitrification rates



300 increased with sediment carbon nitrogen ratios ($r=0.23$, $p<0.01$; Fig. 5b). The C/N
301 ratios can indicate the reactivity of sediment organic material, with lower C/N values
302 generally representing more reactive organic matter (Cheung et al., 2024; Erler et al.,
303 2013). Typically, high denitrification rates are associated with sediments that have
304 lower C/N ratios. However, in this analysis, the opposite trend was observed. One
305 possible explanation is that microbial communities may adapt to use organic matter
306 typically encountered, though the organic matter is not labile (Salk et al., 2017).
307 Denitrification rates showed a weak negative correlation with oxygen penetration
308 depth ($r=-0.29$, $p<0.01$; Fig. 5c), as greater O₂ penetration may be adverse to the
309 occurrence of denitrification (Cheung et al., 2024). Denitrification rates also
310 decreased with water depth ($r=-0.2$, $p<0.01$; Fig. 5d), with most observations
311 occurring at depths shallower than 250 m. Denitrification was positively correlated
312 with higher water temperatures ($r=0.38$, $p<0.01$; Fig. 5e), and negatively correlated
313 with salinity ($r=-0.15$, $p<0.01$; Fig. 5f), with most rates falling within two salinity
314 ranges (0-10 and 30-40). Samples that had a salinity greater than 40 were collected in
315 hypersaline lagoons of tropical regions (Enrich-Prast et al., 2016). The relationship
316 between denitrification and salinity across coastal environments has been summarized
317 by Torregrosa-Crespo et al. (2023) and will not be further elaborated here. There was
318 a weak negative relationship between denitrification rates and dissolved oxygen
319 concentrations ($r=-0.23$, $p<0.01$; Fig. 5g). Overall, higher denitrification rates were
320 recorded in areas with high nitrate concentrations ($r=0.16$, $p<0.01$; Fig. 5h),
321 suggesting the importance of nitrate substrate in regulating denitrification, though



322 some high rates were also observed in sites with low nitrate levels. No significant
323 correlation was found between denitrification rates and ammonium concentrations
324 ($p>0.05$; Fig. 5i).

325

326 Anammox rates showed a weak positive correlation with sediment organic carbon
327 ($r=0.16$, $p<0.05$; Fig. 6a). Although anammox is an autotrophic process that does not
328 require organic carbon as an electron donor (Salk et al., 2017), some studies have
329 reported links between sediment organic carbon content and anammox rates. For
330 example, studies in subtropical mangrove sediments (Meyer et al., 2005) and the
331 Thames estuary (Trimmer et al., 2003) found that higher organic matter stimulated
332 anammox. This correlation may be due to enhanced mineralization leading to
333 increased ammonium production, which indirectly stimulates anammox (Damashek &
334 Francis, 2018), as sediment organic carbon can serve as a proxy for organic carbon
335 mineralization (Song et al., 2016a). Similar to denitrification, high anammox rates
336 were observed at sites with elevated C/N ratios ($r=0.33$, $p<0.01$; Fig. 6b). More
337 research is needed to reveal the influencing mechanisms of organic matter quantity
338 and quality on anammox. No clear trend was found between anammox rates and
339 oxygen penetration depth ($p>0.05$; Fig. 6c), and high anammox rates were observed in
340 shallow waters ($p>0.05$; Fig. 6d). Anammox rates showed a weak positive correlation
341 with temperature ($r=0.19$, $p<0.01$; Fig. 6e). While several studies have suggested that
342 low temperatures could favor anammox (Dalsgaard & Thamdrup, 2002; Rysgaard et
343 al., 2004; Tan et al., 2020), these studies primarily measured anammox potential using



344 anaerobic slurry incubations. Contrary to previous findings, our study showed that
345 actual anammox rates increased with rising temperatures, suggesting a discrepancy
346 between the effects of temperature on actual and potential anammox rates. Future
347 research is needed to investigate the underlying mechanisms for these inconsistent
348 results. Anammox rates decreased with increasing salinity ($r=-0.38$, $p<0.01$; Fig. 6f),
349 and showed no significant relationship with dissolved oxygen ($p>0.05$; Fig. 6g). A
350 weak positive correlation was observed between anammox rates and nitrate
351 concentration ($r=0.41$, $p<0.01$; Fig. 6h), highlighting the importance of substrates in
352 regulating anammox. Although anammox uses nitrite as an electron acceptor rather
353 than nitrate (Graaf et al., 1995), nitrate reduction can produce nitrite, which promotes
354 anammox activity. No relationship was found between anammox rates and
355 ammonium concentration ($p>0.05$; Fig. 6i).

