

1 **Rapid environmental responses to climate-induced hydrographic changes in**
2 **the Baltic Sea entrance**

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13 Abstract

14 The Öresund (the Sound), which is a part of the Danish straits, is linking the marine North Sea
15 and the brackish Baltic Sea. It is a transition zone where ecosystems are subjected to large
16 gradients in terms of salinity, temperature, carbonate chemistry, and dissolved oxygen
17 concentration. In addition to the highly variable environmental conditions, the area is responding
18 to anthropogenic disturbances in e.g. nutrient loading, temperature, and pH. We have
19 reconstructed environmental changes in the Öresund during the last c. 200 years, and especially
20 dissolved oxygen concentration, salinity, organic matter content, and pollution levels, using
21 benthic foraminifera and sediment geochemistry. Five zones with characteristic foraminiferal
22 assemblages were identified, each reflecting the environmental conditions for respective period.

23 The largest changes occurred around 1950, when the foraminiferal assemblage shifted from a
24 low diversity fauna, dominated by the species *Stainforthia fusiformis* to higher diversity and
25 abundance, and dominance of the *Elphidium* species. Concurrently, the grain-size distribution
26 shifted from clayey — to sandier sediment. To explore the causes for the environmental changes,
27 we used time-series of reconstructed wind conditions coupled with large-scale climate variations
28 as recorded by the North Atlantic Oscillation (NAO) index, as well as the ECOSMO II model of
29 currents in the Öresund area. The results indicate increased changes in the water circulation
30 towards stronger currents in the area after the 1950's. The foraminiferal fauna responded quickly
31 (< 10 years) to the environmental changes. Notably, when the wind conditions, and thereby the
32 current system, returned in the 1980's to the previous pattern, the foraminiferal assemblage did
33 not rebound. Instead, the foraminiferal faunas displayed a new equilibrium state.

34 1 – Introduction

35 The Öresund (the Sound) is one part of the Danish straits between Sweden and Denmark.
36 Together with the Great — and Little Belt, they link the open-ocean waters of the North Sea and
37 the brackish waters of the Baltic Sea. The confluence of the water masses creates a north-south
38 gradient as well as a strong vertical stratification of the water in terms of salinity, carbonate
39 chemistry and dissolved oxygen concentration ($[O_2]$) (Leppäranta and Myrberg 2009). The depth
40 of the halocline mainly depends of the outflows from the Baltic Sea; a strong thermocline
41 develops during spring and summer, which further strengthens the vertical stratification. Thus,
42 the ecosystems in the Öresund are exposed — and adapted — to a unique transitional
43 environment. The region is also characterized by intense human activities, with 4 million people
44 living in the vicinity of the Öresund and 85 million people living in the catchment area of the
45 Baltic Sea (HELCOM, 2009). Discharge from agriculture, industry, and urban areas on both the

46 Swedish and Danish sides of the strait, and the considerable impact of marine traffic – the strait
47 is one of the busiest waterways in the world – generate pollution and eutrophication of the water
48 (HELCOM 2009; ICES 2010). Since the 1980's, the implementation of efficient wastewater
49 treatment and measures in agriculture contributed to markedly reduce the amount of nutrients
50 coming from river run-off (Nausch et al. 1999; Carstensen et al. 2006; Rydberg et al. 2006).
51 However, these efforts in decreasing nutrient loads have not resulted in improved water quality,
52 due to the long time scales of biogeochemical cycles to reach equilibrium in the Baltic Sea
53 region (Gustafsson et al. 2012). The Öresund, like most of the Baltic Sea, is still assessed to be
54 eutrophic, and hypoxic events are frequent (Rosenberg et al. 1996; Conley et al. 2007, 2011;
55 HELCOM 2009; Wesslander et al. 2016). Moreover, increasing temperatures and declining pH,
56 linked to global climate change and ocean acidification, have been reported for surface and
57 bottom waters in the area (Andersson et al. 2008; Göransson 2017). As a result, ecosystems in
58 the Öresund are currently under the combined impact of natural and anthropogenic stressors
59 (Henriksson 1969; Göransson et al. 2002; HELCOM 2009; ICES 2010). The multiple stressors
60 currently affecting the environment make this region particularly interesting to study, and also
61 highlight the need to obtain records of decadal and centennial environmental changes. As noted
62 above, both recent human-induced impacts and climate variability have been substantial in the
63 region. Therefore the question arises whether these factors have affected the benthic environment.
64 Furthermore, sediment records of past environmental changes can provide crucial context for
65 ongoing and future predicted changes in the Öresund and Baltic Sea regions.

66 We used the marine sediment record and its contents of foraminifera as well as sediment
67 geochemistry to obtain records of decadal environmental changes. Benthic foraminifera are
68 widely used for environmental reconstructions, based on their rapid response to environmental

69 changes, broad distribution, high densities, and often well-preserved tests (shells) in the sediment
70 (e.g. Sen Gupta 1999b; Murray 2006). For instance, distribution of benthic foraminifera have
71 been used for historical environmental reconstructions of fjords on decadal to centennial
72 timescales on the Swedish west coast (Nordberg et al. 2000; Filipsson and Nordberg 2004a,
73 2004b; Polovodova Asteman and Nordberg 2013; Polovodova Asteman et al. 2015), and in the
74 Kattegat (Seidenkrantz 1993; Christiansen et al. 1996). In the Öresund, living foraminiferal
75 assemblages have been studied (Hansen 1965; Charrieau et al. 2018), but to the best of our
76 knowledge, no studies of past foraminiferal assemblages have been performed. The objective of
77 this study was to reconstruct the environmental conditions of benthic systems during the last two
78 centuries in the Öresund, by using foraminiferal fauna analysis in combination with sediment
79 geochemistry and grain-size. Furthermore, we analyzed long time series of wind conditions in
80 the area to evaluate the coupling between local changes in ecosystem variables and variations in
81 atmospheric and subsequent hydrographic conditions, and a possible link with large-scale
82 variations expressed through the North Atlantic Oscillation (NAO) index. Finally, we compared
83 our data with the model ECOSMO II (Daewel and Schrum 2013; 2017) of currents and water
84 circulation changes in the Öresund area during the period 1948–2013.

85 2 – Study site

86 The Öresund is a 118 km long narrow strait (Figure 1). The water depth in the northern part is on
87 average 24 m but it reaches 53 m south of the Island of Ven. The Öresund is an important link
88 between the North Sea, Skagerrak, Kattegat and the Baltic Sea (Figure 1), and up to 30 % of the
89 water exchange in the region goes through the Öresund (Sayin and Krauß 1996; Leppäranta and
90 Myrberg 2009). The remaining part goes through the Great and Little Belt. The width of the
91 Öresund varies between 4 and 28 km, and the water has overall high current velocities, up to 1.5

92 $\text{m}\cdot\text{s}^{-1}$ at the upper water layer in the northern part (Nielsen 2001). The fully marine Skagerrak
93 consists of water masses from the North Sea and the North Atlantic and in general a thin surface
94 layer with water originating from the Baltic Sea and rivers draining into the sea; the water
95 circulation forms a cyclonic gyre (cf. Erbs-Hansen et al. 2012). Part of the Skagerrak waters
96 reach the Kattegat and the Baltic Sea, where they are successively diluted with the large amounts
97 of freshwater (around $15,000 \text{ m}^3/\text{s}$, Bergström and Carlsson 1994) draining into the Baltic Sea
98 from numerous large rivers. The low-saline Baltic Sea surface water is transported by the Baltic
99 Current, which is typically confined along the Swedish west coast in the Kattegat but may cover
100 a larger surface area towards the west, depending on wind direction. The Baltic Current later
101 joins the Norwegian Coastal Current in the Skagerrak (Figure 1). The large fresh water input and
102 the subsequent large salinity difference between the Kattegat and Baltic Sea result in a two-layer
103 structure in the Öresund (Figure 2) (She et al. 2007; Leppäranta and Myrberg 2009). The water
104 stratification is influenced by the surface water from Arkona Basin (salinity 7.5—8.5), the
105 surface water from the Kattegat upper layer (salinity 18—26) and the lower layer of the Kattegat
106 (salinity 32—34).

107 Salinity, temperature, pH, $[\text{O}_2]$ and nutrient content, here represented by dissolved inorganic
108 nitrogen concentration [DIN] (nitrate + nitrite + ammonium), in the surface and bottom waters of
109 the Öresund vary seasonally (Figure 3, Appendix A). At the surface and bottom water, salinity
110 ranges between ~ 8 and ~ 18 and between ~ 29 and ~ 34 , respectively, and it is more stable between
111 April and July, when the stratification is the strongest (Figure 3). Temperature ranges between
112 ~ 1 °C in February and ~ 19 °C in July in the surface water, while in the bottom water, the lowest
113 temperature is found in March—April with ~ 5 °C, and the highest temperature in October—
114 November with ~ 13 °C. The pH varies between ~ 8.1 and ~ 8.6 in the surface water, and between

115 ~7.8 and ~8.6 in the bottom water, without a clear seasonal pattern (Figure 3). [O₂] in the bottom
116 water reaches ~7 mL.L⁻¹ in January, and it is typically below 2 mL.L⁻¹ in October, approaching
117 hypoxic values. In the surface water, [DIN] can reach ~7 μmol.L⁻¹ in January, and it is ~0
118 μmol.L⁻¹ between April and August (Figure 3).

119 3 – Materials and Methods

120 3.1 Sampling

121 A suite of sediment cores, as well as water samples from the water column, were collected in
122 November 2013 during a cruise with r/v *Skagerak*. Here we present the data from two sediment
123 cores sampled at the Öresund station DV-1 (55°55.59' N, 12°42.66' E) (Figure 1), north of the
124 Island of Ven. The water depth was 45 m, and CTD (Conductivity, Temperature, Depth) casts
125 were taken to measure salinity, temperature and [O₂] in the water column. Water samples were
126 collected at 10, 15, 20, 30 and 43 m from the Niskin bottles for carbonate chemistry analyses.
127 The CTD and carbonate chemistry data are presented in Charrieau et al. (2018). In general, it is
128 challenging to obtain sediment cores in the Öresund, due to the high current velocities up to 1.5
129 m.s⁻¹ (Nielsen 2001), human-induced disturbances, and limited areas of recent sediment
130 deposition (Lumborg 2005), but our site north of Ven represents an accumulation area. The cores
131 (9-cm-inner-diameter) were collected using a GEMAX twin barrel corer. The corer allowed
132 sampling of 30 and 36 cm long sediment cores (referred in this study as core DV1-G and DV1-I,
133 respectively), which were sliced into one centimeter sections. The samples from the DV1-G core
134 were analyzed for carbon and nitrogen content, grain size distribution, and dated using Gamma
135 spectroscopy. The samples from the DV1-I core were analyzed with respect to foraminiferal
136 fauna and carbon and nitrogen content. The distinct carbon content profiles, measured on both

137 cores, were used to correlate the ^{210}Pb dated DV1-G core to the DV1-I core used for
138 foraminiferal analyses.

