| 1  | Repercussions of the Great East Japan Earthquake tsunami on ellipsoidal  |
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| 2  | Alexandrium cysts (Dinophyceae) in Ofunato Bay, Japan  |
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| 4  | Kazumi Matsuoka <sup>1*</sup> , Yuri Ikeda <sup>2</sup> , Shinnosuke Kaga <sup>3</sup> , Yoshimasa Kaga <sup>4</sup> , |
| 5  | Takehiko Ogata <sup>2</sup>  |
| 6  |  |
| 7  | 1: C/O Institute for East China Sea Research, Nagasaki University: 1551-7 Taira-machi,                                 |
| 8  | Nagasaki, 851-2213, Japan  |
| 9  | 2: School of Marine Biosciences, Kitasato University: 1-15-1, Kitasato, Minami-ku, Sagamihara,                         |
| 10 | Kanagawa 252-0373, Japan;  |
| 11 | 3: Iwate Prefectural Government: 10-1, Uchimaru, Morioka, Iwate, 020-8570, Japan                                       |
| 12 | 4: Iwate Fisheries Technology Center: 3-75-3, Hirata, Kamaishi, Iwate 026 0001, Japan                                  |
| 13 |  |
| 14 | 1*: Corresponding author; e-mail kazu-mtk@nagasaki-u.ac.jp,  |
| 15 | Present address; 1403, 370-2 Kuzetakada-cho, Minami-ku, Kyoto, 601-8211, Japan   |
| 16 | Tel: +81-75-932-2062   |
| 17 | <u>2: e-mail yikeda@kitasato-u.ac.jp</u>   |
| 18 | 3: e-mail s-kaga@pref.iwate.jp   |
| 19 | 4: e-mail y-kaga@pref.iwate.jp   |
| 20 | 5: e-mail t.ogata@kitasato-u.ac.jp   |
| 21 |  |
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| 23 | shellfish poisoning, Chilean tsunami   |
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| 25 | Highlight  |

- 1. The ca. upper 25cm sediments of Ofunato Bay were re-deposited by the tsunami of the 2011
  Great East Japan Earthquake.
- 28 2. The maximum density of ellipsoidal *Alexandrium* cysts at the 6–7cm depth was deu to the
- 29 re-suspension of older sediments.
- 30 3. The first PSP incident on 1961 in this bay was resulted from the similar phenomena after the
- 31 Chilean tsunami in 1960.
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#### 34 Abstract

35Shellfish aquaculture in Ofunato Bay, Northeast Japan, was seriously damaged by a tsunami generated by the Great East Japan Earthquake, March 11th 2011, accompanied 36 37 by paralytic shellfish poisoning (PSP) outbreaks caused by Alexandrium tamarense 38 (Dinophyceae). To understand longer future trends of PSP, an investigation was made of 39the historical occurrence and causes of Alexandrium outbreaks after the tsunami. 40 Vertical distributions of Alexandrium cysts in two sediment-cores from Ofunato Bay 41 revealed that the sediments above ca. 25 cm were eroded, re-suspended and 42re-deposited, and they included unusually abundant Alexandrium cysts. This abundance 43of cysts was due to re-deposition of older sediments by the tsunami. The first Ofunato 44Bay PSP incident was in 1961 after the Chilean Earthquake tsunami and was probably caused by similar unusual blooms of Alexandrium germinated from older sediments as 4546 the Great East Japan tsunami, together with nutrient enrichment because of population 47increase at the start of shellfish aquaculture.

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#### 50 Introduction

51Under usual conditions, an ecosystem is in dynamic equilibrium but it may frequently be 52disturbed by unexpected natural and/or artificial events. Consequently, it could be changed to a 53new ecosystem both positively and negatively. Events which can alter marine ecosystems are 54change of water temperature, salinity, light intensity, and water currents. In addition, unusual 55phenomena such as a typhoon/hurricane, extraordinary high or low temperature change can 56 strongly affect the marine ecosystem (eg. Reinolds 2006). A tsunami (high sea wave) caused by 57an earthquake, submarine landslide, or other disturbance is another unusual event that produces 58exceptional conditions. For example, the tsunami following the 2004 Indian Ocean Sumatran 59earthquake resulted in severe environmental impacts, mainly to the coastal zone ecosystems, 60 including mangroves, coastal wetlands, coral reefs, shallow sea floor, and shoreline habitats (eg. 61 Titov et al. 2005, "Impact of Tsunamis on Ecosystems." UN Atlas of the Oceans. Retrieved 10 62 March 2005), and it also changed phytoplankton production (Tang et al. 2009), and a benthic 63 ecosystem (Whanpetch et al. 2010).

The Great East Japan Earthquake on March 11th of 2011 (magnitude 9.0-9.1 Mw, offshore 64 65 of NE Japan) is considered one of the largest earthquakes ever recorded. It generated a wave 66 10m high that swept inland about 10 km, impacting the Fukushima nuclear plant and many 67 coastal communities (https://www.britannica.com/event/Japan-earthquake-and-tsunami-of-68 2011). Many subsequent changes of coastal marine ecosystem and related environmental 69 parameters have been reported previously (eg. Fukuda et al. 2015, Yamada et al. 2015, Okamura 70& Kaga 2016, Naiki et al. 2015, Okamura et al. 2015, Yamamoto et al. 2012, Yamada 2012, 71National Institute for Land and Infrastructure Management, and Building Research Institute 722011). Special interest focused on outbreaks of paralytic shellfish poisoning (PSP) and intensive 73studies were made of the distributions of causative dinoflagellate cysts such as Alexandrium 74tamarense/catenella (Kamiyama et al., 2014, Nishitani et al. 2012, Natsuike et al. 2014, 75Ishikawa et al. 2015).

76These special PSP studies revealed a remarkable increase of Alexandrium 77tamarense/catenella cysts after the tsunami at the Great East Japan Earthquake site, probably 78due to re-suspension of cysts accompanied with bottom sediments and newly introduced by 79plankton blooms after the tsunami. However, no investigation of this subject was made in 80 Ofunato Bay where shellfish aquacultures were active before the Great East Japan Earthquake 81 and where the local fishermen needed information about the future trends of the PSP causative 82 plankton. Additionally, the PSP phenomena in Ofunato Bay are notable in Japanese studies 83 because the first scientific investigation of a PSP event was conducted in Ofunato Bay in 1961 84 (Kawabata et al. 1962), 13 years after a suspected PSP event occurred around Lake Hamana in 85 1948 (Imai & Itakura 2002). However, the causative plankton at Lake Hamana was not known. 86 The PSP causative plankton of Ofunato Bay was identified using preserved samples (Murano 87 1975) in addition to the characteristic symptoms of the PSP (Kawabata et al. 1962). After 1961, 88 the PSP event expanded throughout Japan, persisting until now.

The 1961 PSP event in Ofunato Bay happened just one year after the Chilean tsunami that impacted the Sanriku coast of Northeast Japan in 1960. For many years, it was thought that the Chilean tsunami might be related to the first PSP event in Ofunato Bay but there were no data to support or interpret this hypothesis. New PSP studies of the 2011 tsunami after the Great East Japan Earthquake, which had a similar impact on the coastal environments and marine ecosystems in Ofunato Bay, might now provide a scientific explanation for the PSP events following the Chilean tsunami.

In this study, we investigate the vertical distribution of the cysts of the PSP causative plankton, *Alexandrium tamarense/catenella* and their depositional mode in two sediment cores from Ofunato Bay which contain all the re-deposited layers from the 2011 tsunami of the Great East Japan Earthquake. We also present the basic information on a trend of *Alexandrium tamarense/catenella* thereafter and also to provide a scientific scenario that can reasonably explain the first PSP event in Ofunato Bay following the 1960 Chilean tsunami. 102

#### 103 Environment of Ofunato Bay

104 Ofunato Bay is located at the Pacific coast of Northeast Japan and is one of several bays that 105 characterize ria coasts (Administrator of Kamaishi Port, 1976). This bay is approximately 6 km long from north to south, 2 km wide from east to west, 7.7km<sup>2</sup> in area, 1.24 x 10<sup>8</sup> km<sup>3</sup> in volume, 106 107 40m in maximum depth near the bay mouth, and it has an "L shaped" topography that opens in 108 an east-southeast direction. The Sakari River flows into the north end of this bay. A tsunami 109 break water was started to work in 1963 and built in 1967 after the Chilean tsunami. However, 110 this structure was destroyed by the Great East Japan Earthquake tsunami. Subsequently, a new 111 breakwater is being constructed again at the same site.

In the bay, shellfish aquacultures mainly dealing with scallops and oysters have been developed, both before and after the Great East Japan Earthquake. These aquacultures are one of the important industries around Ofunato Bay. Because of these circumstances, local fishermen are always concerned about PSP phenomena and they need information on the causative plankton and cysts which are called *Alexandrium tamarense* (Lebour) Balech and *A. catenella* Whedon & Kofoid in this region, as explained in the next section.

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#### 119 History of *Alexandrium* blooms and paralytic shellfish poisoning in Ofunato Bay

On May 18<sup>th</sup> of 1961 in Ofunato Bay, a PSP incident happened that was caused by 120 121Akazara-scallops (Chlamys farreri akazara (Kuroda, 1932)), with twenty people becoming 122severely ill and one death (Kawabata et al. 1962). More than ten years later, the causative 123 organism was identified as Gonyaulax sp. which seemed to be similar to Gonyaulax catenella 124Whedon & Kofoid by Murano (1975). Thereafter, Fukuyo (1979) corrected its species 125identification as Gonyaulax excavata (Braarud) Balech and then, as now, as Alexandrium 126 tamarense (Lebour) Balech via Protogonyaulax tamarense (Lebour) Taylor. Furthermore, 127recent progress in the taxonomy of the A. tamarense complex suggests that A. tamarense and A.

*catenella*, forming resting cysts and appearing in Ofunato Bay, should be called as *A. fundyense*Balech and *A. pacificum* Litaker, respectively, if we follow John et al. (2014). However, we use
the names *A. tamarense* and *A. catenella* to avoiding taxonomic confusion in this article. At
present, *A. tamarense* and *A. catenella* are the two major PSP causative dinoflagellates in
Ofunato Bay. *A. tamarense* frequently blooms mainly in spring to early summer and has high
PSP concentrations. On the other hand, *A. catenella* appears in late summer to early autumn
with relatively low PSP concentrations (Kodama et al. 1982, Ogata et al. 1982).