356

357 Numerous studies have found that denitrification was linked to anammox in different
358 habitats, including estuary sediments (Liu et al., 2020), coastal wetland sediments
359 (Gao et al., 2017) and paddy soils (Shan et al., 2016). Consistent with their findings,
360 this work also found denitrification was positively correlated to anammox ($r=0.67$,
361 $p<0.01$; Fig. 7). A majority of denitrifying bacteria are heterotrophic and capable of
362 utilizing organic matter, and the decomposition of organic matter is accompanied by
363 the production of ammonium (Devol, 2015), supplying substrates for anammox. Thus,
364 the relationship between denitrification and anammox may suggest the tight coupling
365 of these two nitrogen removal pathways.



366

367 There was a positive correlation between the contribution of anammox to total N₂
368 production (r_a) and water depth ($r=0.59$, $p<0.01$; Fig. 8d). Previous studies have
369 reported similar findings, including those conducted on the Northeastern New
370 Zealand continental shelf (Cheung et al., 2024), the continental shelf and slope, North
371 Atlantic (Trimmer & Nicholls, 2009). The increased importance of anammox can be
372 attributed to the significant attenuation of denitrification with depth, as the availability
373 of organic carbon, which is essential for denitrification, decreases with increasing
374 water depth (Thamdrup, 2012). In addition to water depth, other factors such as
375 oxygen penetration depth, C/N ratios, and temperature may also influence the relative
376 importance of anammox. The contribution of anammox to total N₂ production (r_a)
377 was positively correlated with oxygen penetration depth ($r=0.7$, $p<0.01$; Fig. 8c). As
378 previously mentioned, denitrification decreases with higher oxygen penetration depth,
379 likely increasing the relative importance of anammox indirectly. Conversely, r_a
380 showed a decreasing trend with elevated C/N ratios ($r=-0.35$, $p<0.01$; Fig. 8b). High
381 C/N ratios may promote denitrification more significantly than anammox because
382 both processes tend to enhance with increasing C/N ratios, leading to a decrease in the
383 relative importance of anammox at sites with high C/N ratios. Additionally, r_a was
384 negatively correlated with temperature ($r=-0.29$, $p<0.01$; Fig. 8e), indicating that
385 denitrification is stimulated at higher temperatures compared to anammox.
386 Temperature-controlled experiments have confirmed that denitrification has a greater
387 optimal temperature than anammox (Canion et al., 2014; Tan et al., 2020). No



388 correlations were found between ra and other environmental factors, including
389 sediment organic carbon, water salinity, dissolved oxygen, nitrate, and ammonium
390 concentrations. (all $p > 0.05$; Fig. 8a, 8f, 8g, 8h, 8i).

391

392 **4 Applications of the database**

393 This database serves as a valuable resource for the broad scientific communities that
394 are interested in nitrogen cycle processes within coastal and marine ecosystems,
395 particularly those focusing on denitrification and anammox. The data is made
396 accessible as a basic database that will lead to a deeper understanding and generate
397 new scientific insights into the nitrogen cycles at the global scale. Potential
398 applications of this database include: (1) serving as a reference for comparing
399 denitrification and anammox rates across local, regional, and global scales in future
400 studies; (2) identifying and comparing the controlling factors of denitrification and
401 anammox at various spatial scales; (3) predicting the global biogeography of
402 denitrification and anammox in coastal and marine systems through machine learning;
403 and (4) providing essential data for the parameterization, validation and enhancement
404 of Earth system biogeochemical models.

405 **5 Conclusions**

406 We compiled a global database of denitrification and anammox measurements
407 obtained from core incubation experiments in coastal and marine sediments. To our



408 knowledge, no efforts have been made to compile actual nitrogen loss rates and
409 associated environmental factors in coastal and marine regions on a global scale. This
410 database offers valuable insights into the spatiotemporal variations and potential
411 controlling factors of denitrification and anammox, along with the contribution of
412 anammox to total N_2 production. It can be used to compare these two nitrogen loss
413 processes, assess the environmental controls on them at regional and global levels,
414 and support the parameterization and development of biogeochemical models.