139 3.2 Chronology

140 The age-depth model was established using ^{210}Pb and ^{137}Cs techniques on samples from the
141 DV1-G core. The samples were measured with an ORTEC HPGe (High-Purity Germanium)
142 Gamma Detector at the Department of Geology at Lund University, Sweden. Corrections for
143 self-absorption were made for ^{210}Pb following Cutshall et al. (1983). The instruments were
144 calibrated against in-house standards and the maximum error was 0.5 year in the measurements.
145 Excess (unsupported) ^{210}Pb was measured down to 23 cm and the age model was calculated
146 based on the Constant Rate of ^{210}Pb Supply (CRS) model (Appleby 2001).

147 3.3 Foraminifera analyses

148 Approximately 10 g of freeze-dried sediment per sample were wet sieved through a 63- μm mesh
149 screen and dried on filter paper at room temperature. Subsequently, the samples were dried
150 sieved through 100- and 500- μm mesh screens and separated into the fractions 100-500 μm and
151 >500 μm . The foraminifera from every second centimeter of the core - plus from additional
152 centimeters around key zones - were picked and sorted under a Nikon microscope (22 samples in
153 total). A minimum of 300 specimens per sample were picked and identified, as recommended by
154 Patterson and Fishbein (1989). If necessary the samples were split with an Otto splitter (Otto
155 1933). For taxonomy at the genus level, we mainly followed Loeblich and Tappan (1964) with
156 some updates from more recent literature, e.g. Tappan and Loeblich (1988). For taxonomy at the
157 species level, we mainly used Feyling-Hanssen (1964), Feyling-Hanssen et al. (1971) and

158 Murray and Alve (2011). For original descriptions of the species, see Ellis and Messina (1940
159 and supplements up to 2013).

160 Recently, the eastern Pacific morphospecies *Nonionella stella* has been presented as an invasive
161 species in the Skagerrak-Kattegat region (Polovodova Asteman and Schönfeld 2015). However,
162 a comparison of *N. stella* DNA sequences from the Santa Barbara Basin (USA) (Bernhard et al.
163 1997) with the Swedish west coast specimens demonstrates that they represent two closely
164 related species but are not conspecific (Deldicq et al. in press). Therefore, we have referred to the
165 species found here as *Nonionella* sp. T1, following Deldicq et al. (2019). The species
166 *Verneuilina media* (here referred to the genus *Eggerelloides*), which has often been reported in
167 previous studies from the Skagerrak-Kattegat area (e.g. Conradsen et al. 1994), was
168 morphologically close to *Eggerelloides scabrus* in the present material, and these two species
169 have been grouped as *E. medius/scabrus*. The taxon *Elphidium excavatum* forma *clavata* (cf.
170 Feyling-Hanssen 1972), was referred to as *Elphidium clavatum* following Darling et al. (2016).
171 *Elphidium clavatum* and *Elphidium selseyense* (Heron-Allen and Earland) were morphologically
172 difficult to separate in this region, as transitional forms occur. The dominant species was *E.*
173 *clavatum*, but we acknowledge that a few individuals of *E. selseyense* could have been included
174 in the counts. The taxon *Ammonia beccarii* was referred to as *Ammonia batava*, following recent
175 molecular work done on the taxon *Ammonia* in the Kattegat region (Groeneveld et al. 2018; Bird
176 et al. 2019)

177 Foraminiferal density was calculated and normalized to the number of specimens per 50 cm³.
178 Data of densities for the first two centimeters of the core are from Charrieau et al. (2018). Some
179 specimens displayed decalcified tests, however the inner organic linings were preserved. These

180 inner organic linings were reported separately and not included in the total foraminiferal counts.

181 Benthic foraminiferal accumulation rates were calculated as follows:

182 $\text{BFAR (number of specimens.cm}^{-2}\text{.yr}^{-1}) = \text{BF} \times \text{SAR},$

183 where BF is the number of benthic foraminifera per cm^3 and SAR is the sediment accumulation

184 rate (cm.yr^{-1}). Foraminiferal species that accounted for >5 % of the total fauna in at least one of

185 the samples were considered as major species, and their density was used in statistical analysis.

186 The Shannon index was calculated to describe the foraminiferal diversity. To determine

187 foraminiferal zones, stratigraphically constrained cluster analysis was performed, using the size-

188 independent Morisita's index to account for the large differences in the densities between

189 samples (e.g. Krebs 1998). A dendrogram was then constructed based on arithmetic averages

190 with the UPGMA method (Unweighted Pair Group Method with Arithmetic Mean).

191 Correspondence analysis was also performed, to determine significant foraminiferal species in

192 each zone. Statistical analyses were performed using the PAST software (Hammer et al. 2001).

193 3.4 Organic matter analyses

194 Total Organic Carbon (TOC) and Total Nitrogen (TN) content were measured for both DV1-G

195 and DV1-I. Approximately 8 mg of freeze-dried sediment was homogenized for each centimeter

196 and placed in silver capsules. Removal of inorganic carbon was carried out by in-situ

197 acidification (2M HCl) method based on Brodie et al. (2011). TOC and TN content were

198 analyzed on a Costech ECS 4010 Elemental Analyzer at the Department of Geology, Lund

199 University. The instrument was calibrated against in-house standards. The analytical precisions

200 showed a reproducibility of 0.2 % and 0.03 % for TOC and TN contents, respectively. The molar

201 C/N ratio was calculated.

202 3.5 Grain-size analyses

203 Grain-size analyses were performed on core DV1-G using 3.5 to 5 g of freeze-dried sediment for
204 each centimeter. Organic matter was removed by adding 15 mL of 30 % H₂O₂ and heating
205 during 3 to 4 minutes until the reaction ceased. After the samples had cooled down, 10 mL of
206 10 % HCl was added to remove carbonates; thereafter the sediment was washed with milli-Q
207 until its pH was neutral. In the last step, biogenic silica was removed by boiling the sediment in
208 100 mL 8 % NaOH, and then washed until neutral pH was reached. The sand fraction (>63 µm)
209 was separated by sieving and the mass fraction of sand of each sample was calculated. Grain
210 sizes <63 µm were analyzed by laser diffraction using a Sedigraph III Particle Size Analyzer at
211 the Department of Geology, Lund University. The data were categorized into three size groups,
212 <4 µm (clay), 4–63 µm (silt) and 63–2000 µm (sand).

213 3.6 Climate data and numerical modeling

214 Data from the dataset High Resolution Atmospheric Forcing Fields (HiResAFF) covering the
215 time period 1850–2008 (Schenk and Zorita 2012; Schenk 2015) were used to study the variations
216 of near-surface (10 m) wind conditions during the winter half of the year (October to March).
217 The daily dataset can be downloaded from WDC Climate (Schenk 2017). Wind conditions over
218 the Öresund are represented by the closest grid point of HiResAFF at 55° N and 12.5° E. The
219 North Atlantic Oscillation (NAO) index as defined by Jones et al. (1997) for boreal winter
220 (December to March) was used, with updates taken from the Climate Research Unit (CRU,
221 <https://crudata.uea.ac.uk/cru/data/nao/>). To allow comparison, the NAO and wind data were
222 normalized relative to the period 1850–2008. Changes in the currents through the Öresund and
223 the Kattegat were taken from the fully coupled physical biogeochemical model ECOSMO II

224 (Daewel and Schrum 2013, 2017), which was forced by NCEP/NCAR reanalysis data and covers
225 the period 1950–2013. On model ECOSMO II, the simulated South-North currents are
226 represented as VAV (vertically averaged V- component) and the simulated West-East currents as
227 UAV (vertically averaged U - component).

228 4 – Results

229 4.1 Age model

230 The unsupported ^{210}Pb showed a decreasing trend with depth in the DV1-G core (Figures 4A,
231 4B). The peak observed in the ^{137}Cs around 9 cm corresponds to the Chernobyl accident in 1986
232 (Figure 4C). The unsupported ^{210}Pb allowed direct dating of the core between 2013 and 1913.
233 The sedimentation rate ranged between 1 and 5.6 mm.y^{-1} , with an average of 2.2 mm.y^{-1} , and
234 was decreased with depth. The ages of the lower part of the sediment record were deduced by
235 linear extrapolation based on a sedimentation rate of 1.4 mm.y^{-1} , corresponding to the linear
236 mean sedimentation rate between the years 1913 and 1946 (Figure 4D).

237 4.2 Foraminiferal assemblages and sediment features

238 The foraminiferal assemblages were composed of 76 species from the porcelaneous, hyalines
239 and agglutinated forms (0.3, 54.5 and 45.2 %, respectively) (Appendix B). Eleven foraminiferal
240 species had relative abundance higher than 5 % in at least one sample and were considered as
241 major species (Plate 1, Figure 5).

242 The cluster analysis revealed three main foraminiferal zones (FOR-A, FOR-B, and FOR-C)
243 (Figures 5, 6). The correspondence analysis resulted in three factors explaining 92 % of the
244 variance, and in assemblages consisting in seven significant species, presented in order of

245 contribution: *Nonionella* sp. T1, *Nonionoides turgida*, *Ammonia batava*, *Stainforthia fusiformis*,
246 *Elphidium albiumbilicatum*, *E. clavatum* and *Elphidium magellanicum* (Table 1). Based both on
247 the cluster and the correspondence analyses, five subzones could be separated to which we
248 assigned dates according to the age model: FOR-A1 (1807–1870), FOR-A2 (1870–1953), FOR-
249 B1 (1953–1998), FOR-B2 (1998–2009), and FOR-C (2009–2013) (Figures 5, 6).