135 According to Kaga et al. (2012) the history of toxin concentration in shellfish from Ofunato 136Bay is as follows. In the early 1960's, toxin level in the Akazara-scallop and/or Yesso giant 137 scallop (Pecten yessoensis Jay 1857) was around 100 to 1000 MU/g digestive gland (DG). In 138 the late 1960's to the early 1970's, this value dropped to 10-100 MU/g DG, and thereafter, it 139 has been between 100 and 1000, sometimes 2000 MU/g DG, with a sharp drop (less than 10 140MU/g DG) in 1985. During the huge bloom of A. tamarense in May to June of 2011 after the 141Great East Japan Earthquake tsunami in 2011, accumulation of PSP toxins through the ingestion 142of A. tamarense was studied in the blue mussel Mytilus galloprovincialis Lamarck 1819, Yesso giant scallops and other species in Ofunato Bay. On May 13th of 2013, the toxin concentration in 143 144 scallops of Ofunato Bay was recorded as 4,647 MU/g DG, which was the maximum value for 145the bay (Kaga et al. 2012, 2013).

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#### **Material and Method**

#### 148 Core Sampling sites

Two core-sampling stations were selected in Ofunato Bay (Fig. 1). One was near Shizu where the Iwate Fisheries Experimental Station and Iwate Fisheries Technology Center have continued the PSP monitoring since 1970. Approximately 800m north from this site (St. O in Fig. 1), Port and Airport Research Institute (2001) collected a core sample and measured the sedimentation rate using the <sup>210</sup>Pb and <sup>137</sup>Cs methods before the Great East Japan Earthquake. After the 154 tsunami, the Kamaishi Office of Tohoku Regional Development Bureau (2012) collected a 155 sediment core at the same station of St. O and named that core as "P". The other station was 156 near Nagasawa where dense concentrations of ellipsoidal *Alexandrium* cysts were found in a 157 previous investigation by Iwate Prefectural Fisheries Experimental Station (Iwate Prefectural 158 Fisheries Experimental Station 2013).

Two core samples were taken by divers using a transparent acrylic pipe with 6 cm diameter. A 64cm-long core was collected near Shizu station (N 39°02.335", E 141°43.935", water depth 32m) on October 23 of 2012 and a 76cm-long core was taken near Nagasawa (N 39°02.689", E 141°43.594", water depth 21m) on August 27 of 2013. Both cores were kept vertically in dark conditions, transported to a laboratory of Iwate Fisheries Technology Center and preserved in a refrigerator of 4°C before observation and sub-sampling for further examination on water content, grain size and palynomorphs including dinoflagellate cyst extraction.

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#### 167 Sub-sampling of the cores

After observation of the sediment facies, the cores kept at Iwate Fisheries Technology Center were cut at 0.5 to 1cm intervals for the top 10cms, and at 1cm intervals below the 10cm depth. Sub-samples were stored in plastic bags and kept in a refrigerator of 4 °C before palynomorph analysis, sediment grain size analysis, and measurement of water contents.

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#### 173 Particle size analysis

For sediment particle size analysis, a Laser diffraction type particle size distribution measuring apparatus (SALD – 3100, Shimadzu Co. Ltd, Kyoto, Japan) was employed. Classification of sediment grains was followed with JIS A1204: gravel (2000 $\mu$ m>), sand (75–2000 $\mu$ m), silt (5–75 $\mu$ m), clay (<5 $\mu$ m). This classification is usually employed as test method for particle size distribution of soils by the Japanese Geotechnological Siciety. Median particle size and mode values were also calculated. 180

#### 181 Water contents

For calculating water contents, the following formula in Matsuoka and Fukuyo (2000) wasprovided: 1g of wet samples was placed on a Petri dish and put in low temperature drying oven

184 (EYELA NDO-400, Tokyo Rikakikai Co. LTD, Tokyo, Japan) at 70°C for 12 hours.

185  $V = \{ (Wm - Wd) / Wm \} \times 100$ 

- 186 V: water content (%), Wm: mud in wet condition (g), Wd: mud in dry condition (g)
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#### 188 Extraction of palynomorphs including dinoflagellate cysts

189For palynomorph extraction, the analytical method of Matsuoka and Fukuyo (2000) was 190 followed: after removing calcareous and silicate grains using 30% HCl and 37% HF 191 respectively at room temperature, the remaining organic material was sieved with stainless steel 192screens of 125µm and 20µm mesh opening size. The residue trapped on the 20µm screens was 193recovered and kept in a 20ml glass bottle. A 0.5ml aliquot from the 20ml glass bottle was 194 provided for observation and palynomorphs, mainly dinoflagellate cysts, were counted under an 195 inverted optical microscope Olympus IX 50 (Olympus Co Ltd., Tokyo, Japan) with 400 196 magnification. Such observations were performed twice for every sub-sample. Observation and 197 counting of marine palynomorphs including dinoflagellate cysts were carried out at every 1cm 198 interval above the 25cm depth and every 5cm interval below this depth for both cores. All 199 marine palynomorphs remains were represented as cyst numbers per dry weight of sediment.

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#### 201 Identification of dinoflagellate cysts

Dinoflagellate cyst identification mainly followed Matsuoka & Fukuyo (2000) and Zonneveld & Pospelova (2015). Although ellipsoidal cysts with transparent walls have been known to be produced by *Alexandrium acatenella* (Whedon & Kofoid) Balech, *A. catenella* and *A. tamarense*, it is mostly impossible for differentiating these cysts as species based on only cyst morphology. However, cyst of *A. acatenella* has been recorded only one time from surface
sediment of Hiroshima Bay by Yoshida et al. (2003). Therefore, in this study we treated these
cysts as cysts of ellipsoidal *Alexandrium*, that included cysts of both *Alexandrium catenella* and *A. tamarense*. Dinoflagellate cyst concentrations including the cyst of ellipsoidal *Alexandrium*were based on cysts filled with protoplasm and of empty.

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#### Results

#### 213 Sediment

214The core taken at the Shizu station (Shizu core) was 64cm in total length. When the core sample 215was collected (Fig. 2A), on the surface many organisms belonging to the Ampharetidae 216 (polychaetes) inhabited. Many small burrows probably formed by these polychaetes were also 217 observed around 24.5cm depth below the surface. The sediment from the top to 2.5cm depth 218was mud of dark olive color, and from 2.5 to 24.5cm depth, it was rather soft and grey, with a 219dark olive mottled structure yielding many small shell fragments. The sediment from 6 to 9cm 220depth was sandy. Below 24.5cm depth, the sediment became slightly firm in texture. Around 22122cm depth, a thin layer consisting of small shell fragments was observed. The sediment above 222the 25cm depth became coarser than that below this depth. Between 32 and 38cm depth, the 223sediment was greenish black to greenish grey in color. Around 41cm depth, the sediment had a 224little sulfide smell of rotten eggs. Below 45cm, the sediment was rather homogeneous and 225greenish grey in color and it included scattered small shell fragments.

The core collected at the Nagasawa station was 76cm in total length and generally characterized by abundant shell fragments throughout the section (Fig. 2B). The sediment was grayish to olive black mud, interspersed with small shell fragment layers at the 6, 9 and 17cm depths. From the 16 to 20cm depth, there were lots of coarse sand grains. Many Ampharetidae polychaetes were also observed on the surface and their slender burrows intruded to nearly the 25cm depth. Below approximately the 20cm depth, the sediment became a harder, greenish dark

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grey mud with a marbled color. From 33–37cm depth, a shell fragment layer was observed.
From the 55cm depth to the bottom, the sediment became coarse and included larger shell
fragments and cobbles and barnacle shell fragments of ca. 2cm size were found around the
60cm depth.

236In both cores, fine sediment particles of silt and clay generally comprise more than 80 % of 237the weight; however, at several horizons, amounts of coarse sand grains increased. These sandy 238horizons were at the 2.5, 6, and 24cm depth in the Shizu core (Fig. 3), and at 2, 5, 13–22, 40, 45, 23955-75cm depth in the Nagasawa core (Fig. 4). Water content in the Shizu core was clearly 240divided into two parts (Fig. 3). In the upper part above 22cm, water content ranged between 60 241and 70% except for the top part, and in the lower part, it decreased below 60%. On the other 242hand, in the Nagasawa core, in the upper part above 17cm, the water content was generally 243constant around 65 %, although below 17cm depth, it decreased to less than 60% and below 24462cm depth, it was less than 40% (Fig. 4). Median particle size  $(md\phi)$  in the Shizu core was 245also divided into two parts by the 22cm depth. The upper part above 17cm was around 15 µm 246and the lower part below 23cm was around 12µm. Between17 and 23cm in the Shizu core, the 247median particle size increased to around 20µm. In the Nagasawa core, median particle size 248varied between 8 and 25µm. In particular, in the 26 to 32cm and 36 to 44cm intervals, median 249particle size was less than 10µm. The mode of the Shizu core sediments showed a similar trend 250to the median particle size and it was rather stable, generally 10 to 20µm except for the 22-23cm 251depth. On the other hand, the mode of the Nagasawa core sediment particle size was larger and 252variable (8.8-26.5µm) except for 71-72cm depth due to scattered larger shell fragments.

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### 254 Dinoflagellate cysts and other marine palynomorphs

255 Various marine palynomorphs composed of dinoflagellate cysts, acritarchs, archeomonads, 256 tintinnomorphs, microforaminiferal linings, and crustacean remains were abundantly preserved 257 in these two core sediments. Among these palynomorphs, dinoflagellate cysts were most abundant and composed of 3 gymnodinioid, 11 gonyaulacoid, and 27 peridinioid species.
Among them, there were 13 photo/mixotrophic species and 28 heterotrophic species. Important
dinoflagellate cysts were shown in Fig. 5. Acritarchs included two species: *Baltisphaeridium* sp.
and *Halodinium major* Bujak. Three different uniserial, biserial and coiled types of
microforaminiferal linings were also observed. These palynomorphs were listed in Table 1 and
their abundance were recorded in Supplemental Tables 1 and 2.