415 **Data availability**

416 The data used in this study are openly available in Figshare repository at
417 <https://doi.org/10.6084/m9.figshare.27745770.v3> (Chang et al., 2024).

418 **Author contributions**

419 SJK and EHT conceived the research. YKC and EHT compiled the data. YKC, EHT,
420 DZG, CL and SJK participated in the data analysis. All co-authors contributed to the
421 writing and reviewing of this manuscript.

422 **Competing interests**

423 None declared.

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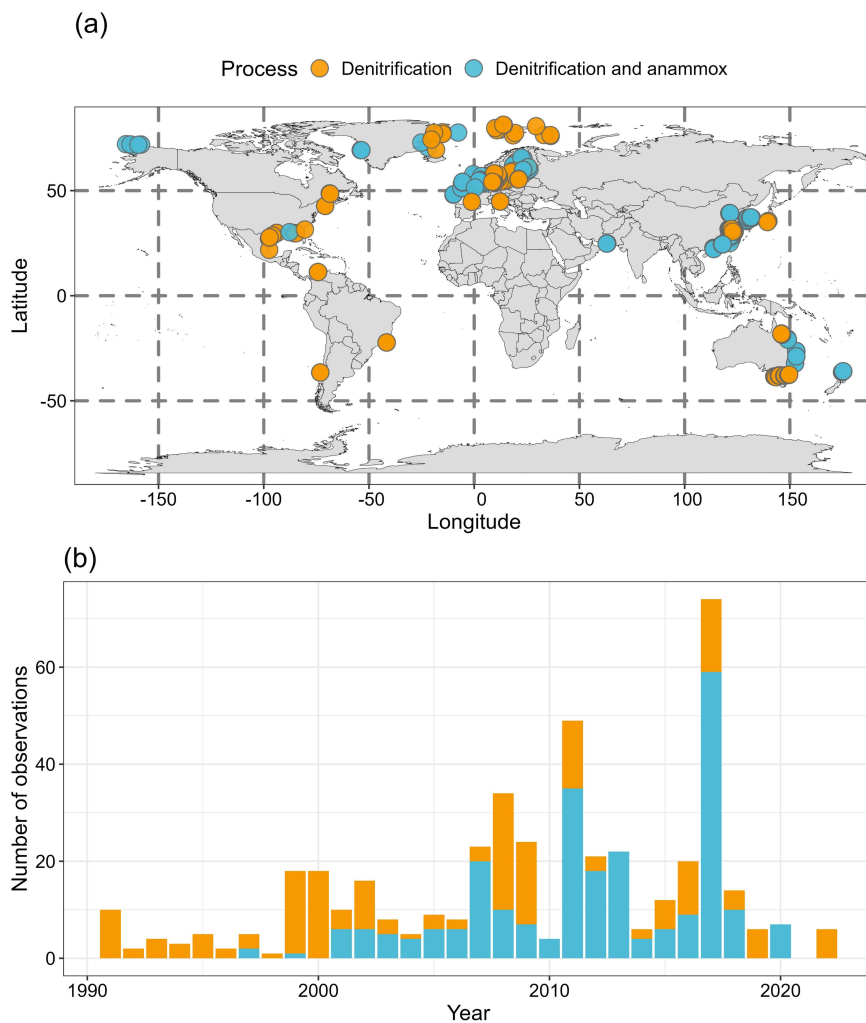
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792 **Figures and Table**

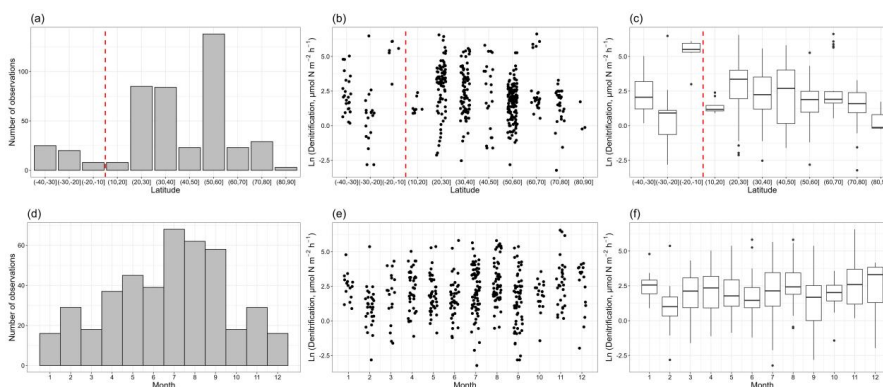
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795 **Figure 1** Map showing the sampling sites distribution of nitrogen loss rate
796 measurements (a) and the number of rate observations each year (b). Orange solid
797 points denote that only denitrification rates were measured. Cyan solid points denote
798 that both denitrification and anammox rates were measured.