250 4.2.1 Zone FOR-A1 (1807–1870)

251 The foraminiferal accumulation rate (BFAR) was on average 5 ± 3 specimens.cm⁻².y⁻¹ in zone
252 FOR-A1 (Figure 5). The Shannon index was stable and low, around 1.77 ± 0.1 (Figure 5). The
253 agglutinated species *Eggerelloides medius/scabrus* and the hyaline species *Stainforthia*
254 *fusiformis* made major contributions to the assemblages (relative abundances up to 53 % and
255 34 %, respectively; Figure 5A). *Ammonia batava*, the three *Elphidium* species (*E.*
256 *albiumbilicatum*, *E. clavatum*, and *E. magellanicum*), *Nonionellina labradorica* and the
257 agglutinated species *Reophax subfusiformis* were also major species with abundances up to 7 %.
258 The TOC and C/N values on this period were stable and were on average 3.36 % and 8.8 %,
259 respectively (Figure 7). The clay size fraction dominated the sediment at the end of this period
260 with a mean value of 63 %, and the sand content was around 7 % (Figure 7).

261 4.2.2 Zone FOR-A2 (1870–1953)

262 The BFAR was on average 9 ± 5 specimens.cm⁻².y⁻¹ in zone FOR-A2 (Figure 5). The Shannon
263 index was stable and low, around 1.94 ± 0.15 (Figure 5). *Stainforthia fusiformis* dominated the
264 assemblage with relative abundances up to 56 % and BFAR up to 608 specimens.cm⁻².y⁻¹
265 (Figures 5A, 5B), which is the highest BFAR observed for this species along the core.
266 *Eggerelloides medius/scabrus* was still very abundant, up to 48 % (Figure 5A). *Ammonia batava*,

267 the three *Elphidium* species and *N. labradorica* were present but with lower abundances than in
268 the zone FOR-A1 (maximum 5 %). *Bulimina marginata* started to be more abundant with an
269 average relative abundance of 2 % in the zone. *Reophax subfusiformis* was still a part of the
270 assemblage and ranged between 1 and 8 %. The TOC and C/N values were stable and were on
271 average 3.5 % and 8.74 %, respectively (Figure 7). The clay size fraction dominated the
272 sediment during this period with a mean value of 63 %, and the sand content was around 6 %
273 (Figure 7).

274 4.2.3 Zone FOR-B1 (1953–1998)

275 The BFAR increased massively during the zone FOR-B1 with on average 54 ± 31 specimens.cm⁻².y⁻¹
276 and with a peak at 93 specimens.cm⁻².y⁻¹ around 1965 (Figure 5). It is lower during the
277 second part of the zone. The Shannon index was higher than in previous zones and it
278 progressively increased towards the top of the zone (Shannon index average 2.34 ± 0.3) (Figure
279 5). The highest BFAR along the core were observed for all the dominant species of the previous
280 zone FOR-A2, except for *S. fusiformis* (Figure 5B). The zone was then also characterized by a
281 drastic drop in the relative abundance of *S. fusiformis* from 31 to 2 % (Figure 5A).

282 *Eggerelloides medius/scabrus* gradually decreased in the zone, with relative abundances from
283 49 to 24 %. The highest relative abundance of *A. batava* for the entire record was in this zone but
284 it was slowly decreasing as well, from 10 to 3 %. The *Elphidium* species were more abundant
285 than in the FOR-A zones and their relative abundance was increasing, especially for *E. clavatum*
286 (increasing up to 23 %). *Bulimina marginata*, *N. labradorica* and *R. subfusiformis* had a relative
287 abundance between 2 and 6 %. A period of lower TOC values was observed during zone FOR-
288 B1 between 1953 and 1981, with an average of 2.38 % (Figure 7). On the same period, the sand
289 content showed a pronounced increase, with an average of 24 % (Figure 7).

290 4.2.4 Zone FOR-B2 (1998–2009)

291 In zone FOR-B2 the BFAR was still high, on average 55 ± 6 specimens.cm⁻².y⁻¹ (Figure 5). The
292 Shannon index was high with an average of 2.8 ± 0.2 (Figure 5). The dominant species in the
293 zone were *E. clavatum* (up to 25 %) and *Eggerelloides medius/scabrus* (up to 15 %; Figure 5A).
294 The other two *Elphidium* species reached their highest relative abundances over the core (up to
295 6 %). *Nonionella* sp. T1, which had not occurred in the record until now, appeared in this zone
296 with a relative abundance of 1 %. *Nonionoides turgida*, which was present in very low
297 abundances along the core, had a mean abundance of 1 % in the zone (Figure 6A). *Stainforthia*
298 *fusiformis* was present with up to 9 % in relative abundance and a BFAR higher than in zone
299 FOR-B1 (up to 570 specimens.cm⁻².y⁻¹). *Ammonia batava*, *B. marginata*, *N. labradorica*, and *R.*
300 *subfusiformis* were present and ranged between 2 and 8 %. The TOC values were increasing,
301 with on average 3.05 % (Figure 7). The sediment was dominated by the clay fraction that was
302 increasing (mean value of 58 %), and the sand content was around 17 % (Figure 7).

303 4.2.5 Zone FOR-C (2009–2013)

304 The BFAR was lower than in previous zones FOR-B1 and FOR-B2, with on average 21 ± 5
305 specimens.cm⁻².y⁻¹ (Figure 5). The Shannon index was the highest during FOR-C (Shannon
306 index average 2.93 ± 0.07) (Figure 5). *Nonionella* sp. T1 was a dominant specie in the zone with
307 a strong increase in relative abundance (from 1 to 14 %) and in BFAR (from 61 to 137
308 specimens.cm⁻².y⁻¹) (Figures 5A, 5B). *Elphidium clavatum* and *R. subfusiformis* were also
309 dominant species with abundances up to 13%. *Nonionoides turgida* had its highest relative
310 abundance and BFAR over the core during the zone, with up to 9 % and 342 specimens.cm⁻².y⁻¹,
311 respectively (Figures 5A, 5B). *Eggerelloides medius/scabrus* had its lowest relative abundance

312 over the core (up to 9 %). *Bulimina marginata*, the other two *Elphidium* species, *N. labradorica*
313 and *S. fusiformis* were still present (between 1 and 6 %), while *Ammonia batava* was absent
314 during the zone. The TOC and C/N values were on average 3.71 % and 8.17 %, respectively
315 (Figure 7). The clay size fraction dominated the sediment with a mean value of 66 % and the
316 sand fraction was 7 % (Figure 7).

317 4.2.6 Inner organic linings

318 Decalcified specimens were few and ranged between 0 and 4 specimens.cm⁻².y⁻¹ with an average
319 of 1 specimen.cm⁻².y⁻¹ (Fig. 5). They were observed throughout the core and especially during
320 zone FOR-B2, and the morphology of the remaining inner organic linings allowed the
321 identification of the taxon *Ammonia* (Plate 1).

322 4.3 Simulated data from model ECOSMO II

323 The VAV (vertically averaged South-North current velocity) through the Öresund from the
324 model ECOSMO II showed a reversed pattern compared to the UAV (vertically averaged West-
325 East current velocity) through the Kattegat (Figure 8). Thus, higher VAV through the Öresund
326 translates to an increase in the East to West flow in the Kattegat (lower UAV), suggesting a
327 stronger outflow from the Baltic Sea. The VAV through the Öresund had the lowest values
328 around 1955 (Figure 8), followed by a shift to very high values, which dominated throughout
329 1960–70. A comparable period with increased outflow from the Baltic into the Kattegat re-
330 occurred during the period 1993–2000.

331 5 – Discussion

332 Our environmental interpretations of the foraminiferal assemblages were based on the ecological
333 characteristics of each major species (Table 2). Based on our environmental reconstructions, we
334 could infer environmental changes regarding [O₂], salinity, organic matter content, and pollution
335 levels. Furthermore, we linked local environmental changes to larger atmospheric and
336 hydrographic conditions.

337 5.1 1807 – 1870

338 All the major species found in this period are tolerant to low oxygen conditions, especially the
339 two main species: *S. fusiformis* and *E. medius/scabrus* (Table 2). *Stainforthia fusiformis* is an
340 opportunistic species used to hypoxic and potentially anoxic conditions (Alve 1994), and *E.*
341 *medius/scabrus* specimens have been found alive down to 10 cm in the sediment, where no
342 oxygen was available (Cesbron et al. 2016). *Stainforthia fusiformis* and *N. labradorica* are also
343 able to denitrify (Piña-Ochoa et al. 2010). The fact that species tolerant to low oxygen conditions
344 dominated, and the presence of species that have the capacity to denitrify, suggest that low
345 oxygen conditions were prevailing during this period. Furthermore, *S. fusiformis* prefers organic
346 rich substrate and clayey sediment, which was measured in our core during this time period
347 (Figure 7). The low species diversity, as indicated by the low Shannon index in this section of
348 the core, can sometimes be linked with low salinity (Sen Gupta 1999a). Most of the major
349 species found during this period, such as the *Elphidium* species, *R. subfusiformis* and *A. batava*
350 tolerate lower salinities, and are typical of brackish environments (Table 2). The low occurrence
351 of *B. marginata*, a typical marine species, also suggests a salinity lower than in the open ocean.
352 However, the salinity was probably not below ~30, which is the lower limit for *N. labradorica*
353 and *S. fusiformis*, which were present throughout the period (Figure 5, Table 2). In summary, this

354 period appears to have been characterized by low [O₂], high organic matter content, and salinity
355 around 30.