264In the Shizu core, the most dominant dinoflagellate cyst was Alexandrium 265tamarense/catenella (ellipsoidal Alexandrium cyst), followed by Spiniferites bulloideus s.l. and 266 Brigantedinium spp. The average number of ellipsoidal Alexandrium cyst was recorded as 11.4 267 x  $10^3$  cysts/g above the 20cm depth, with a maximum of 26.3 x  $10^3$  cysts/g at 18–19cm depth 268(Fig. 6). Cysts filled with protoplasm were observed at the 50-51cm depth and empty forms 269were seen at 60–61cm depth. However, the cyst density below 26cm was lower:  $1.6 \times 10^3$  cyst/g 270on average. The density of the ellipsoidal Alexandrium cyst in sediments above 25cm depth, 271fluctuated much more than below this level. Average cyst density of the gonyaulacoid species 272was 2.2 x  $10^3$  cyst/g above the 20cm depth, and 1.5 x  $10^3$  cyst/g below this depth. Spiniferites 273elongatus and Lingulodinium machaerophorum occurred rarely throughout the core. 274Tuberculodinium vancampoae was observed sporadically, even below 20cm depth. Average 275cyst density of the peridinioid species except for organic linings of phototrophic Scrippsiella 276spp. cysts was 4.5 x  $10^3$  cyst/g average above 20cm depth, and 3.2 x  $10^3$  cyst/g below this depth. 277 Brigantedinium species, mostly B. simplex, were most abundant. Selenopemphix quanta and the 278cyst of Protoperidinium leonis constantly occurred. Votadinium spinosum, Dubridinium 279caperatum, and cyst of Protoperidinium americanum were few in number but they constantly 280occurred throughout the core. Heterotrophic gymnodinioid species, Polykrikos kofoidii and 281Polykrikos schwartzii were commonly present.

In the Nagasawa core, the most dominant dinoflagellate cyst was also the ellipsoidal *Alexandrium* cyst, followed by *Spiniferites bulloideus* s.l. and *Brigantedinium* spp. (Fig. 7). The

ellipsoidal *Alexandrium* cyst was recorded with an average of 76.8 x  $10^3$  cyst/g above 20cm depth, with a maximum of 155.21 x  $10^3$  cyst/g at 6–7cm depth; they showed also fluctuations as in the Shizu core. Cysts filled with protoplasm were found at 74cm depth, but the cyst density below 20cm depth became lower, averaging 1.12 x  $10^3$  cyst/g. Average cyst densities of the gonyaulacoid species was 4.1 x  $10^3$  cysts/g above 20cm depth, and 1.7 x  $10^3$  cysts/g below this depth. Average cyst density of the peridinioid species except for organic linings of *Scrippsiella* spp. cysts were 7.9 x  $10^3$  cysts/g above 20cm depth and 3.9 x  $10^3$  cysts/g below this depth.

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#### 292 **Discussion**

#### 293 1. Re-depositional processes reflected in sediment graded-bedding

294Based on the water contents, grain size distribution and development of worm burrows formed 295by polychaetes, the core sediments collected at Shizu and Nagasawa sites were divided into two 296parts separated by a transitional zone. In the Shizu core, the transition was seen from 20 to 297 26cm depth where sand content slightly increased. Slender worm burrows reached down to 298around the 25cm depth. In the Nagasawa core, the transitional zone was rather unclear with 299 regard to water content. However, worm burrows in this core were also continuously observed 300 to the 25cm depth. This means that the thickness of re-deposited sediments reached around the 301 25cm depth in both cores. The thickness of re-deposited sediments was nearly same of the depth 302 at Station P of the Kamaishi Office of Tohoku Regional Development Bureau (2013) which was 303 located at approximately 800m north of the Shizu core site (Data provided by Kamaishi Office 304 of Tohoku Regional Development Bureau 2013). However, the water contents around 20-25cm 305depth in both cores did not changed from the below (undisturbed) sediments. This suggested 306 that these sediments were considered as mixture parts of re-suspended and non-eroded 307 sediments. Accordingly, the surface sediments from the sea floor to near 25cm depth in the 308 central part of Ofunato Bay were re-deposited after erosion and subsequent re-suspension by the 309 tsunami of the 2011 Great East Japan Earthquake.

310 Above the transitional part of the Shizu core, graded-bedding and reverse graded-bedding 311 also appeared repeatedly at 3 to 5cm intervals. If the velocity of tsunami flow increased with the 312rising current, the sediment gradation was reversed as shown by Fujiwara et al. (2003). In the 313 Shizu core, normal and reverse gradations were observed repeatedly. Fujiwara et al. (2003) 314 showed that normal and reverse gradations in tsunami deposits were often observed and were 315 produced by different events in a single tsunami based on investigation of fossil beds in Boso 316 Peninsula of the East Japan. Such sedimentary features in the Shizu core might reflect the 317 number of surges associated with the tsunami event. The Port and Airport Research Institute 318 (2011) showed that several tsunami events (multiple surges) were recorded along the Sanriku 319 coast at the time of the Great East Japan Earthquake. Such depositional features are also 320 reflected in ellipsoidal Alexandrium cyst densities as discussed below. In fact, five tsunami 321 events at the time of the Great East Japan Earthquake were recorded off Kamaishi Bay located 322 at 26km NNE from Ofunato Bay (Port and Airport Research Institute, 2011). Since the tsunami 323 giant waves were repeated in approximately one-hour interval, re-suspended sediment particles 324 including ellipsoidal Alexandrium cysts filled with protoplasm would have started to sink to the 325 seafloor after each surge. However, before complete deposition on the sea floor, the next 326 tsunami surge disturbed the seafloor again. Such seawater motion would be reflected in 327 characteristics of the re-deposited sediments and ellipsoidal Alexandrium cysts of Ofunato Bay 328 as fluctuations in the sediment gradation and density of ellipsoidal Alexandrium cysts.

When the tsunami came into the bay, bottom sediments were eroded, and dinoflagellate cysts including ellipsoidal *Alexandrium* cysts were re-suspended and transported together with other sediment particles. After the tsunami encroachment, re-suspended particles were deposited according to their specific gravities. Since the specific gravity of fine muddy sediments of md  $\varphi$ (= ca. 2.5µm) saturated with sea-water (sludge) and the cysts of *A. tamarense* filled with protoplasm are 1.18-1.51g/cm<sup>3</sup> (Nakagawa et al. 2004) and 1.24 g/ cm<sup>3</sup> (Anderson et al. 1985) respectively, these ellipsoidal *Alexandrium* cysts generally behaved as fine muddy sediment

336 particles but slower than coarse silt particles as suggested in Kamiyama et al. (2014). Therefore, 337 these ellipsoidal Alexandrium cysts would probably sink faster than finer sediment particles 338 (fluid mud) with a specific gravity of approximately 1.11g/cm<sup>3</sup> (Nakagawa et al. 2014; 339 Kamiyama et al. 2014). However, in the Shizu and the Nagasawa cores, the vertical depositional 340 profile of ellipsoidal Alexandrium cyst concentration showed several peaks in the re-suspended 341 layer, as well as other sediment particles (Figs. 3, 4). Such vertical distributions of the 342 ellipsoidal Alexandrium cysts reflect the several different tsunami events at the Great East Japan 343 Earthquake.

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#### 345 **2.** Extremely unusual occurrences of *Alexandrium* cysts

346 Abundance of Alexandrium, other gonyaulacoid, and peridinioid cysts shows remarkable 347 changes around 20cm depth in both cores. The effect of re-suspension and re-deposition of other 348 gonyaulacoid and peridinioid cysts is nearly twice as high compared to the lower part of the 349 records (Figs. 6, 7). As discussed previously, the sediment above the 20 to 25cm depth of both 350 cores were re-suspended and re-deposited after the tsunami. Ellipsoidal Alexandrium cysts filled 351with protoplasts drastically increased in the re-deposited sediment as did the other dinoflagellate 352cysts, but the aspect of the Alexandrium cysts is remarkably different from other dinoflagellate 353cyst groups. In the Shizu core, the density of ellipsoidal *Alexandrium* cysts averages  $11.4 \times 10^3$ 354cyst/g above 20cm depth, with a maximum amount of 26.3 x  $10^3$  cyst/g at 18–19cm depth. In 355the Nagasawa core, the cyst density of Alexandrium filled with protoplasm showed a slightly different distribution with larger average of 76.8 x 10<sup>3</sup> cyst/g above 20cm depth and a maximum 356 357 of 155.21 x  $10^3$  cyst/g at 6–7cm depth.

In the first report of a PSP causative dinoflagellate called *A. tamarense* in Ofunato Bay (Fukuyo, 1980), the ellipsoidal cyst concentration maximum in surface sediments was recorded as 2,189 cysts/cm<sup>3</sup> at the center of the bay, near the Nagasawa station. Thereafter, cyst concentrations recorded annually by Iwate Prefectural Fisheries Station (1985–1989 and 1991)

362 ranged from 175 to 1200 cysts/cm<sup>3</sup> (Sekiguchi et al. 1998). Unfortunately, since these data were 363 reported in cysts per wet sediment volume, it is impossible to directly compare these 364 concentrations with our data measured as cysts per dry sediment weight. However, we can 365 convert those data cysts per wet sediment volume to dry sediment weight using the data of the 366 specific gravity of sediments provided by Kamaishi Office of Tohoku Regional Development 367 Bureau (2013). The specific density of dry sediments of the top 2cm of the core collected in the 368 center of the bay was measured as 0.289g/cm<sup>3</sup>. Using this conversion value, the cyst 369 concentrations of Fukuyo (1980) and Iwate Prefectural Fisheries Station (1985-1989, 1991) can 370 be recalculated from cysts/cm<sup>3</sup> to cysts/g of dry sediment. This re-calculation indicates that 371 from 197980 to 1991, cyst concentrations in the center of Ofunato bay are estimated to be 951 372 to 4,276 cysts/g.