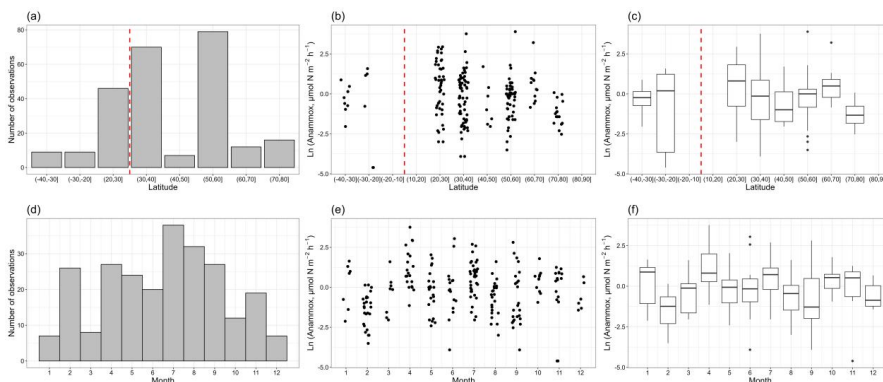
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801 **Figure 2** The observation numbers of denitrification (a, d) and denitrification rates (b, 802
 803 c, e, f) with latitudinal bands and months. A vertical dashed red line delimits the
 804 Southern Hemisphere and the Northern Hemisphere. Tops and bottoms of boxes in
 805 box plots denote the 25th and 75th percentiles, respectively. The horizontal lines
 806 inside the box plots represent the medians. Whiskers mark the minimum and
 807 maximum values within 1.5 times the interquartile range, with black points
 808 representing outliers beyond that range.

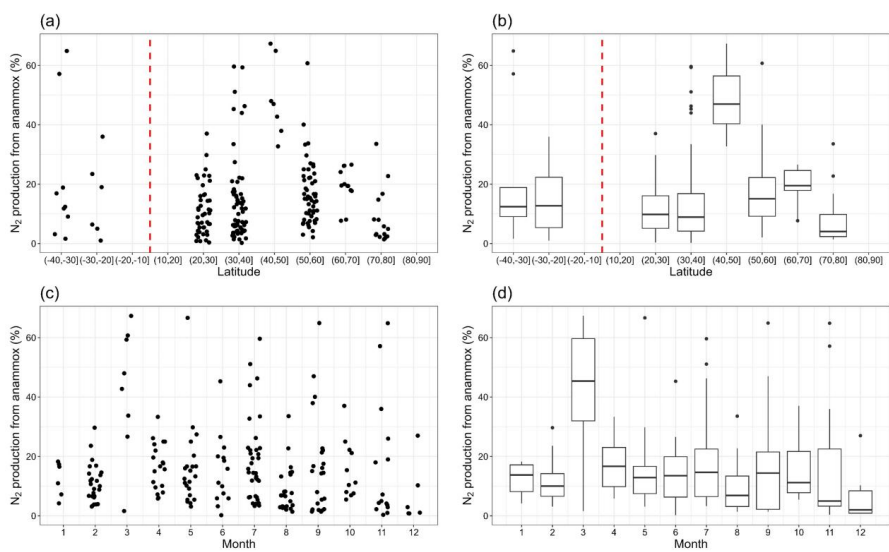
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810 **Figure 3** The observation numbers of anammox (a, d) and anammox rates (b, c, e, f)
 811 with latitudinal bands and months.