356 5.2 1870 – 1953

357 *Stainforthia fusiformis* was largely dominating the assemblage during this period, which may
358 suggest even lower oxygen conditions than during the previous period. This would also go along
359 with the low species diversity, which is sometimes linked to low salinity. In the Öresund, low
360 salinity can be caused by less influence of more saline marine waters from the Kattegat, and
361 changes in the water transport through the strait is a possible explanation for both lower salinity
362 and oxygen levels. However, the occurrence of the marine species *B. marginata* suggests that the
363 salinity was at least ~30 (Table 2). Low oxygen can also be associated with high organic matter
364 contents, since oxygen is consumed during remineralization of organic matter. However, the
365 TOC levels observed in our core in this zone were high, but not higher than in the previous zone
366 (Figure 7). At the time of the industrial revolution, the Öresund, as the Baltic Sea in general, was
367 used as a sewage recipient for a mixture of domestic and industrial wastes, industrial cooling
368 water and drainage water (Henriksson 1968), and the amount of marine traffic increased
369 considerably during this time period. Across the Baltic Sea, this notably caused increased
370 deposition of heavy metals (Borg and Jonsson 1996). This diverse type of pollution could have
371 modified the water properties, for example regarding the carbonate chemistry and pH. Indeed,
372 this zone is characterized by the presence of organic linings in the core (see also section 5.6).
373 Moreover, heavy metals, fuel ash (black carbon) and pesticides have been demonstrated to
374 generally have a negative effect on foraminiferal abundance and diversity (Yanko et al 1999;
375 Geslin et al. 2002). Pollution and low oxygen concentration could explain the low species BFAR
376 and diversity as well as the dissolution of tests during this period. Some species that were present,

377 i.e. the agglutinated species *E. medius/scabrus* and *R. subfusiformis*, are known to be tolerant to
378 various kind of pollution (Table 2).

379 5.3 1953 – 1998

380 The large increase in general BFAR from 1953 suggests either more favorable growth conditions
381 or significant deposition of transported specimens into the area. The coarser grain size observed
382 during this period indicates possible changes in the current system, which could affect both
383 growing conditions and transport of specimens (Figure 7). However, the dating of our core
384 showed continuous sediment accumulation without any interruption during this period (Figure 4).
385 Moreover, all the new dominating species were already present in the core, even if in lower
386 relative abundances (Figure 5A). This indicates that the BFAR increase is most likely not due to
387 specimens transport, but rather as a result of a change in substrate and environmental conditions
388 that became favorable for a different foraminiferal assemblage. The higher foraminiferal
389 diversity compared to previous periods and the decrease in the relative abundance of *S.*
390 *fusiformis* may indicate more oxic conditions. *Elphidium clavatum* has been found in coarse
391 sediment in the area (Bergsten et al. 1996), and other species that tolerate sandy environments
392 and varying TOC dominated the assemblage, such as *A. batava*, the other species in the
393 *Elphidium* species, *B. marginata*, and *E. medius/scabrus*. Furthermore, anthropogenic activities
394 such as agricultural practices were intensified during this period until the 1980s, which resulted
395 in increased nutrient loads and resulting eutrophication (i.e. Rydberg et al. 2006). The increase in
396 organic matter may have been beneficial for foraminifera as food source. Food webs and species
397 interaction like intra and inter competition might also have been modified, giving the advantage
398 to some species such as the *Elphidium* species to develop in these new environmental conditions.

399 The temporal coincidence with the shifts seen in the sediment record and the anomalous wind
400 conditions suggests a notable change of the currents through the Öresund (Figures 8, 9). The
401 simulated currents through the Öresund confirm such an abrupt change characterized by a shift
402 from very limited outflow from the Baltic to the Kattegat before ~1960 to more than a decade of
403 high relative outflow (high VAV) from the Öresund to the Kattegat and high current velocities
404 (Figure 8). While the simulation only covers the period after 1950, the analysis of wind
405 conditions and the NAO index suggest that the anomalies in the current and sediment pattern
406 from ~mid 1950's might have been unprecedented since at least the middle of the 19th century
407 (Figure 9). The shift in local sediment properties and the shift to higher BFAR and species
408 diversity suggest a combination of anomalous currents during a period of unusually negative
409 NAO index and the abrupt first advection of anthropogenic eutrophication from the Baltic Sea
410 towards the Kattegat. Consistent with our findings, long-term variations in Large Volume
411 Changes in the Baltic Sea (LVC, Lehmann and Post 2015; Lehmann et al. 2017), which are
412 calculated from >29 cm (~100 km³) daily sea-level changes at Landsort (58.74° N; 17.87° E) for
413 1887–2015, show an unusual cluster of both, more frequent and also larger LVCs during the
414 1970's to 1980's relative to the entire time period. Notably, this period coincides with the most
415 dramatic shift in foraminiferal BFAR and species diversity as well as an increase in sand content.
416 The period before the “regime shift” of the 1950's to 1960's is dominated by very infrequent and
417 few large LVC events. After the shift, the 1990's show also very few or partly no LVC events
418 with generally record-low Major Baltic Inflow events.

419 Thus, during this period, the ecosystems were affected both by climatic effects through
420 sedimentation changes, and human impact. At the end of the period, after ~1980, the general
421 BFAR was lower during a short time (Figures 5, 9). This could be linked to the measures that

422 were taken in agriculture and water treatments in order to reduce the nutrients discharge
423 (Carstensen et al. 2006; Conley et al. 2007), which could have reduced the food input.
424 Interestingly, when the sedimentation pattern changes again and the sand content decreases
425 markedly (Figure 7), the new species in the foraminiferal fauna do not return to previous relative
426 abundances as one could have expected (Figure 5A). This suggests that once the foraminiferal
427 fauna was established in the Öresund area after the ~1953 shift, it created a new state of
428 equilibrium.

429 5.4 1998 – 2009

430 The foraminiferal assemblage in this zone was similar to the previous one, with high BFAR, high
431 diversity, and the *Elphidium* species as dominating species. This period is, however,
432 characterized by the appearance of two new major species: *N. turgida* and *Nonionella* sp. T1.
433 *Nonionella* sp. T1 is suggested to be an invasive species in the region which arrived by ship
434 ballast tanks around 1985, and rapidly expanded to the Kattegat and Öresund (Polovodova
435 Asteman and Schönfeld 2015). According to our dated core, the species arrived in the Öresund
436 ~2000 CE (Figure 5). The species is also present on the south coast of Norway since ~2009
437 (Deldicq et al. 2019), but additional genetic analyses are necessary to have a better overview of
438 the species' origin and expansion. *Nonionoides turgida* is an opportunistic species that prefers
439 high levels of organic matter in the sediment, as observed in our core during this period (Figure
440 7). The increase in the *S. fusiformis* BFAR suggest lower [O₂] than in the previous zone, which
441 was indeed a general trend in the Danish waters during this time period (Conley et al. 2007). The
442 salinity was probably marine during this period, as suggested by the high occurrence of the
443 marine species *B. marginata* (Figure 5). This period was then characterized by low [O₂], high
444 organic matter content, and open ocean salinity.

445 5.5 2009 – 2013

446 The ability of *Nonionella* sp. T1 to denitrify and its tolerance to varying environment may
447 explain its rapid increase during this period. The increase of *N. turgida* also suggests higher
448 levels of organic matter in the sediment. The dominance of these two species and the lower
449 BFAR compared to previous periods suggest low oxygen levels. This period is thus characterized
450 by low [O₂], high organic matter content, and open ocean salinity.

451 5.6 Dissolution

452 The inner organic linings of the taxon *Ammonia* were observed (in low numbers, < 5 units) along
453 the whole core, except in the top two centimeters (Figure 5). Inner organic linings of the taxa
454 *Ammonia* and/or *Elphidium* were noticed in previous studies among dead fauna in the region
455 (Jarke 1961; Hermelin 1987: Baltic Sea; Christiansen et al. 1996; Murray and Alve 1999:
456 Kattegat and Skagerrak; Filipsson and Nordberg 2004b: Koljö Fjord). Dissolution of calcareous
457 foraminiferal tests has been considered as a taphonomic process, affecting the test of the
458 specimens after their death (Martin 1999; Berkeley et al. 2007). However, living decalcified
459 foraminifera have been observed in their natural environment in the south Baltic Sea (Charrieau
460 et al. 2018) and the Arcachon Bay, France (Cesbron et al. 2016) and, proving that test dissolution
461 can also occur while the specimens live. In any case, low pH and low calcium carbonate
462 saturation are suggested as involved in the observed dissolution (Jarke 1961; Christiansen et al.
463 1996; Murray and Alve 1999; Cesbron et al. 2016; Charrieau et al. 2018). Test dissolution may
464 occur in all calcitic species, but only the organic linings of *Ammonia* were found in our study,
465 probably because these were more robust to physical stress such as abrasion.

466 6 – Conclusion

467 In this study, we described an environmental record from the Öresund, based on benthic
468 foraminifera – and geochemical data and we link the results with reconstructed wind data, NAO
469 index and currents from a hydrodynamic model. Five foraminiferal zones were differentiated and
470 associated with environmental changes in terms of salinity, [O₂], and organic matter content. The
471 main event is a major shift in the foraminiferal assemblage ~1950, when the BFAR massively
472 increased and *S. fusiformis* stopped dominating the assemblage. This period also corresponds to
473 an increase in grain-size, resulting in a higher sand content. The grain-size distribution suggests
474 changes in the current velocities which are confirmed by simulated current velocity through the
475 Öresund. Human activities through increased eutrophication also influenced the foraminiferal
476 fauna changes during this period. Organic linings of *Ammonia* were observed throughout the
477 core, probably linked to low pH and calcium carbonate saturation, affecting test preservation.

478 The long-term reconstruction of sediment – and ecosystem parameters since ~1807 suggests that
479 the onset of increased anthropogenic eutrophication of the eastern Kattegat started with an abrupt
480 shift ~1960 during a period of strongly negative NAO index. With unusually calm wind
481 conditions during the winter half and increased easterly winds, the conditions were ideal for
482 larger Baltic outflow events which is a prerequisite for more frequent and stronger major Baltic
483 inflow events (Lehmann et al. 2017), as calculated from LVC events during this period. Our
484 high-resolution sediment record points towards the importance of considering also large Baltic
485 outflow events for the Kattegat environment. Since the Baltic Sea is much more eutrophic, less
486 oxygenated and less saline, large outflow events may have a significant impact also on the
487 Kattegat ecosystem. Periods with a negative NAO or conditions with intense atmospheric
488 blocking over Scandinavia like in 2018 may also increase the influence of Baltic Sea's
489 environmental problems into the Kattegat region.