373 As shown in Table 2, the highest previously recorded concentration of the Alexandrium 374 tamarense complex cysts was 220,872 cysts/g in Bedford Basin of Nova Scotia, Canada 375 (Lacasse et al. 2013). In Japan, the highest previously known concentration of ellipsoidal 376 Alexandrium cysts was 8,900 cysts/cm<sup>3</sup> in Aso Bay of Tsushima Island (Matsuoka & Lee 1994). 377 After the Great East Japan Earthquake ellipsoidal Alexandrium cysts were recorded as 8,190 378 cysts/cm<sup>3</sup> in Sendai Bay (Kamiyama et al. 2014). In comparison with these previous data, the 379 concentration of ellipsoidal Alexandrium cysts filled with protoplasm at the 6-7 cm depth of the 380 Nagasawa core (155.21 x 10<sup>3</sup>cyst/g) is the highest record presently known in Japan. Similar 381 extremely high ellipsoidal Alexandrium cyst concentrations were also recorded in several other 382bays, Kesennuma Bay (Ishikawa et al. 2014) and Funka Bay (Natsuike et al. 2014) after the 383 tsunami at the Great East Japan Earthquake.

However, two points should be carefully considered in explaining the unusual abundances of ellipsoidal *Alexandrium* cysts filled with protoplasm in the tsunami-impacted area of the Great East Japan Earthquake. The first point is the effect of sediment erosion, re-suspension, and re-deposition by the tsunami as suggested by Kamiyama et al. (2012). The second point 388 concerns the introduction of newly produced cysts during huge blooms of A. tamarense after the 389 tsunami (Natsuike et al. 2014, Ishikawa et al. 2015, Ogata et al. 2016). In the Shizu and 390 Nagasawa cores, the top 1cm sediments deposited after the huge bloom of A. tamarense 391 happened on May to June of 2011 (6,554 and 9,619cysts g<sup>-1</sup>, respectively) contained more 392 abundant ellipsoidal Alexandrium cysts compared with the period before 2011 (ca. 4,276cysts/g 393 as shown in our records). These Alexandrium cysts preserved in the top 1cm of the Shizu and 394 Nagasawa cores included ellipsoidal Alexandrium cysts newly produced during the bloom of 395 May to June of 2011. Such deposition of ellipsoidal Alexandrium cysts in the surface of 396 sediments was common to Sendai Bay, Kesennuma Bay and Funka Bay, but different from the 397 deeper parts of the Shizu and the Nagasawa cores.

398 Since higher cyst concentrations were recorded below the 2cm depth of the Shizu and 399 Nagasawa cores, there was no possibility that new ellipsoidal Alxandrium cysts produced one 400 year after the tsunami were preserved in deeper sediments. Therefore, explanation for such 401 extremely abundant occurrences of ellipsoidal Alexandrium cysts can be focused on different 402 eco-physiological and sedimentological factors: a higher cyst-forming capability of A. 403 tamarense and deeper erosion and transport of re-suspended and well-sorted sediments by the 404 tsunami. These events happened before the next bloom of Alexandrium tamarense on spring of 4052012.

406

#### 407 2-1. Higher cyst forming capability of Alexandrium tamarense

In Ofunato Bay, *Alexandrium tamarense* and *A. catenella* which can produce ellipsoidal cysts are known to be present (eg. Fukuyo 1982). *A. tamarense* usually forms blooms during the relatively lower sea surface temperature period (ca. 10°C, January to June, mainly May to June). On the other hand, *A. catenella* blooms are often observed during the relatively higher sea surface temperature period (ca. 15°C, July to December, mainly November (eg. Ogata et al. 1982). Of these two species, the cell density and toxin level of PSP in *A. tamarense* is usually 414 higher than A. catenella in Ofunato Bay (Kodama et al. 1982). Thus, A. tamarense is the more 415important causative PSP Alexandrium species in this bay. Incidence of cyst formation for A. 416 tamarense has been reported as 20% (Anderson et al. 1984), 30% (Ichimi et al. 2001), 23.8 ± 417 5.1 % at 14°C (Nagai et al. 2008) and 17% (Kaga 2010). These cyst production numbers are 418 significantly higher than for other photosynthetic gonyaulacacean dinoflagellates, including A. 419 catenella 1.2-2.1% (Matsuoka & Takeuchi 1995), Gonyaulax digitale (Pouchet) Kofoid (= 420 Spiniferites bentori (Rossignol) Wall et Dale) 5.9% (Wall & Dale 1968), Alexandrium 421*tamiyavanichii* Balech  $6.2 \pm 1.1\%$  in (Nagai et al. 2008), and a heterotrophic cyst-forming 422dinoflagellate (Protoperidinium oblongum (Aurivillius) Parke et Dodge 1.4% (Wall & Dale 4231968). This means that A. tamarense is one of most prolific cyst-producing species among the 424bloom-forming dinoflagellates.

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#### 426 2-2 Large blooms of <u>Alexandrium tamarense</u> frequently formed in the bay

427After the first species identification of A. tamarense as an important PSP causative organism in 428 Ofunato Bay, monitoring for this species has continued until now at the Shizu site by Iwate 429 Prefectural Fisheries Experimental Station and subsequent Iwate Fisheries Technology Center. 430 Before the 2011 Great East Japan Earthquake, an A. tamarense cell density of less than 10,000 431cells/L was recorded in 1979, 1983–1988, 1991–1992, 1996, 1999, 2001–2002, 2004–2007, and 4322009-2010. Over 10,000 but less than 50,000 cells/L was reported in 1980-1982, 1990, 1994, 4331997, 2000, 2003, and 2008, and over 500,00 cells /L was found in 1989, 1995 and 1998. The 434 highest known density reported was 96,200 cells/L in 1998 (Sekiguchi et al. 1996; Iwate 435Fisheries Technology Center 2013). The other PSP causative species, A. catenella which 436 usually appears in sea surface temperatures above 15°C, is reported to have lower cell densities 437 (eg. maximum of 20,300 cells/L according to Ogata et al. 1982) and amounts have never 438exceeded that of A. tamarense. Therefore, most of ellipsoidal Alexandrium cysts in Ofunato Bay 439 are probably formed by A. tamarense, although ellipsoidal cysts can be produced by both A.

440 *tamarense* and *A. catenella*. From the viewpoint of cyst-forming capability and cell density 441 during blooms, *A. catenella* seems to be lower than those of *A. tamarense*. Therefore, *A.* 442 *tamarense* is the most important and key species for producing unusually abundant cysts 443 compared with other cyst-forming species, and it also produced most of the resting cysts 444 preserved in the sediments of Ofunato Bay.

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#### 446 **2-3** Deeply eroded, well sorted, and re-deposited bottom sediments

447It is impossible to know the depositional age of Shizu and Nagasawa cores based on radioactive 448 elements such as <sup>210</sup>Pb and <sup>137</sup>Cs measurements, because the upper part of these cores were 449 disturbed by the tsunami of the Great East Japan Earthquake. However, the sediment core of 45050cm length of the P site, which was located at approximately 800m north of the Shizu site, was collected on September of 2012 and investigated for <sup>210</sup>Pb and <sup>137</sup>Cs concentrations by 451452Administrator of Kamaishi Port (2013). The upper 21cm of this core was also re-deposited by 453the several tsunamis including the Great East Japan Earthquake on 2011, Chilean tsunami on 4541960, probably Showa Sanriku tsunami on 1933 and Meiji Sanriku tsunami on 1986. However, 455based on the sedimentation rate  $(0.2 \text{ g/cm}^2/\text{y})$  in another core collected on 2000 at the same site 456(called the O site) provided by the Port and Airport Research Institute and Japan NUS (2001), the depositional age of the 21cm depth in the P core was estimated as 1885 year by 457458Administrator of Kamaishi Port (2013). The un-disturbed lower parts of all these four cores 459(Shizu, Nagasawa, P and O cores collected at the center of Ofunato Bay) were not different in 460 sediment facies, having sediments mainly consisted of silt with small shell fragments. Since the 461 re-suspensiond depth by the 2011 tsunami of the Great East Japan Earthquake was 20-25cm in 462 the Shizu and Nagasawa cores, the un-disturbed lower sediments seem to be deposited over 463 more than 100 years. The first report of ellipsoidal Alexandrium cysts from the surface 464 sediments of Ofunato Bay was only in 1980 (Fukuyo 1980). However, small amounts of 465ellipsoidal Alexandrium cysts either filled with protoplasm or empty (ca. 210 cysts/g in Shizu

466 core and ca. 260 cysts/g in Nagasawa core) were recovered from deeper un-disturbed sediments 467 of the cores that were obviously deposited before 1980 according to depositional age of 468 sediments below 20-25cm depth. This means that A. tamarense and/or A. catenella might have 469 inhabited Ofunato Bay for over 100 years. This conclusion is partially supported by the 470evidence that appearance of A. tamarense and/or A. catenella in Funka Bay, North Japan, more 471than 100 years ago (Miyazono et al., 2012). The 25cm thick interval of sediments re-suspended 472and re-deposited by the tsunami could cover more than 125 years. Although it is impossible to 473 know the exact number of ellipsoidal Alexandrium cysts re-suspended after the tsunami in all 474Ofunato Bay area, the most likely explanation is that large amounts of ellipsoidal Alexandrium 475cysts preserved in the sediments before disturbed by the tsunami were eroded, re-suspended and 476 re-deposited in Ofunato Bay.

477Such unusual re-deposition of ellipsoidal Alexandrium cysts was also observed in Sendai 478 Bay after the 2011 Great East Japan tsunami (Kamiyama et al. 2014). However, the cyst density 479of ellipsoidal Alexandrium cyst in Ofunato Bay was much higher than that of Sendai Bay. The 480 mechanism for accumulation of these Alexandrium cysts by the tsunami seems to be the same at 481 Ofunato Bay and Sendai Bay. However, different cyst densities preserved in the older, 482non-disturbed sediments due to the pre-tsunami bloom histories of A. tamarense might explain 483 the different cyst densities in the re-deposited sediments of both bays. Therefore, knowledge of 484 the vertical distribution and germination viability of living cysts in the sediments is important 485 for predicting future blooms, because living cysts in deeper bottom sediments can still be 486 capable of germination when brought back to the surface (Mizushima and Matsuoka, 2004).