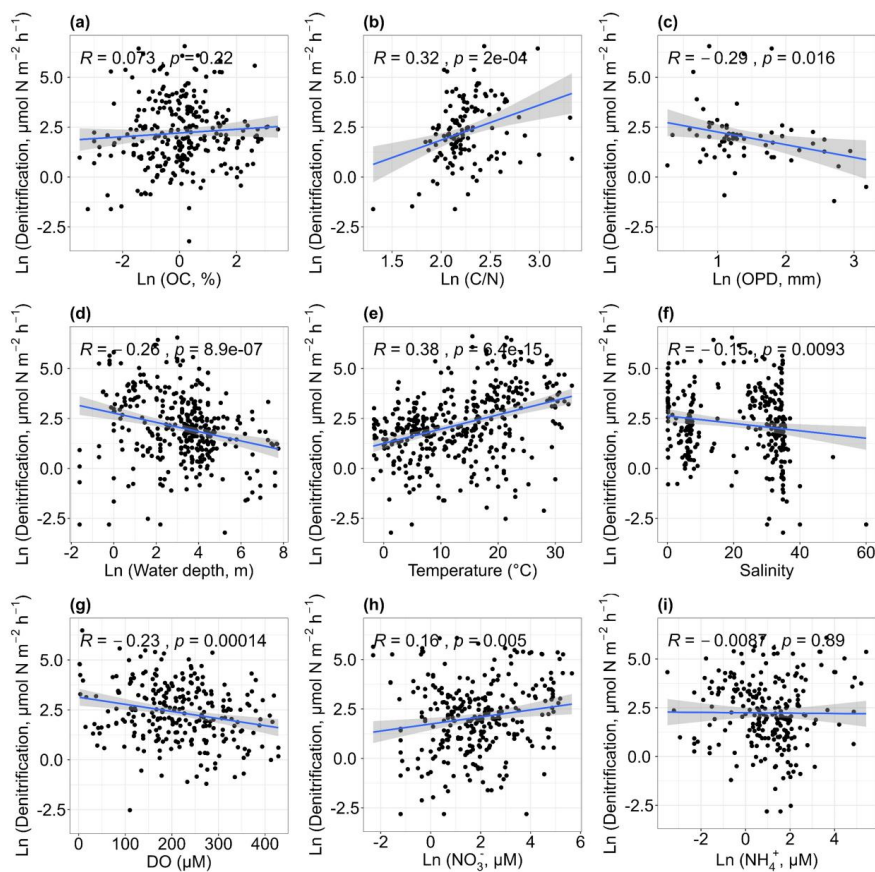
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814 **Figure 4** The contribution of anammox to total N_2 production with latitudinal bands
815 (a, b) and months (c, d).

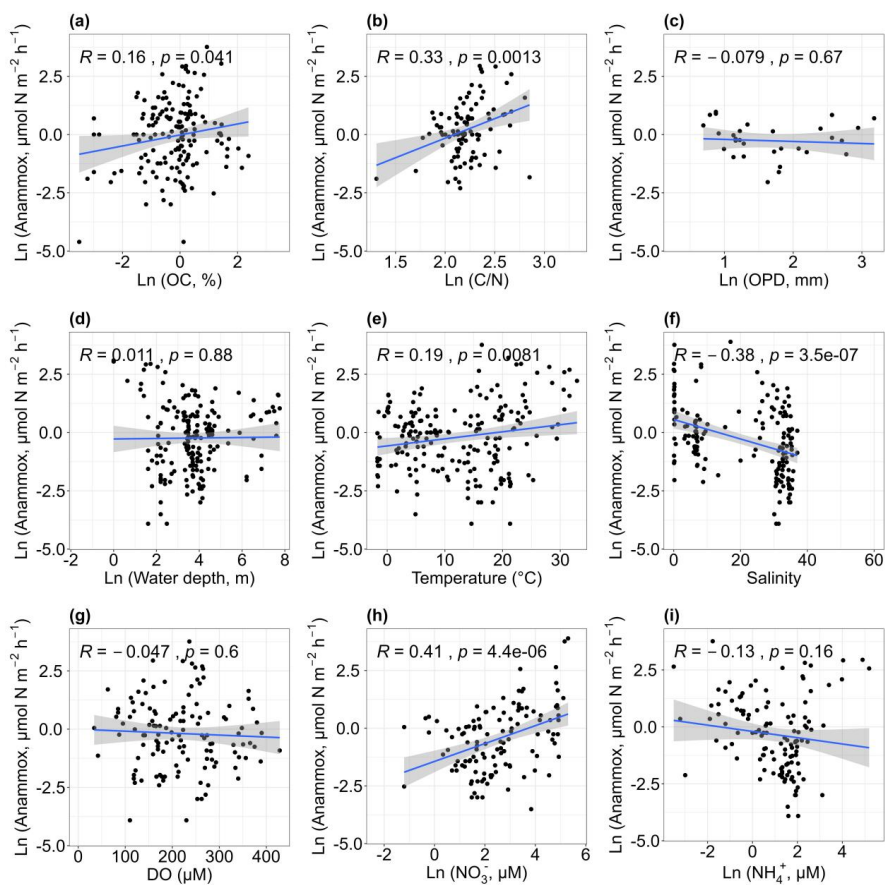
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818 **Figure 5** Relationships between denitrification rates and organic carbon [OC, (a)],
819 carbon-nitrogen ratios [C/N, (b)], oxygen penetration depth [OPD, (c)], water depth
820 (d), temperature (e), salinity (f), dissolved oxygen [DO, (g)], nitrate concentrations
821 [NO₃⁻, (h)] and ammonium concentrations [NH₄⁺, (i)].