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499 Supplementary data

500 Appendix A, with time series of salinity, temperature and dissolved oxygen concentration at the
501 bottom water of the Öresund, and Appendix B, with total foraminiferal faunas normalized to 50
502 cm³ along the DV core, are available in the online version of the article.

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714 Figures

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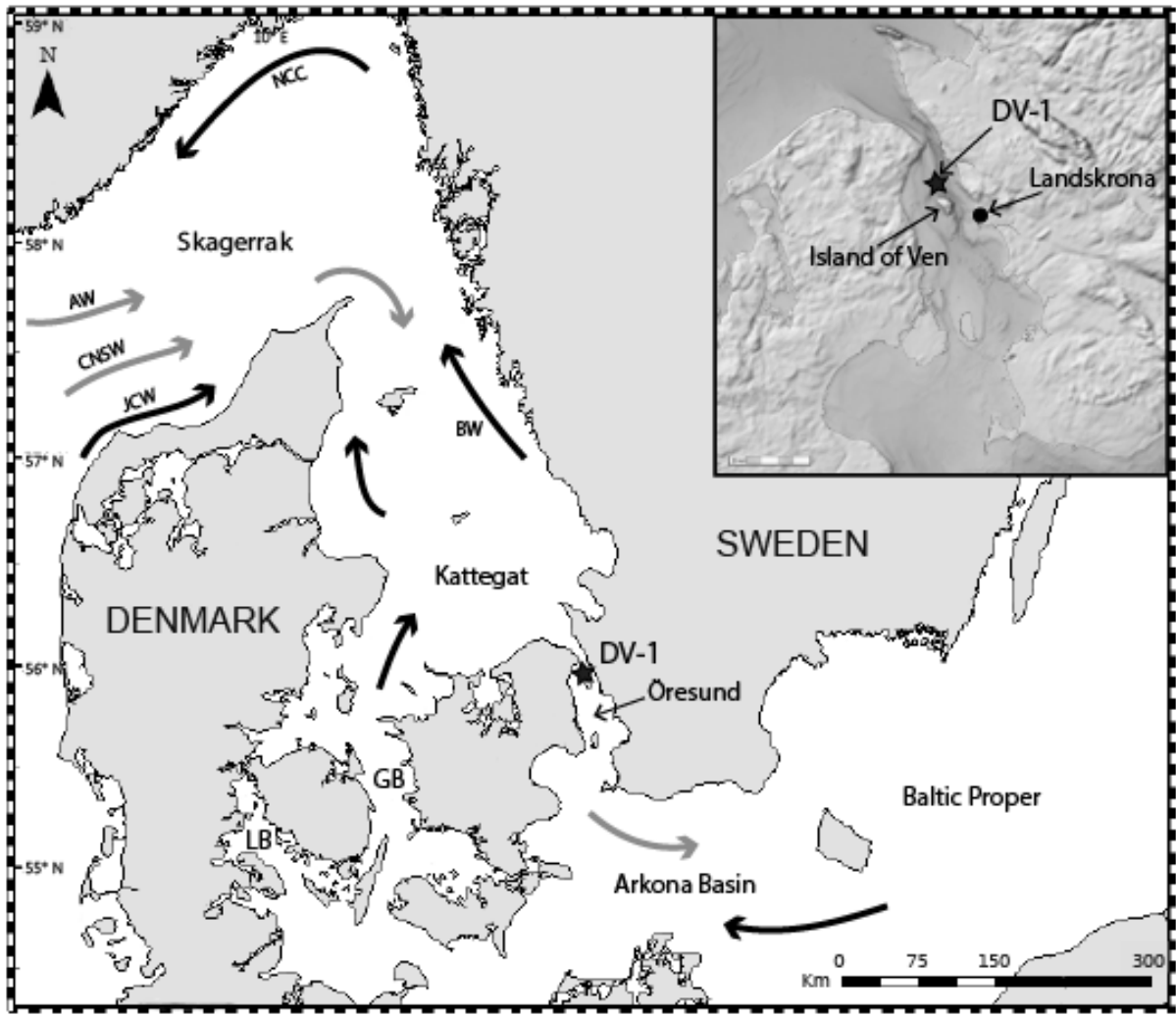
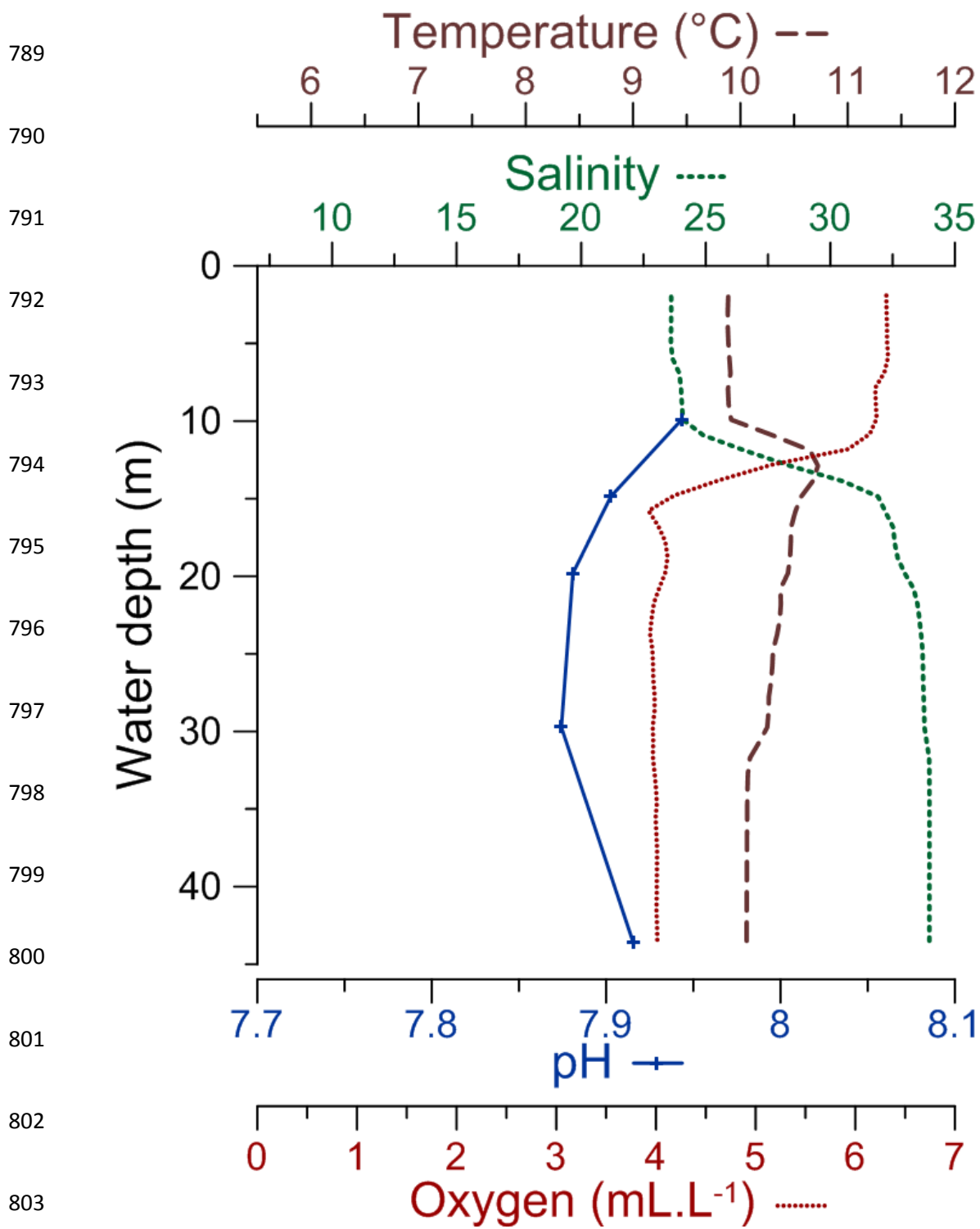
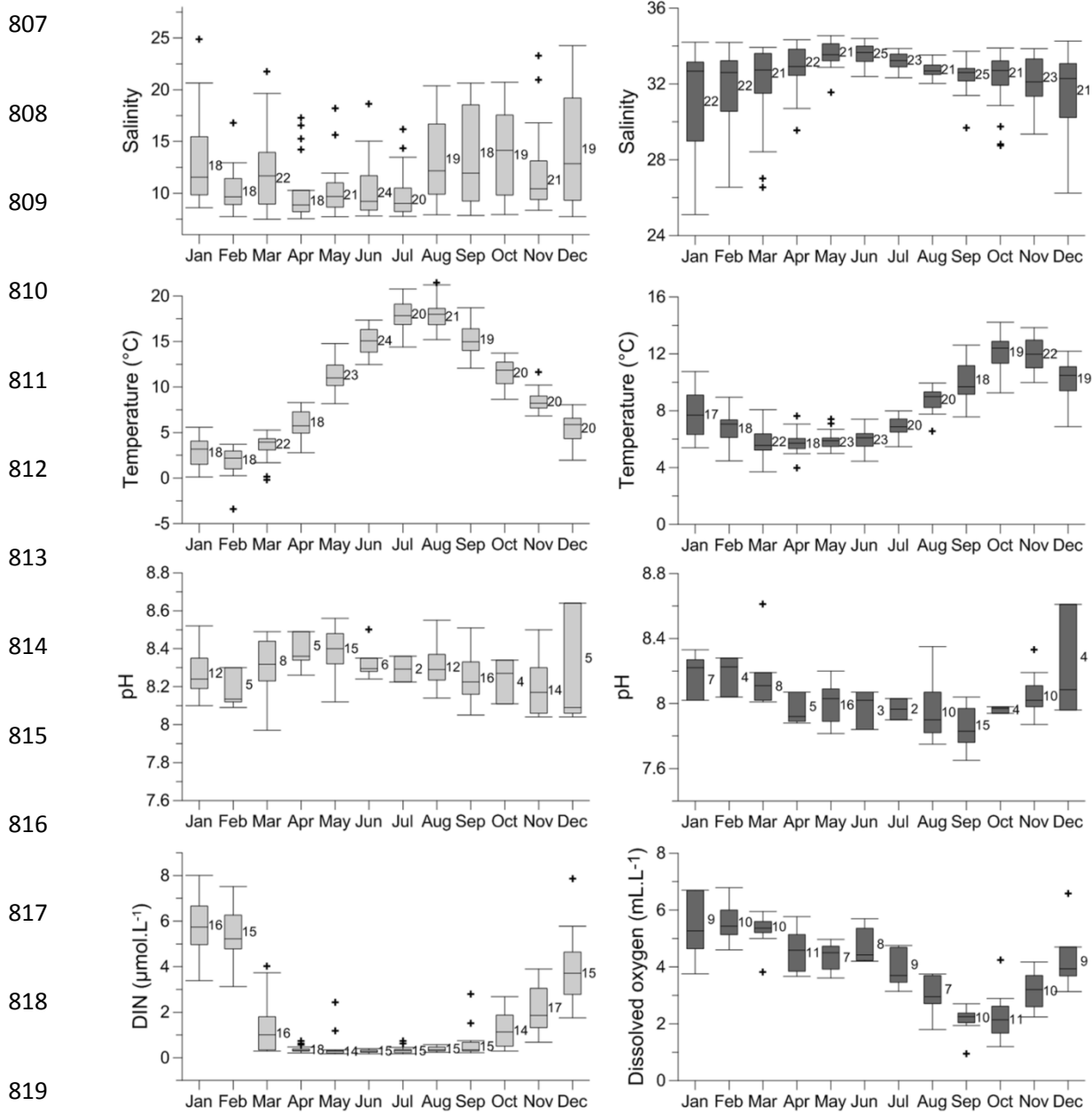