In summary, the highly unusual abundance of ellipsoidal *Alexandrium* cysts (mainly *A. tamarense*) in the Shizu and Nagasawa cores was caused mainly by deep erosion (more than 20 cm thick), re-suspension, and re-deposition of bottom sediments by the tsunami at the Great East Japan Earthquake, combined with the higher cyst production and more frequent bloom-forming characteristics of *A. tamarense* in Ofunato Bay. This explanation is supported

492by additional information on tsunami-related sedimentation processes. Nishi et al. (2013) 493 studied change of sea floor environments and the mechanism of flow dynamics in several inner 494 bays including Ofunato Bay along the Sanriku Coast. In the central part of Ofunato Bay, 495 ignition loss was lower in sediments re-deposited after the tsunami (Nishi et al. 2013). In our 496 cores, re-deposited sediment became coarser as shown by median particle size increasing from 497 10 to 15µm. In addition, our two core sites were reported as sediment accumulation areas after 498 the tsunami (Kamaishi Office of Tohoku Regional Development Bureau 2012). These data 499 suggest that during the tsunami wave surges, re-suspended sediments were sorted, with the finer 500 silt to clay sediments containing organic materials being exported from the bay by the return 501flow of the tsunami. As a result, coarser silt particles were concentrated in the center of the bay. 502A similar change of sediment particles were also observed in Otsuchi and funakosi bays along 503 the sanriku coast (Seike et al. 2013. Since the size of ellipsoidal Alexandrium cysts (35-56µm x 504 23-35µm in size) is like that of coarse silt, these cysts were preserved in re-deposited sediments.

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#### 506 **3. Relation between shellfish aquaculture and tsunami events**

507In Ofunato Bay, the first mass PSP incident happened on May of 1961 due to ingestion of 508 Akazara-scallops that attached to cultured oysters (Kawabata et al., 1962), and later the 509causative plankton was identified as Gonyaulax sp. (Murano 1975). Fishermen in Ofunato Bay 510 often ate Akazara-scallops, starting around March; however, this PSP incident occurred in May 511of 1961. Because the year 1961 followed the 5.5m-high Chilean tsunami surge into Ofunato 512Bay on May 23, 1960, a relation between the 1961 PSP incident and the Chilean tsunami event 513was suspected. According to our core studies, a tsunami could erode and re-suspend the bottom 514sediment, including ellipsoidal Alexandrium cysts which were then re-deposited on the sea floor 515surface after the tsunami. Such huge amounts of cysts might have seeded of the unusually large 516A. tamarense blooms that occurred along the coast of East and North Japan during June 2011. 517The tsunami waves caused by the 2011 Great East Japan Earthquake on March 11, 2011 were

also observed in Alaska 6-9 hours later, with a height of 1.5m, and 2m-high waves reached Chile 21 hours later. A PSP outbreak was noticed in Southeast Alaska in May to June 2011 (MMWR 2011), about 2 months after being hit by the tsunami of the Great East Japan Earthquake. At present, however, it is uncertain if this PSP event was caused by the processes similar to those in Ofunato Bay and Sendai Bay, because in Alaska PSP incidents frequently occur in May to June with non-tsunami events.

524The Chilean tsunami attacked Ofunato Bay with a wave of 1.9-5.7m height, which was 525slightly lower than that of the Great East Japan Earthquake tsunami, but the effect on the bottom 526 sediment seemed to be nearly the same as the Great East Japan Earthquake tsunami. 527Unfortunately, the sediments re-deposited by the Chilean tsunami did not remain in the present 528location because the Great East Japan Earthquake tsunami eroded and re-suspended these 529sediments as described above. However, it is reasonable to postulate that re-suspended 530 ellipsoidal Alexandrium cysts filled with protoplasm would become the seeding source for the 531forthcoming 1961 bloom because ellipsoidal Alexandrium cysts were preserved in sediments 532before the Chilean tsunami. Consequently, the first PSP incident in Ofunato Bay might be the 533 result of the Alexandrium bloom triggered by germination of living cysts one year after of the 534Chilean tsunami in almost the same manner as the events following the 2011 tsunami of the 535Great East Japan Earthquake.

Another environmental issues should be discussed here. In Ofunato Bay, the oyster aquaculture first started at Akasaki on 1887 (Miyazawa & Hayakawa 1994) but no paralytic shellfish poisoning was ever reported before the Chilean tsunami despite occurrences of several huge earthquake-generated tsunamis that surged into this bay (Sekiguchi 2010). This lack of pre-1961 PSP outbreaks may reflect the low scale of shellfish aquaculture industry development in this bay before the Chilean tsunami. Interestingly, the opening of the scallop culture industry in Ofunato Bay took place in 1961, just after the Chilean tsunami. This historical circumstance, together with the environmental situation imposed by the tsunami of the Great East Jana
Earthquake, provided the following account of the first (1961) PSP incident in Ofunato Bay.

545When the Chilean tsunami wave hit Ofunato Bay on May 24 of 1960, many ellipsoidal 546 Alexandrium cysts preserved in the old sediments might have been re-suspended and 547re-deposited on the sea floor. It is very difficult to know whether in 1960 these re-deposited 548cysts could germinate or not (probably not), because the re-deposited sediments were later 549re-disturbed by the Great East Japan Earthquake tsunami. At the next spring of 1961, the bloom 550season of A. tamarense, some of these cysts might have germinated into vegetative cells which 551might form dense blooms of A. tamarense. These dense blooms were also supported by water 552quality changes such as eutrophication in this bay because the water quality in this enclosed bay 553 is easily affected by increased population around the bay (eg. Matsuoka 1999). In the case of 554Ofunato Bay, the population had increased nearly 10% per year since 1920 and reached 555approximately 400,000 by 1980 (http://demography.blog.fc2.com/blog-entry-4839.html based 556on National census data). This population increase was accompanied with transportation of 557 waste-water from land and resulting increase of nutrients (eutrophication). Unfortunately before 558 1971, there was no regulation of industrial and domestic wastewater from lands in Ofunato Bay. 559 Phytoplankton population might be also changed by such water quality change in Ofunato Bay 560around 1960's. The vegetative cells forming dense blooms of A. tamarense would be consumed 561by cultured and/or natural shellfish (oysters and scallops) in the bay. A part of these toxicated 562shellfish (Akazara-scallops) were consumed by local people who then suffered with paralytic 563 shellfish poisoning. During the bloom, vegetative cells of A. tamarense produced many new 564resting cysts, which also remained within the bay because of the semi-enclosed natural shape. 565Thereafter, these preserved cysts could act as seeding sources for subsequent blooms. In 566 addition, since around 1960 oyster cultures have became popular in Ofunato Bay, and local 567 people would be more scientifically aware of PSP events. The scientific work informs us that 568unpleasant phenomena for human society, such PSP, will often manifest after catastrophic

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natural disasters (the tsunami in this case) and will be worsened by artificial changes by rapid increase of populations and shellfish aquacultures for natural environments.

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572 Summary

573 1. The sediment above ca. 25cm depth from the sea floor in the center of Ofunato Bay was
574 re-deposited by the tsunami of the 2011 Great East Japan Earthquake.

575 2. The fluctuation of vertical distribution of sediment grain size and density of ellipsoidal
 576 *Alexandrium* cysts in bay sediment cores might reflect the multiple surges of the tsunami.

577 3. The maximum density of ellipsoidal Alexandrium cysts recorded at the 6-7cm depth in the

578 Nagasawa core was the result of the deep erosion and re-suspension of older sediments and
579 the specific cyst-forming capability of *Alexandrium tamarense*.

4. The first PSP incident on 1961 in Ofunato Bay probably resulted from huge blooms of *Alexandrium tamarense* initiating from resting cysts which were moved to the sea floor by deep erosion and re-suspension of sediments involving ellipsoidal *Alexandrium* cysts filled with fresh protoplasm after the Chilean tsunami in 1960, combined with deterioration of water quality due to rapid increase of population and development of shellfish aquacultures.

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| 802 | Explanations | of Figures. | Tables and | Supp | lemental | Data |
|-----|--------------|-------------|------------|------|----------|------|
|     |              |             |            |      |          |      |

803

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804 Figure 1
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- 805 Coring sites in Ofunato Bay, Northeast Japan
- 806 The Shizu coring site was located at ca. 800m north of the core O series site by Port and Airport
- 807 Research Institute and Japan NUS (2001) and of the Station P in Kamaishi Office of Tohoku
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- 809

#### 810 Figure 2

811 Sediment characteristics of the Shizu core (A) and Nagasawa core (B). Vertical lines indicate 812 the depth of sediments which were re-deposited at the Great East Japan Earthquake tsunami on 813 March 11 0d 2011. Vertical curved lines showed slender tubes formed by polychaetes, which 814 may indicate the re-deposited parts of sediments.

815

#### 816 Figure 3

817 Sediment characteristics; grain size composition, median particle size, and water content of the 818 Shizu core. Two horizontal dashed lines indicate changes of median particle size and water 819 content. The interval might suggest a transitional zone between re-deposited part by the Great 820 East Japan Earthquake tsunami and un-disturbed part. Lines with arrow indicate increase or 821 decrease of median particle size and water content.

822

#### 823 Figure 4

824 Sediment characteristics; grain size composition, median particle size, and water content of the 825 Nagasawa core. Two horizontal dashed lines indicate changes of median particle size, and water 826 content. The interval might suggest a transitional zone between re-deposited part by the Great 827 East Japan Earthquake tsunami and un-disturbed part. Lines with arrow indicate increase or

| 828 | decrease of median particle size and water content. Within the | un-disturbed part, media | in particle |
|-----|--|--------------------------|-------------|
| 829 | size was rather unstable.                                      |                          |             |

830

| 831 Figu | re | 5 |
|----------|----|---|
|----------|----|---|

832 Dominant dinoflagellate cysts in the Shizu and the Nagasawa core sediments.

833 1-4; Ellipsoidal Alexandrium cysts (1 & 3 cysts filled with protoplasm, 2 & 4 empty cysts), 5
834 Spiniferites bulloideus (Cookson et Deflandre) Sarjeant, 6 Brigantedinium sp. indet.
835 (Protoperidinium cyst filled with protoplasm), 7 Votadinium spinosum Reid (=cyst of
836 Protoperidinium claudicans (Paulsen) Balech)

837

#### 838 Figure 6

839 Vertical distributions of total dinoflagellate cysts and major groups in the Shizu Core.

840 *Alexandrium* cysts included all of cysts filled with protoplasm and empty cysts. Dashed lines 841 show the 5cm interval of observed samples. Shadow indicates a transitional zone between the

842 re-deposited and un-disturbed sediments after the Great East Japan Earthquake tsunami.