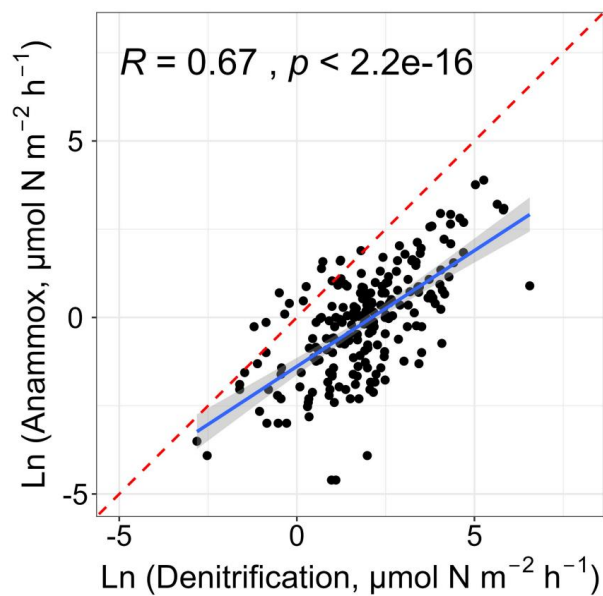
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824 **Figure 6** Relationships between anammox rates and organic carbon [OC, (a)],
825 carbon-nitrogen ratios [C/N, (b)], oxygen penetration depth [OPD, (c)], water depth
826 (d), temperature (e), salinity (f), dissolved oxygen [DO, (g)], nitrate concentrations
827 [NO₃⁻, (h)] and ammonium concentrations [NH₄⁺, (i)].

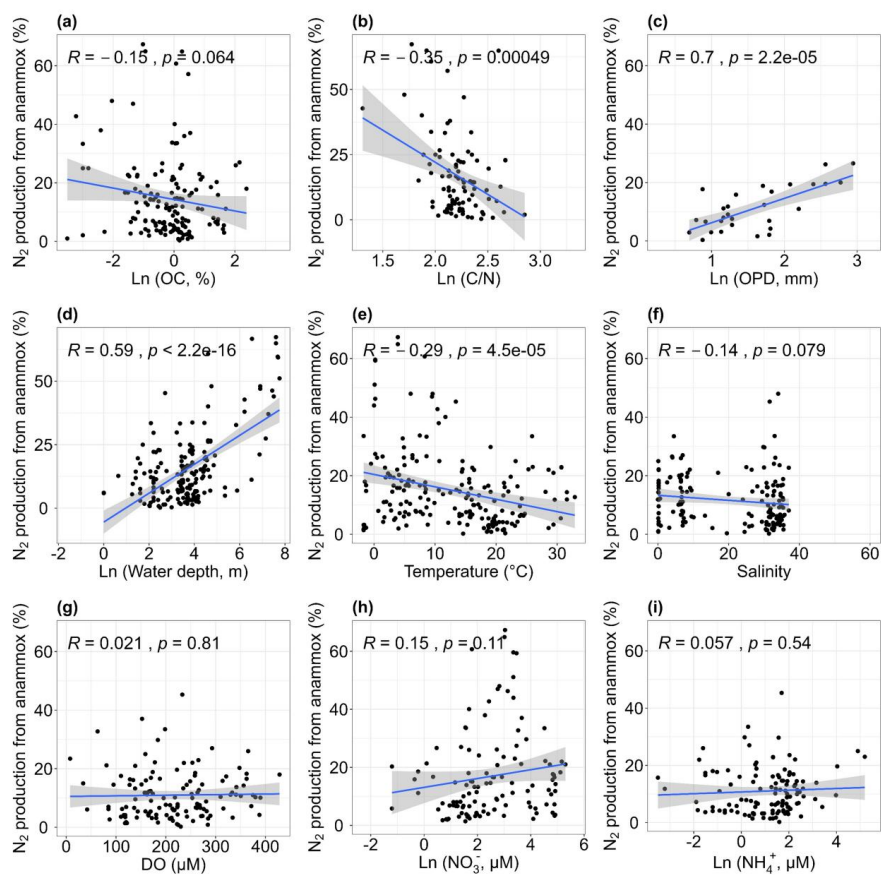
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830 **Figure 7** Relationships between denitrification and anammox rates. The blue solid
831 line and red dashed line denote the linear regression and 1:1 line, respectively.