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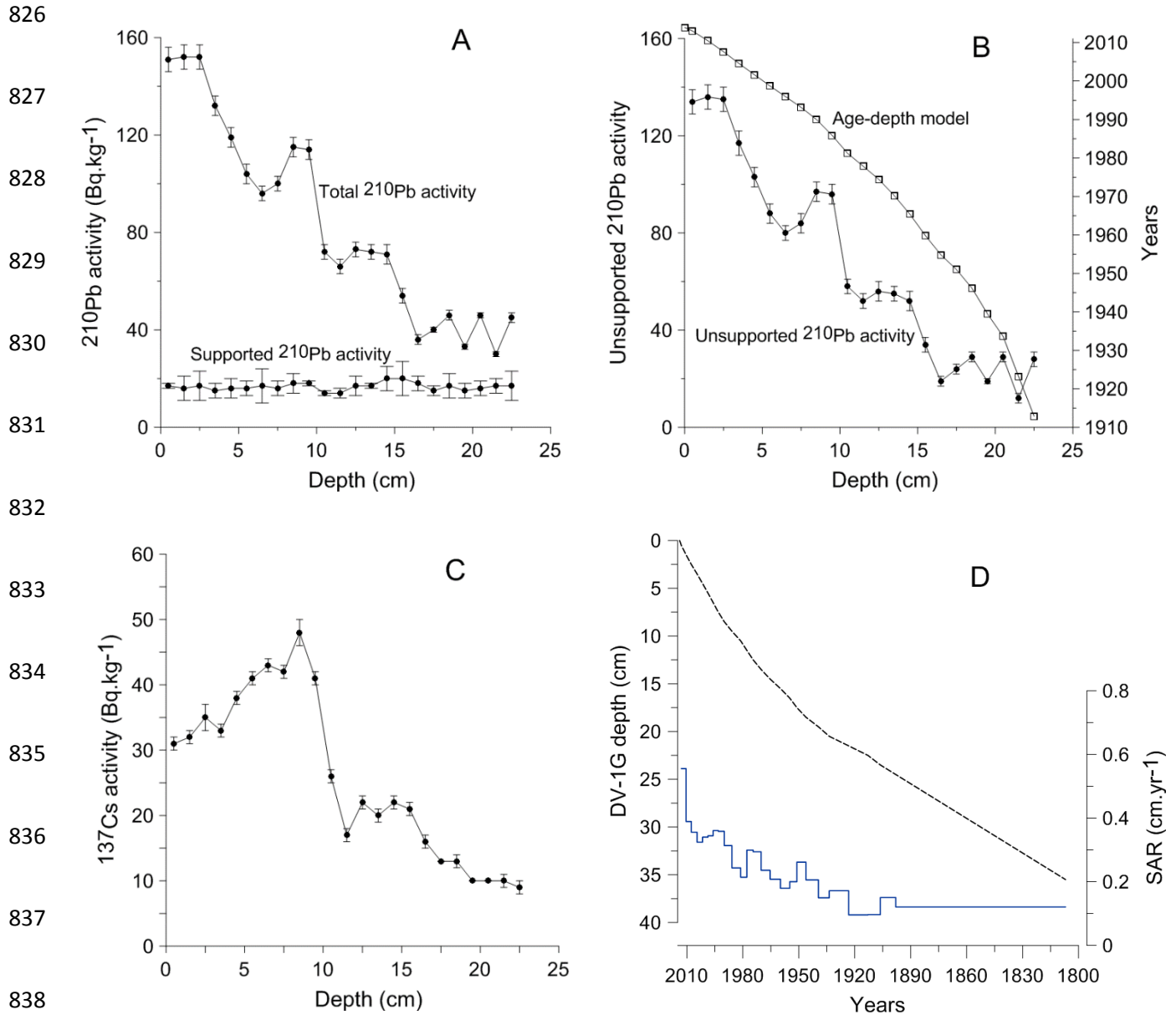


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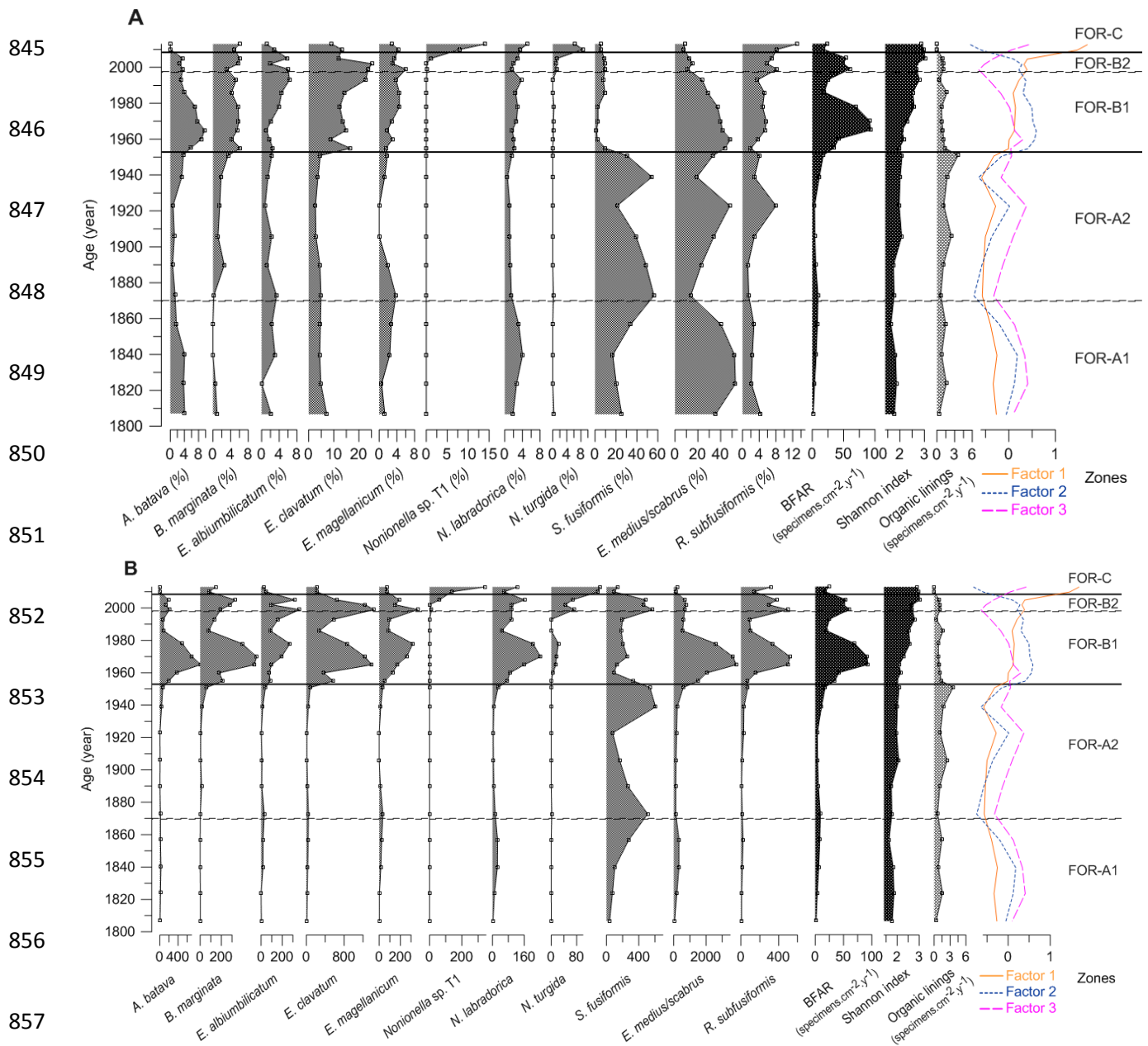