843

844 **Figure 7** 

Vertical distributions of total dinoflagellate cysts and major groups in the Nagasawa Core. *Alexandrium* cysts included all of cysts filled with protoplasm and empty cysts. Dashed lines show the 5cm interval of observed samples. Shadow indicates a transitional zone between the

848 re-deposited and un-disturbed sediments after the Great East Japan Earthquake tsunami.

849

850 Table 1

851 List of dinoflagellate cysts, acritarchs and other palynomorphs observed in the Shizu and

852 Nagasawa cores

853

| 854 | Table 2   |
|-----|---|
| 855 | Cyst densities of Alexandrium tamarense/cetenella including A. fudyense in the previous     |
| 856 | records.  |
| 857 |   |
| 858 | Table 3   |
| 859 | Hieghst occurrence of ellepsoidal Alexandrium cysts in Ofunato Bay (cysts/cm <sup>3</sup> ) |
| 860 |   |
| 861 | Supplemental Data 1   |
| 862 | Occurrence of dinoflagellate cysts in the Shizu core  |
| 863 |   |
| 864 | Supplemental Data 2   |
| 865 | Occurrence of dinoflagellate cysts in the Nagasawa core                                     |
| 866 |   |

#### Figure 1



#### Figure 2









Figure 3

## Figure 5



## Figure 6







Cysts/g sediment dry weight

#### Table 1 List of dinoflagellate cysts, acritarchs and other palynomorphs

#### Photo/mixotrophic dinoflagellate cysts:

Alexandrium tamarense/catenella (as ellipsoidal Alexandrium cyst) Alexandrium fraterculus/minutum/pseudogoniaulax<sup>#</sup> (as spherical Alexandrium cyst) Spiniferites bulloideus (Deflandre & Cookson) Sarjeant ls\* Spiniferites delicates Reid\* Spiniferites mirabilis (Rossignol) Sarjeant\* Spiniferites ramosus (Ehrenberg) Mantel\* Spiniferites hyperacanthus (Deflandre & Cookson) Cookson & Eisenack\* Spiniferites elongatus Reid\* Protoceratium reticulatum (Claparéde & Lachmann) Bütschli = Operculodinium centrocarpum (Deflandre & Cookson) Wall sensu Wall & Dale\* Lingulodinium machaerophorum (Claparéde & Lachmann) Wall\* Tuverculodinium vancampoae (Rossignol) Wall \* Scrippsiella spp. Polykrikos hartmannii Zimmermann Heterotrophic dinoflagellate cysts: Polykrikos kofoidii Charron Polykrikos schwartzii Bütschli Niea acanthocysta (Kawami, Iwataki & Matsuoka) T. Liu, K.N. Mertens & H. Gu Protoperidinium leonis (Pavillard) Balech Protoperidinium obtusum (Karsten) Park & Dodge Protoperidinium latissinum (Balech) Balech Protoperidinium latidorsale (P.-A. Dangeard) Balech Protoperidinium paraoblongum Sarai, Yamaguchi, Kawami & Matsuoka Protoperidinium steidingerae Balech Protoperidinium subinerme (Paulsen) A. R. Loeblich Protoperidinium americanum (Gran & Braarud) Balech Brigantedinium cariacoense (Wall) Reid\* Briantedinium simplex (Wall) Reid\* Brigantedinium majusculum Reid\* Brigantedinium irregulare Matsuoka ex Head

Stelladinium stellatum (Wall) Reid \*

Dubridinium caperatum Reid\*

Echinidinium aculeatum Zonneveld\*

Echinidinium spp\*.

Selenopemphix nephroides Benedek) Benedek & Sarjeant\*

Selenopenphix quanta (Bradford) Matsuoka (large form)\*

Selenopenphix quanta (Bradford) Matsuoka (small form)\*

Trinovantedinium applanatum (Reid) Bujak & Davies\*

Votadinium spinosum Reid\*

Votadinium calvum Reid\*

(\* cyst-based biological name; # based on observation of plankton forms by Kaga et al. 2006)

#### Acritarchs:

Baltisphaeridium sp.

Halodinium major Bujak

#### Other marine palynomorphs:

Archeomonas sp.

Tintinnopsis sp.

Microforaminiferal linings (uni-serial type, bi-serial type, coiled type)

Crustacean resting eggs and body fragments

 Table 2

 Cyst densities of Alexandrium tamarense/cetenella including A. fudyense

| Area   | Maximum cyst density              | References                    |
|--|-----------------------------------|-------------------------------|
|  | Surface sediment                  |                               |
| Gulf of Maine, USA/Canada                      | $2,000 \text{ cysts cm}^{-3}$     | Anderson et al. (2005)*       |
| Okhotsuku Sea and Soya Strait, Hokkaido, Japan | 2,022 cysts wet g <sup>-1</sup>   | Shimada and Miyazono (2005)   |
| Ise Bay, central Japan                         | $2,254 \text{ cysts cm}^{-3}$     | Ishikawa et al. (2007)        |
| Funka Bay, Hokkaido, Japan                     | 2,568 cysts wet g <sup>-1</sup>   | Shimada and Miyazono (2005)   |
| Yellow Sea                                     | 3,778 cysts dry g <sup>-1</sup>   | Cho & Matsuoka (2001)         |
| Hiroshima Bay, West Japan                      | 4,454 cysts $cm^{-3}$             | Yamaguchi et al. (2002)       |
| Osaka Bay, West Japan                          | 5,683 cysts cm <sup>-3</sup>      | Yamamoto et al. (2009)        |
| Mikawa Bay, Central Japan                      | $7,311 \text{ cysts cm}^{-3}$     | Ishikawa et al. (2007)        |
| Bay of Fundy, Canada                           | 7,440 cysts cm <sup>-3</sup>      | Martin and Wildish (1994)*    |
| Tokuyama Bay, West Japan                       | 8,137 cysts cm <sup>-3</sup>      | Yamaguchi et al. (2002)       |
| Aso Bay, Tsushima Island, West Japan           | 8,900 cysts cm <sup>-3</sup>      | Matsuoka & Lee (1994)         |
| Chukchi Sea, the Arctic Ocean                  | $10,600 \text{ cysts cm}^{-3}$    | Natsuike et al. (2013)        |
| Puget Sound, Washington, USA                   | 12,125 cysts cm <sup>-3</sup>     | Horner et al. $(2011)^{\#}$   |
| Bedford Basin, Nova Scotia, Canada             | 220,872 cysts dry $g^{-1}$        | Lacasse et al. (2013)**       |
|  | Surface sediment after the tsuna  | mi                            |
| Sendai Bay, Northeast Japan                    | 8,190 cysts cm <sup>-3</sup>      | Kamiyama et al (2014)         |
| Kesennnuma Bay, Northeast Japan                | 6,383 cysts cm <sup>-3</sup>      | Ishikawa et al. (2015)        |
| Kesennnuma Bay, Northeast Japan                | 4,170 cysts cm <sup>-3</sup>      | Nishitani et al. (2012)       |
| Funka Bay, Hokkaido, Japan                     | $3,200 \text{ cysts wet g}^{-1}$  | Natsuike et al. (2014)        |
| Ofunato Bay, Northeast Japan                   | 62,000 cysts dry g <sup>-1</sup>  | Ikeda et al. (2013)           |
|  | Core sediment after the tsunam    | i                             |
| Ofunato Bay, Northeast Japan                   | 155,210 cysts dry g <sup>-1</sup> | This study                    |
|  |                                   | * A. fundyense                |
|  |                                   | ** A. tamarense               |
|  |                                   | <sup>#</sup> A. catenella     |
|  |                                   | No mark A.                    |
|  |                                   | <i>tamarense/catenella</i> or |

| Age     | Innter Bay | Central Bay   | Bay Mouth | Reference         |
|---------|------------|---------------|-----------|-------------------|
| 1980    | 642        | 1326          | 62        | Fukuyo (1980)     |
| 1985    | 143        | 490           | 25        | Iwate PFES (1986) |
| 1986    | 185        | 955           | 20        | Iwate PFES (1987) |
| 1987    | 320        | 490           | ND        | Iwate PFES (1988) |
| 1988    | 1050       | 1200          | 30        | Iwate PFES (1989) |
| 1990    | 433        | 1000          | 500       | Iwate PFES (1990) |
| 1992    |            | No iformation |           | Iwate PFES (1993) |
| 1998    | 430        | 620           | ND        | Iwate FTC (1999)  |
| 1999    | 467        | 438           | 40        | Iwate FTC (1999)  |
| Average | 458        | 815           | 113       |                   |
|         |            |               |           |                   |

Table 3 Hieghst occurrence of elongate Alexandrium cysts in Ofunato Bay (cysts/cm<sup>3</sup>)