832



833

834 **Figure 8** Relationships between the relative contribution of anammox to total N_2
835 production and organic carbon [OC, (a)], carbon-nitrogen ratios [C/N, (b)], oxygen
836 penetration depth [OPD, (c)], water depth (d), temperature (e), salinity (f), dissolved
837 oxygen [DO, (g)], nitrate concentrations [NO_3^- , (h)] and ammonium concentrations
838 [NH_4^+ , (i)].



839 **Table 1** Summary of the observations of actual nitrogen loss rates. The locations,
 840 observation numbers, core incubation methods and references are listed.

Sampling locations	Observation numbers	Core incubations	References
Aarhus Bight, Denmark	2	Intact core incubations	(Nielsen and Glud, 1996)
Arabian Sea	4	Intact core incubations	(Sokoll et al., 2012)
Arctic fjord (Svalbard, Norway)	3	Intact core incubations	(Gihring et al., 2010b)
Bassin d'Arcachon coastal lagoon	3	Intact core incubations	(Welsh et al., 2000)
Casino, NSW, Australia	2	Intact core incubations	(Erler et al., 2008)
central Sagami Bay, Japan	1	Intact core incubations	(Glud et al., 2009)
Changjiang estuary and its adjacent East China Sea	7	Intact core incubations	(Song et al., 2021)
Changjiang River Estuary and Jiulong River Estuary, China	23	Intact core incubations	(Tan et al., 2022)
Changjiang River Estuary, China	22	Continuous-flow experiments	(Liu et al., 2020)
Changjiang River Estuary, China	14	Continuous-flow experiments	(Liu et al., 2019)
Coast of Finland, northern Baltic Sea	10	Intact core incubations	(Hellemann et al., 2020)
Coast of Victoria, Australia	11	Intact core incubations	(Kessler et al., 2018)
Coastal area of the Gulf of Gdańsk	6	Intact core incubations	(Benelli et al., 2024)
Coastal lagoons, France	6	Intact core incubations	(Rysgaard et al., 1996b)
Coastal sediments, Greenland	11	Intact core incubations	(Rysgaard et al., 2004)
Continental shelf and slope, North Atlantic	12	Intact core incubations	(Trimmer and Nicholls, 2009)
Continental shelf region off central Chile	5	Intact core incubations	(Farias et al., 2004)
Danshuei River in northern Taiwan, China	1	Intact core incubations	(Hsu and Kao, 2013)
East China Sea	2	Intact core incubations	(Song et al., 2016)
Elbe Estuary, North Frisian	5	Intact core incubations	(Deek et al.,