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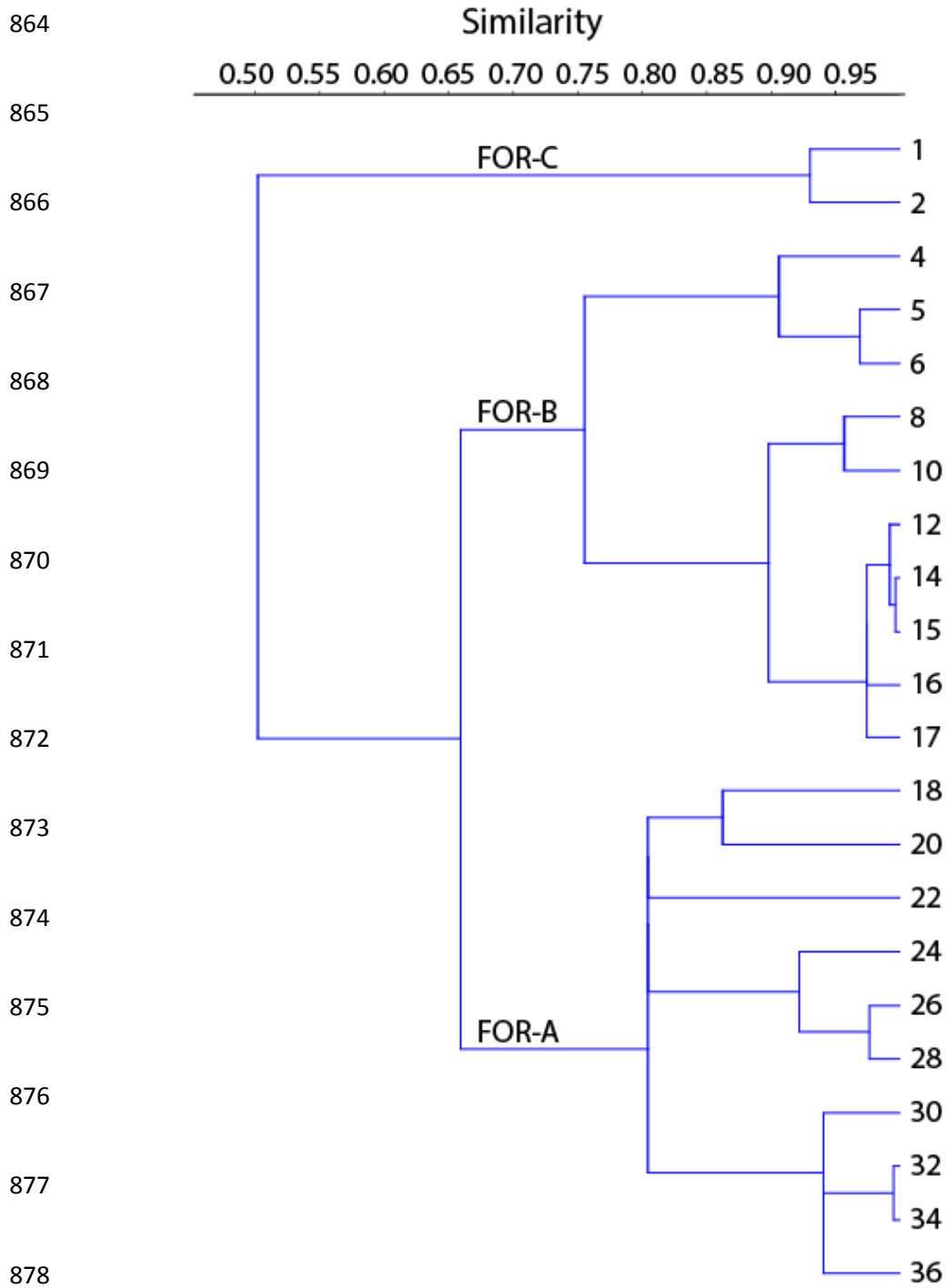


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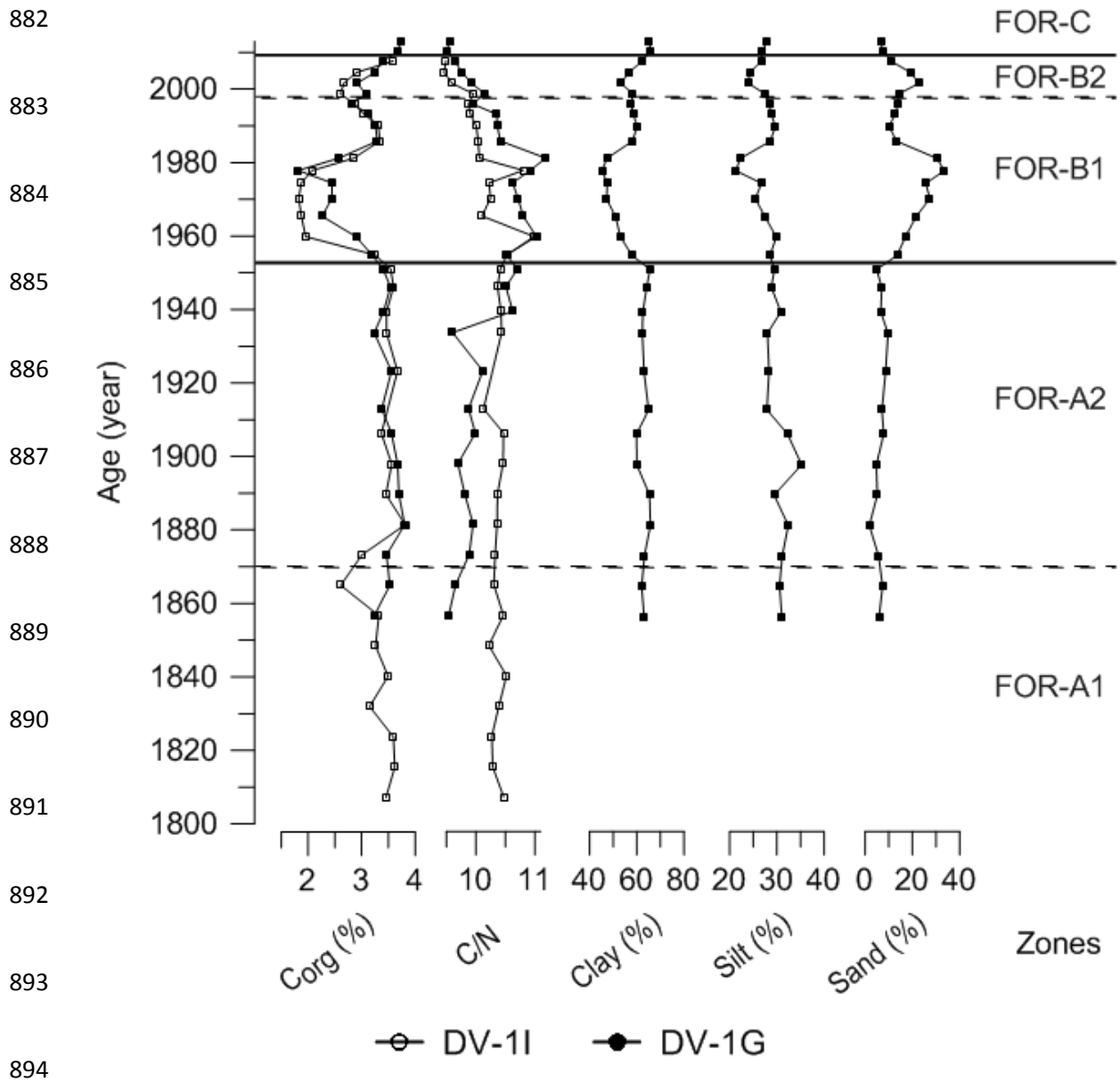