PFES: Prefectural Fisheries Experimental Station FTC: Fisheries Techology Center

## Supplemental Data 1 Shizu Core

| Sample dent                       | h     |         |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |         |       |       |       |
|-----------------------------------|-------|---------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|---------|-------|-------|-------|
| Species                           | 0-0.5 | 0.5-1 1 | -1.5  | 1.5-2 | 2-2.5 | 3-3.5 | 4-4.5 | 5-5.5 | 6-6.5 | 7-7.5 | 8-8.5 | 9-9.5 | 10-11 | 11-12 | 12-13 | 13-14 | 14-15 | 15-16 | 16-17 | 17-18 | 18-19 | 19-20 | 20-21 | 21-22 | 22-23 | 23-24 | 24-25 | 25-26 | 30-31 | 35-36 | 40-41 | 45-46 | 50-51 : | 55-56 | 60-61 | 64-65 |
| Alexandrium                       |       |         |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |         |       |       |       |
| Alexandrium tamarense / catenella | 4857  | 6393    | 6575  | 18588 | 15833 | 12971 | 9315  | 6098  | 5333  | 5902  | 8473  | 12300 | 12829 | 10500 | 10457 | 5512  | 8777  | 16171 | 12264 | 10971 | 26294 | 16685 | 20303 | 4093  | 1952  | 3204  | 844   | 558   | 71    | 62    |       | 63    | 60      |       |       | -     |
| Empty-A tamarense                 | 428   | 214     | 425   | 794   | 472   | 143   | 368   | 268   | 214   | 73    | 184   | 425   | 229   | 475   | 257   | 51    | 416   | 1257  | 323   | 1028  | 558   | 171   | 181   | 250   | 333   |       |       | 46    | 142   | 125   |       |       | 20      | 96    | 21    |       |
| Alexandrium affine / leei         | 785   | 286     | 450   | 794   | 555   | 1000  | 921   | 317   | 238   | 390   | 394   | 525   | 514   | 475   | 257   | 282   | 361   | 629   | 852   | 685   | 970   | 428   | 787   | 375   | 95    | 45    | 111   | 162   | 47    | 120   |       |       | 20      |       |       |       |
| Empty-A.affine / leei             | 142   |         |       |       |       |       | 105   | 98    | 24    | 48    |       |       | 28    |       | /     |       |       | 57    |       |       |       |       |       | 31    | 23    |       |       |       | 23    |       |       |       | 80      | 115   | 42    | 40    |
|                                   |       |         |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |         |       |       |       |
| Other gonyaulacoid                |       |         |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |         |       |       |       |
| Spiniferites                      |       |         |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       | 86    | 147   |       |       | 57    | 30    |       |       |       |       |       | 47    | 83    |       | 21    |         |       | 21    | 20    |
| Spiniferites mirabilis            |       |         |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       | 29    |       |       |       |       | 31    | 47    |       |       |       |       |       | 21    |       | 20      | 173   | 21    | 20    |
| Spiniferites ramosus              |       | 36      |       | 58    | 166   | 57    | 105   | 415   | 286   | 292   | 316   | 350   |       |       |       |       |       | 29    | 147   |       | 29    |       |       | 62    | 71    | 22    |       | 139   | 333   | 166   | 489   | 638   | 220     | 269   | 191   | 142   |
| Spiniferites hyperacanthus        | 142   | 36      | 25    |       |       |       |       |       |       |       |       | 50    |       | 50    | 29    | 28    | 55    | 5     |       |       | 29    |       |       |       |       | с     |       |       | 47    | 20    | 21    | 21    |         | 19    |       |       |
| Spiniferites elongatus            | 214   | 36      |       |       | 55    |       | 52    |       | 48    |       |       | 25    |       |       |       |       | 27    | 7     | 29    |       |       | 28    | 30    |       |       | 22    | 22    |       | 71    | 41    | 42    | 21    | 40      | 38    |       | 102   |
| Spiniferites bentorii             |       |         |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       | 125   |       |       |         |       |       |       |
| Spiniferites bulloideus           | 4357  | 2036    | 1900  | 2588  | 2333  | 1971  | 2157  | 1829  | 1524  | 1171  | 1158  | 1725  | 2371  | 2075  | 1914  | 1467  | 2972  | 2371  | 1794  | 2771  | 2441  | 1600  | 1181  | 1125  | 1309  | 1500  | 911   | 1069  | 2166  | 1604  | 1021  | 617   | 300     | 1711  | 787   | 612   |
| Protoceratium reticulatum         |       |         |       |       |       |       | 78    | 49    | 24    | 24    | 132   |       |       |       |       |       |       | 29    |       | 171   |       |       | 30    | 281   |       |       | 22    | 46    | 47    |       | 21    | 21    | 80      |       | 85    | 81    |
| Tuberculodinium vancampoae        |       |         |       |       |       | 86    | 26    | 24    |       |       |       | 25    |       | 25    |       |       | 55    | 5 29  |       | 57    | 29    | 28    |       | 31    | 23    | 22    | 22    | 23    | 95    | 20    | 42    | 21    |         | 38    | 21    | 20    |
|                                   |       |         |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |         |       |       |       |
| Gymnodinioid                      |       |         |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |         |       |       |       |
| Pheopolykrikos hartmannii         |       |         |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       | 83    |       |       |         |       |       |       |
| Polykrikos kofoidii               | 285   |         | 50    | 58    | 55    | 86    | 26    |       | 71    | 24    | 53    | 125   | 57    | 25    | 29    | 51    | 27    | 29    | 29    | 57    | 29    | 28    | 60    | 62    | 71    | 45    | 44    | 69    | 142   |       | 42    | 21    | 40      | 76    | 127   | 81    |
| Polykrikos schwartzii             | 214   | 214     | 175   | 205   | 83    | 57    | 52    | 146   | 167   | 73    | 105   | 175   | 171   | 150   | 114   | 51    | 194   | 200   | 205   | 371   | 352   |       | 151   | 156   | 71    | 45    | 44    |       | 404   | 62    | 212   | 127   | 260     | 211   | 170   | 163   |
|                                   |       |         |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |         |       |       |       |
| Peridinioid                       |       |         |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |         |       |       |       |
| Protoperidinium conicum           | 285   | 36      | 75    | 117   | 166   | 57    | 263   | 146   | 71    | 146   | 60    | 125   | 143   | 100   | 171   | 153   | 305   | 5 229 | 58    | 57    | 323   | 228   | 212   | 250   | 95    | 68    | 88    | 23    | 285   |       | 106   | 63    | 80      | 76    | 297   | 326   |
| Protoperidinium leonis            | 1714  | 571     | 575   | 617   | 527   | 543   | 526   | 537   | 1048  | 902   | 605   | 900   | 657   | 625   | 886   | 615   | 361   | 1114  | 382   | 1028  | 1058  | 428   | 696   | 531   | 71    | 272   | 400   | 279   | 595   | 145   | 212   | 170   | 280     | 403   | 127   | 265   |
| Protoperidinium obtusum           | 142   | 36      |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       | 22    |       |       | 354   |       |       |         |       |       |       |
| Protoperidinium claudicans        | 214   | 107     | 75    | 58    | 111   | 57    | 78    | 122   | 71    | 24    | 105   | 100   | 57    | 50    | 57    | 51    | 27    | 86    | 29    | 85    | 88    |       |       | 125   | 71    | 68    | 22    | 69    | 190   |       | 63    | 106   | 120     | 288   | 170   | 142   |
| Votadinium calvum                 |       |         |       | 58    |       |       |       |       |       |       |       |       |       | 75    |       | 51    | 27    | 7     | 58    | 28    | 88    |       | 60    | 62    |       |       | 22    |       | 95    | 125   | 21    | 21    | 20      | 57    | 21    |       |
| Protoperidinium latidorsale       |       |         |       |       |       |       | 26    |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       |       | 83    |       |       |         |       |       |       |
| Protoperidinium inaequale         |       |         |       |       |       |       | 52    |       |       |       | 79    |       | 28    |       |       |       |       |       |       |       |       |       | 30    |       |       |       |       | 23    | 47    |       | 21    |       | 20      |       |       |       |
| Protoperidinium steidingerae      |       |         |       |       |       | 28    | 26    |       |       |       |       |       |       |       |       |       |       | 28    |       |       |       |       |       |       |       |       |       | 23    |       |       |       |       |         |       |       |       |
| Protoperidinium symmetricum       | 214   | 71      | 25    | 88    |       |       |       | 73    | 47    | 73    |       |       | 28    |       |       |       |       |       |       | 28    |       |       | 60    |       |       |       |       | 23    | 71    |       | 63    | 21    |         |       | 42    |       |
| Protoperidinium subinerme         | 214   | 36      | 25    | 29    |       | 28    | 52    | 49    | 48    | 24    | 53    | 50    |       | 100   |       | 51    |       | 28    | 29    | 28    | 29    |       | 30    |       | 23    | 68    | 22    | 23    | 166   | 41    | 85    | 42    | 60      | 38    | 85    | 102   |
| Protoperidinium americatum        | 285   |         | 125   | 88    | 194   | 86    | 131   |       | 95    | 24    | 53    | 175   | 171   | 150   | 29    | 102   |       | 171   |       | 85    | 58    | 114   | 60    | 62    | 71    | 22    | 155   | 116   | 166   | 62.5  | 276   | 106   | 120     | 192   | 106   | 81    |
| Stelladinium stellatum            | 71    | 36      |       |       |       |       |       |       |       |       | 26    |       |       | 100   |       |       |       | 28    |       |       |       | 28    |       |       |       |       |       | 23    |       | 125   |       |       |         |       |       |       |
| Trinovantedinium applanatum       | 71    |         |       |       |       | 28    |       |       |       |       |       | 50    | 29    |       |       |       | 27    | 7     | 88    |       |       |       | 30    |       |       | 22    |       | 23    | 23    |       |       |       |         | 19    | 21    | 20    |
| Bri gantedinium spp.              | 4071  | 1714    | 1325  | 1882  | 1638  | 1000  | 763   | 780   | 1976  | 1122  | 1184  | 1325  | 1057  | 1875  | 2371  | 1307  | 2027  | 2029  | 911   | 1142  | 2176  | 1000  | 1060  | 1093  | 976   | 1181  | 822   | 418   | 523   | 62.5  | 468   | 510   | 500     | 365   | 489   | 346   |
| Bri gantedinium simplex           | 3500  | 607     | 975   | 1176  | 777   | 1200  | 1210  | 1171  | 881   | 1122  | 1132  | 1125  | 1314  | 725   | 1029  | 923   | 944   | 1371  | 588   | 914   | 1176  | 1628  | 1000  | 937   | 1928  | 590   | 511   | 860   | 2166  | 479   | 1361  | 1319  | 1140    | 1653  | 1702  | 897   |
| Brigantedinium cariacoense        | 71    | 36      | 75    |       | 166   |       | 78    | 98    | 167   | - 98  | 263   | 150   | 314   | 300   | 114   | 153   | 138   | 257   | 205   | 457   | 176   | 114   | 272   | 187   | 119   | 181   | 200   | 162   | 285   | 1312  | 255   | 170   | 340     | 96    | 276   | 61    |
| Brigantedinium majusculum         | 642   | 36      | 700   | 647   | 888   | 514   | 368   | 317   | 857   | 244   | 110   | 400   | 343   | 50    |       |       |       | 285   | 29    | 171   | 205   | 171   | 151   | 31    | 47    |       |       |       | 190   | 187   | 106   | 127   | 160     | 346   | 255   | 244   |
| Brigantedinium irregulare         |       |         |       |       |       |       | 26    |       |       |       |       |       |       | 75    |       | 25    |       |       |       |       |       | 28    | 30    | 31    |       | 45    | 22    |       |       | 83    | 63    |       |         |       | 21    | 20    |
| Dubridinium caperatum             | 357   | 107     | 175   | 205   | 277   | 114   | 105   | 122   | 357   | 98    | 105   | 350   | 314   | 200   | 200   | 179   | 222   | 200   | 205   | 171   | 264   | 85    | 30    | 218   | 166   | 90    | 111   | 93    | 214   |       | 63    | 127   | 40      | 192   | 340   | 551   |
| Scrippsiella spp.                 | 642   | 357     | 225   | 176   | 83    | 314   | 131   | 195   | 190   | 49    | 289   | 125   | 57    |       | 86    |       |       | 200   | 117   |       | 29    |       | 30    |       |       | 22    |       | 69    | 809   | 145   | 63    |       |         |       |       |       |
| Total (cysts/g)                   | 23917 | ####    | 13975 | 28226 | 24379 | 20340 | 17040 | 12853 | 13738 | ##### | ##### | 20600 | 20711 | 18200 | 18001 | 11052 | 16962 | 26913 | 18518 | 20305 | 36401 | 22849 | 26504 | 10024 | 7562  | 7534  | 4417  | 4339  | 9460  | 5595  | 5137  | 4353  | 4020    | 6471  | 5438  | 4336  |