Wadden Sea		incubations	2013)
Fjords in Svalbard and northern Norway	5	Intact core incubations	(Glud et al., 1998)
Georgia continental shelf, USA	2	Intact core incubations	(Vance-Harris and Ingall, 2005)
Great Barrier Reef lagoon	2	Intact core incubations	(Erler et al., 2013)
Gulf of Bothnia, Baltic Sea	7	Intact core incubations	(Bonaglia et al., 2017)
Gulf of Finland	5	Intact core incubations	(Susanna, 2007)
Gulf of Finland, Baltic Sea	11	Intact core incubations	(Jäntti and Hietanen, 2012)
Gulf of Finland, Baltic Sea	13	Intact core incubations	(Jäntti et al., 2011)
Gulf of Finland, Baltic Sea	5	Intact core incubations	(Hietanen and Kuparinen, 2008)
Gulf of Mexico	6	Intact core incubations	(Gihring et al., 2010a)
Gullmarsfjorden, Sweden and Thames Estuary, England	2	Intact core incubations	(Trimmer et al., 2006)
Hypoxic zone off the Changjiang River estuary, China	9	Intact core incubations	(Yang et al., 2022)
Jinpu Bay, China	12	Continuous-flow experiments	(Yin et al., 2015)
Jiulong River Estuary, China	2	Intact core incubations	(Wan et al., 2023)
Kattegat and Skagerrak	10	Intact core incubations	(Rysgaard et al., 2001)
Lawrence estuary	1	Intact core incubations	(Crowe et al., 2012)
Little Lagoon, USA	1	Continuous-flow experiments	(Bernard et al., 2015)
Noosa River estuary, Australia	5	Intact core incubations	(Chen et al., 2021)
North Sea	9	Intact core incubations	(Rosales Villa et al., 2019)
North Sea	1	Intact core incubations	(Fan et al., 2015)
North Sea	8	Intact core incubations	(Bale et al., 2014)
North Sea	16	Intact core incubations	(Neubacher et al., 2011)
Northeast Chukchi Sea	5	Continuous-flow	(McTigue et al.,



Northeastern New Zealand continental shelf	7	experiments	2016)
Northern Baltic Proper	17	Intact core incubations	(Cheung et al., 2024)
Northern East China Sea, China	16	Intact core incubations	(Bonaglia et al., 2014a)
Norwegian Trench, Skagerrak	4	Continuous-flow experiments	(Chang et al., 2021)
Öre Estuary, Swedish	6	Intact core incubations	(Trimmer et al., 2013)
Pearl River Estuary, China	5	Intact core incubations	(Hellemann et al., 2017)
Pearl River Estuary, China	5	Intact core incubations	(Tan et al., 2019)
Plum Island Sound, Massachusetts	4	Intact core incubations	(Koop-Jakobsen and Giblin, 2010)
Randers Fjord and Norsminde Fjord, Denmark	2	Intact core incubations	(Risgaard-Petersen et al., 2004)
Randers Fjord, Young Sound and Skagerrak, Danmark	3	Intact core incubations	(Risgaard-Petersen et al., 2003)
Sacca di Goro lagoon, Italy	6	Intact core incubations	(Magri et al., 2020)
Southern and central Baltic Sea	12	Intact core incubations	(Deutsch et al., 2010)
Southern Finland	5	Intact core incubations	(Uusheimo et al., 2018)
St. George Island, Gulf of Mexico, Hausstrand, German Wadden Sea and Spitsbergen island, Svalbard	5	Intact core incubations	(Canion et al., 2014)
St. Joseph Bay, USA	4	Continuous-flow experiments	(Hoffman et al., 2019)
St. Lawrence Estuary, Canada	3	Intact core incubations	(Poulin et al., 2007)
Stockholm Archipelago, Baltic Sea	1	Intact core incubations	(Bonaglia et al., 2014b)
Svalbard, Norway	10	Intact core incubations	(Blackburn et al., 1996)
Taganga Bay, Colombia Caribbean	8	Intact core incubations	(Arroyave Gómez et al., 2020)
Tama Estuary, Japan	2	Continuous-flow experiments	(Usui et al., 2001)
Texas estuaries, USA	26	Continuous-flow experiments	(Gardner et al., 2006)
The Baltic Sea	1	Intact core incubations	(Bonaglia et al.,



		incubations	2013)
		Intact core	(Bartoli et al.,
		incubations	2021)
		Intact core	(Enrich-Prast et
		incubations	al., 2016a)
		Intact core	(Adame et al.,
		incubations	2019b)
		Intact core	(Na et al., 2018)
		incubations	
		Intact core	(Erler et al., 2017)
		incubations	
		Intact core	(Rysgaard et al.,
		incubations	1996a)
841	Continuous-flow experiments denote continuous flow experiments combined with core		
842	incubations		