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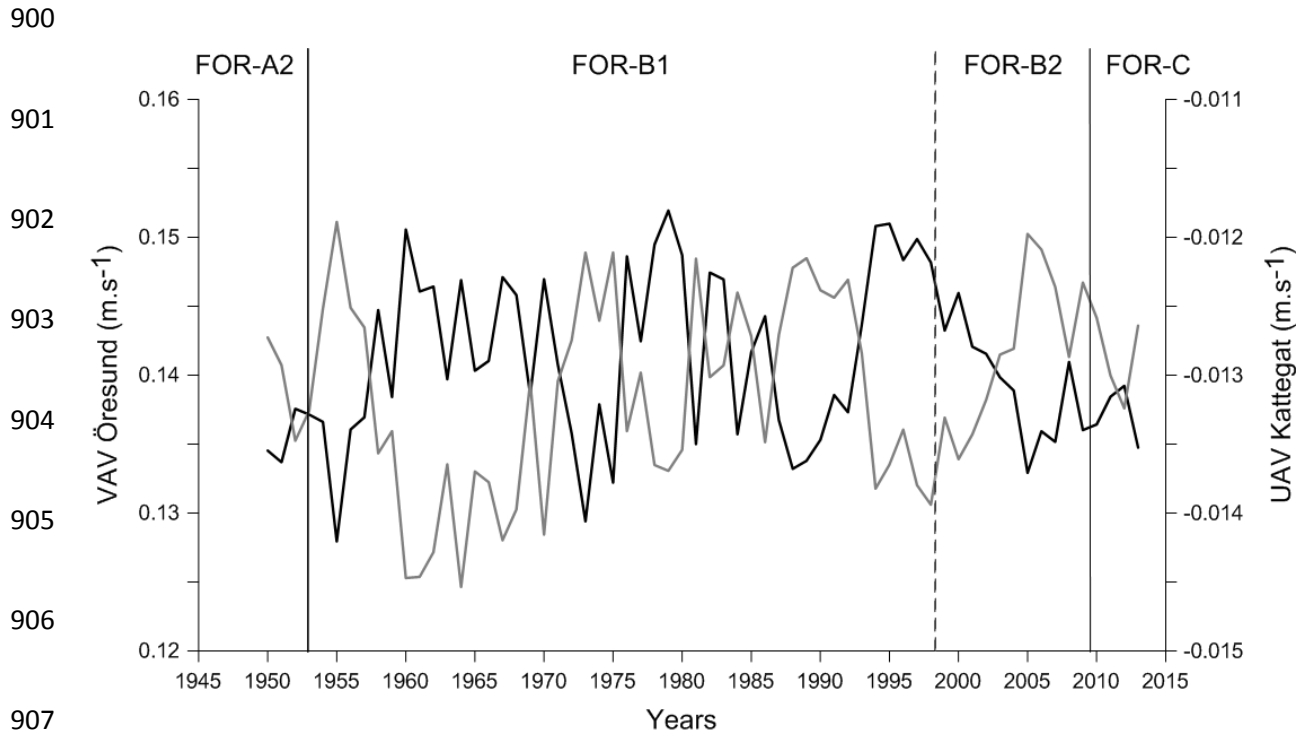
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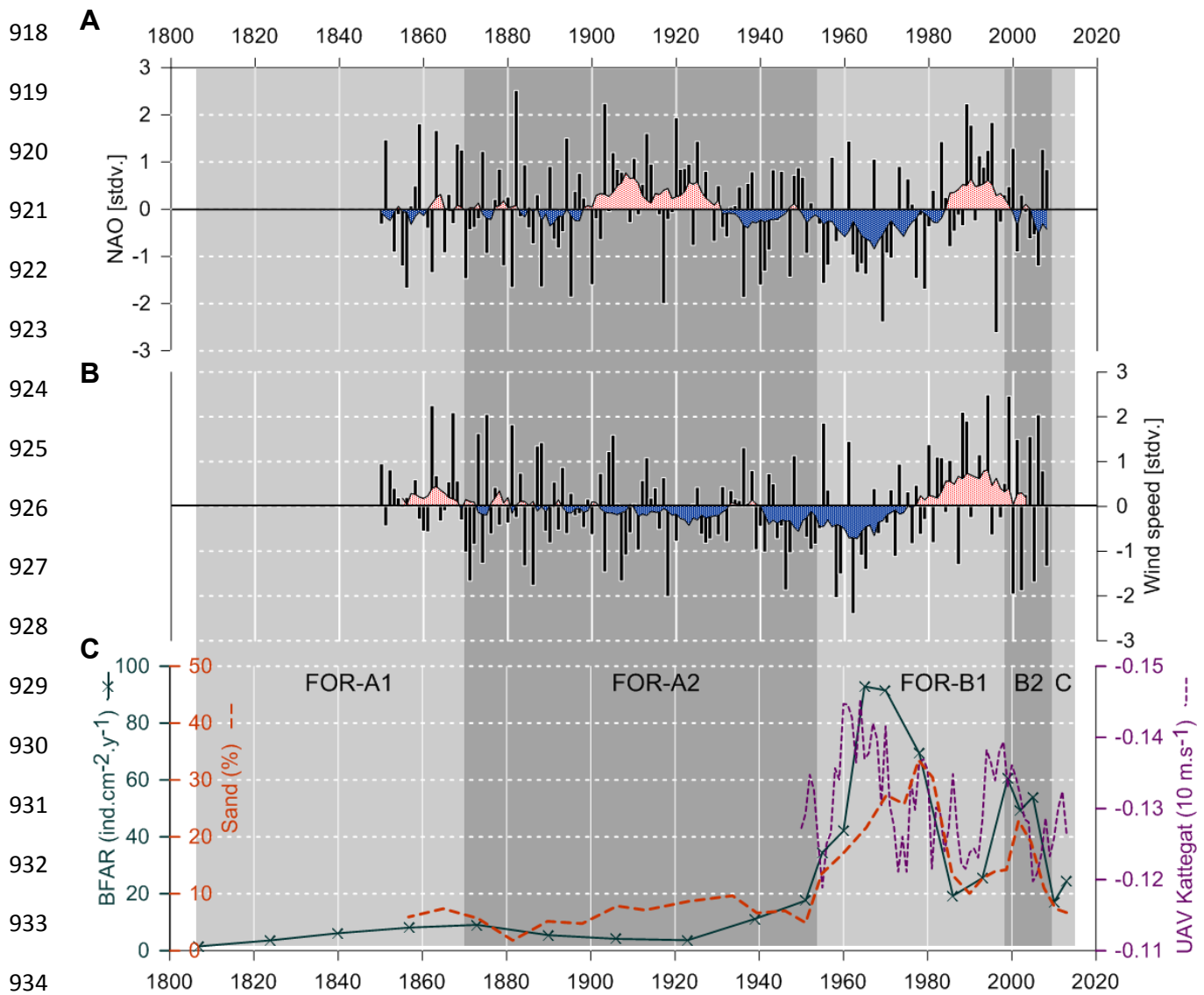
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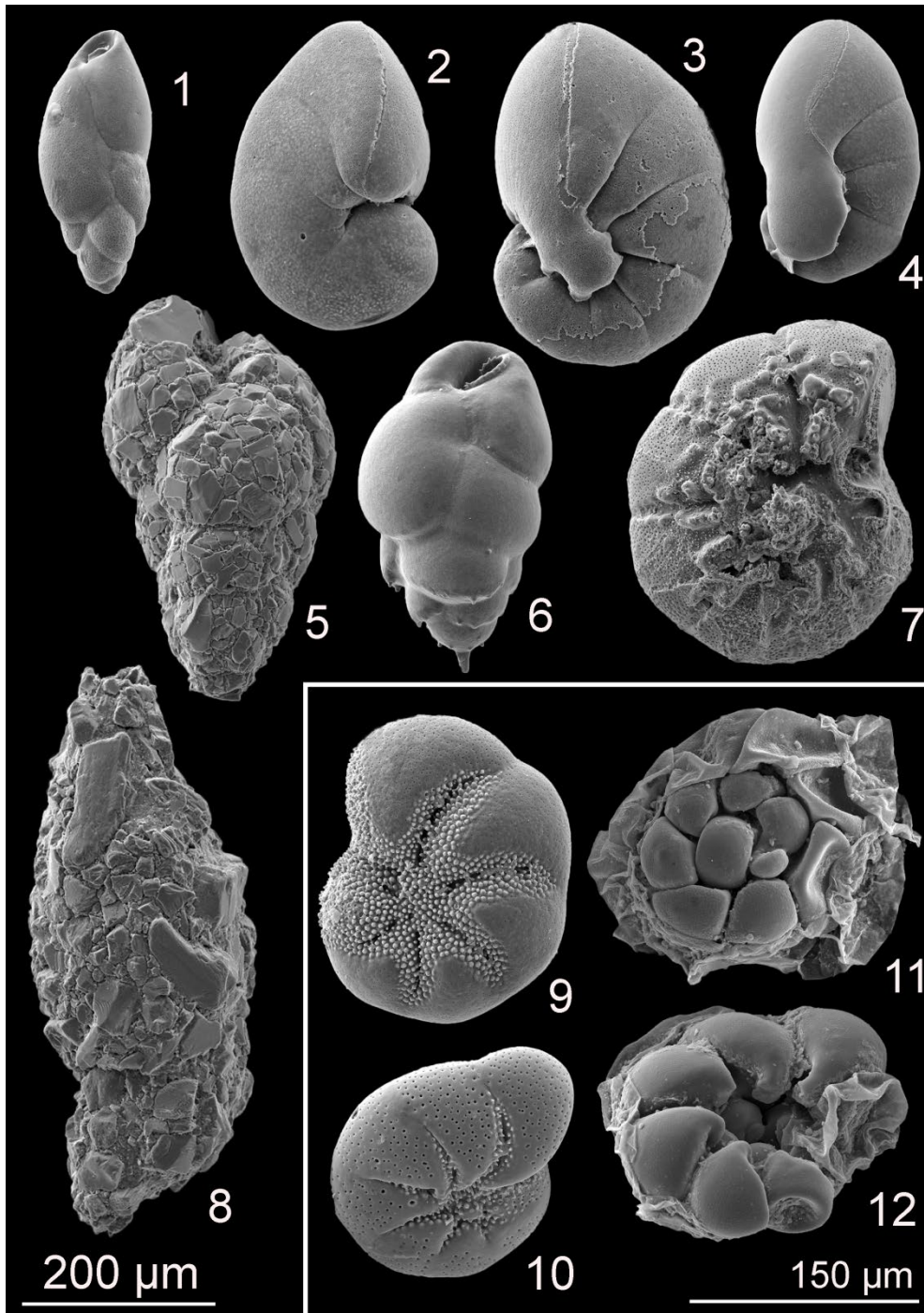


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Factor	Total variance (%)	Significant species	Score
1	48.18	<i>Nonionella</i> sp. T1	5.10
		<i>Nonionoides turgida</i>	4.14
2	30.88	<i>Ammonia batava</i>	1.34
		<i>Stainforthia fusiformis</i>	-1.41
3	13.36	<i>Elphidium albiumbilicatum</i>	-1.65
		<i>Elphidium clavatum</i>	-1.57
		<i>Elphidium magellanicum</i>	-1.32

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975 Table 2. Ecological significance of the benthic foraminiferal assemblages (major species).

Species	Ecological significance	Reference
<i>Ammonia batava</i>	Salinity 15-35, T 0-29 ⁰ C, high tolerance to varying substrate and TOC	Alve and Murray (1999); Murray (2006)
<i>Bulimina marginata</i>	Tolerates low oxygen conditions, salinity 30-35, T 5-13 ⁰ C, muddy sand, prefers organic rich substrates	Conradsen (1993); Murray (2006)
<i>Elphidium albiumbilicatum</i>	Salinity 16-26, typical brackish species	Alve and Murray (1999)
<i>Elphidium clavatum</i>	Tolerates low oxygen conditions, salinity 10-35, T 0-7 ⁰ C, high tolerance to varying substrate and TOC, subtidal	Conradsen (1993); Alve and Murray (1999); Murray (2006)
<i>Elphidium magellanicum</i>	Coastal species	Sen Gupta (1999)
<i>Nonionella stella</i> /aff. <i>stella</i>	Tolerates low oxygen conditions, kleptoplastidy, able of denitrification, invasive in the Skagerrak-Kattegat	Piña-Ochoa et al. (2010); Bernhard et al. (2012); Charrieau et al. (2018)
<i>Nonionellina labradorica</i>	Salinity >30, T 4-14 ⁰ C, high latitudes, kleptoplastidy, able of denitrification	Cedhagen (1991)
<i>Nonionoides turgida</i>	Opportunistic species, tolerates low oxygen conditions, prefers high food availability	Van der Zwaan and Jorissen (1991)
<i>Stainforthia fusiformis</i>	Opportunistic species, tolerates very low oxygen conditions, salinity >30, able of denitrification, prefers organic rich substrates, fast reproduction cycle	Alve (1994); Filipsson and Nordberg (2004); Piña-Ochoa et al. (2010)
<i>Eggerelloides medius/scabrus</i>	High tolerance to hypoxia, salinity 20-35, T 8-14 ⁰ C, sandy-muddy sand, tolerance to various kind of pollution	Alve and Murray (1999); Alve (1990); Murray (2006); Cesbron et al. (2016)
<i>Reophax subfusiformis</i>	Tolerance to environmental variations and fuel ash	Sen Gupta (1999)

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