#### Supplemental Data 2

Nagasawa Core

| Species                          | 0-0.5 | 0.5-1 | 1-1.5 | 1.5-2  | 2-2.5 | 2.5-3 | 3-3.5 | 3.5-4 | 4-4.5  | 4.5-5 | 5-6   | 6-7    | 7-8    | 8-9    | 9-10  | 10-11   | 11-12  | 12-13  | 13-14  | 14-15 | 15-16 | 16-17  | 17-18 | 18-19   | 19-20 | 20-21 | 21-22 | 22-23 | 23-24 | 24-25 | 25-26 | 30-31 | 35-36 | 40-41    | 45-46     | 50-51 | 55-56  | 60-61     | 65-66      | 70-71 | 74-75 |
|----------------------------------|-------|-------|-------|--------|-------|-------|-------|-------|--------|-------|-------|--------|--------|--------|-------|---------|--------|--------|--------|-------|-------|--------|-------|---------|-------|-------|-------|-------|-------|-------|-------|-------|-------|----------|-----------|-------|--|-----------|------------|-------|-------|
| Alexandrium                      |       |       |       |        | 1     |       |       | Î     |        |       |       |        |        |        |       | 1       | 1      |        |        | 1     |       |        | Î     | Î       |       |       |       |       |       |       |       | Î     |       | 1        | $\square$ | i     | ( The second sec |           | Ê          |       |       |
| Alexandrium tamarense / catenell | 14    | 41100 | 80931 | 115182 | 38010 | 80117 | 38057 | 81088 | 109857 | 39324 | 73342 | 155218 | 109571 | 140852 | 5412  | 9 78176 | 115333 | 97527  | 86058  | 60470 | 71722 | 122054 | 31857 | 24444   | 3090  | 5204  | 1808  | 6520  | 180   | 520   | 836   | 265   | 218   | 17       | 38        | 118   | 125  | 67        | 14         | 44    | 55    |
| Alexandrium affine/leei          | 14    | 2700  | 241   | 636    | 210   | 29    | 228   | 382   | 171    | 189   | 85    | 62     | 314    | 441    | 90    | 5       | 194    | 194    | 88     | 264   | 361   | 54     | 261   | 88      | 113   | 142   | 148   | 229   | 40    | 100   | 20    | 20    | ) 73  | 86       | 76        | 196   | 71   | (         | 14         | 73    | 18    |
|                                  | 28    | 43800 | 81172 | 115818 | 38220 | 80146 | 38285 | 81470 | 110029 | 39514 | 73427 | 155280 | 109885 | 141293 | 5422  | 5 78176 | 115527 | 97721  | 86146  | 60734 | 72083 | 122108 | 32118 | 3 24532 | 3203  | 5346  | 1956  | 6749  | 220   | 620   | 856   | 285   | 291   | 103      | 114       | 314   | 196  | 67        | 28         | 117   | 73    |
| Other gonvaulacoid               |       |       |       |        |       |       |       |       |        |       |       |        |        |        |       |         |        |        |        |       |       |        |       |         |       |       |       |       |       |       |       | 1     |       |          |           |       | $ \rightarrow$   |           |            | _     |       |
| Spiniferites spp.                |       |       |       |        |       |       |       | 1     |        |       |       |        |        |        |       | 58      | 3      |        |        |       |       |        | 1     | 1       |       |       |       | 20    |       |       | 20    | )     |       |          |           |       |  | $ \neg  $ |            |       |       |
| Spiniferites mirabilis           |       |       |       |        |       |       |       |       |        |       |       |        |        |        |       |         |        |        |        |       |       |        |       |         |       |       |       |       |       |       |       |       |       | 1        |           |       | 17   | (         |            |       | 18    |
| Spiniferites ramosus             | 41    |       | 24    | 61     |       | 58    |       |       | 86     | 27    |       |        |        | 58     | 3 6   | 4 29    | )      | 55     |        |       | 27    | 54     | 47    | 133     | 68    | 41    |       |       |       | 20    | 40    | 40    | 0 18  | ,        | 38        | 59    | 107  | 16        | 41         | 308   | 271   |
| Spiniferites hyperacanthus       | 68    |       | 24    |        |       |       |       |       |        |       |       | 31     |        |        |       |         |        |        | 29     |       |       |        |       |         |       |       |       | 62    | 20    | 20    | 20    | 122   | 2 15  | ,        | 57        | 59    | 89   | 152       | 68         |       | 31    |
| Spiniferites elongatus           |       | 50    |       | 30     | 33    |       |       | 29    | 29     | 54    |       |        |        |        | 32    | 2 29    | )      |        |        |       |       | 54     |       | 44      | 90    | 61    | 63    | 62    | 120   | 60    | 40    | 40    | ) 55  | ,        | 19        | 20    | 89   | 33        |            | 44    | 55    |
| Spiniferites bulloideus          | 757   | 5650  | 3034  | 3848   | 2700  | 2235  | 1428  | 2764  | 3629   | 3351  | 6342  | 5156   | 3742   | 3941   | 532   | 2 5911  | 4222   | 7111   | 6235   | 4382  | 9472  | 1378   | 1547  | 1422    | 4136  | 3836  | 1212  | 1270  | 680   | 780   | 1122  | 1285  | 1345  | 844      | 576       | 882   | 1803   | 1000      | 560        | 2044  | 2500  |
| Protoceratium reticulatum        |       | 50    |       |        |       | 29    | 1     | 58    | 29     |       | 28    |        |        |        |       |         | 27     | 55     |        |       |       | 27     |       |         | 68    | 40    | 63    |       |       | 20    |       |       | 18    | 17       |           | 20    | 17   | 50        |            |       |       |
| Pyrophacus steinii               |       | 50    |       |        |       | 29    | 1     |       | 29     |       |       | 31     |        |        |       |         |        |        | 58     | 29    | 27    | 27     |       | 22      | 90    | 40    | 106   | 62    | 40    | 20    | 40    | 20    | 0 18  | ,        | 57        | 39    | 17   | 16        |            | 29    |       |
|                                  | 866   | 5800  | 3082  | 3939   | 2733  | 2351  | 1428  | 2851  | 3800   | 3432  | 6370  | 5218   | 3742   | 3999   | 541   | 8 6027  | 4249   | 7221   | 6322   | 4411  | 9526  | 1540   | 1594  | 1621    | 4452  | 4018  | 1444  | 1476  | 860   | 920   | 1282  | 1507  | 1469  | 861      | 747       | 1078  | 2139   | 1267      | 669        | 2425  | 288   |
| Gymnodinioid                     |       |       |       |        |       |       |       |       |        |       |       |        |        |        |       |         |        |        |        |       |       |        |       |         |       |       |       |       |       |       |       |       |       | 1        |           | , — † | $\square$  |           |            |       | -     |
| Pheopolykrikos hartmannii        |       |       |       |        |       |       |       |       |        |       |       |        |        |        |       |         |        |        |        |       |       |        |       |         |       |       |       |       |       |       |       |       | 1     | <u> </u> | 1         |       |  |           |            |       |       |
| Polykrikos kofoidii              | 108   | 150   | 517   | 121    | 133   | 117   | 28    | 3 29  | 200    | 216   | 85    | 218    | 57     | 29     | 19    | 3       | 55     | 55     | 176    | 58    | 84    | 81     | 119   | 111     | 159   | 40    | 63    | 166   | 180   | 160   | 163   | 265   | 182   | 137      | 173       | 196   | 142  | 101       | 106        | 43    | 160   |
| Polykrikos schwartzii            | 95    | 550   | 1897  | 91     | 233   | 352   | 400   | 205   | 229    | 568   | 371   | 562    | 457    | 117    | 32    | 2 235   | 305    | 277    | 470    | 382   | 333   | 162    | 95    | 5 222   | 590   | 285   | 191   | 270   | 260   | 200   | 163   | 183   | 182   | 137      | 192       | 98    | 178  | 118       | 95         | 161   | 148   |
|                                  | 203   | 700   | 2414  | 212    | 366   | 469   | 428   | 234   | 429    | 784   | 456   | 780    | 514    | 146    | 5 51  | 5 235   | 360    | 332    | 646    | 440   | 417   | 243    | 214   | 333     | 749   | 325   | 254   | 436   | 440   | 360   | 326   | 448   | 364   | 274      | 365       | 294   | 320  | 219       | 201        | 204   | 314   |
| Peridinioid                      |       |       |       |        |       |       |       |       |        |       |       |        |        |        |       |         |        |        |        |       |       |        |       |         |       |       |       |       |       |       |       |       |       | <u> </u> |           |       |  |           | <u>t t</u> |       |       |
| Protoperidinium conicum          | 121   | 200   | 448   | 242    | 233   | 117   | 228   | 323   | 200    | 270   | 428   | 187    | 371    | 112    | 354   | 4 352   | 194    | 83     | 264    | 264   | 111   | 189    | 190   | 311     | 227   | 122   | 191   | 395   | 320   | 300   | 428   | 693   | 255   | 137      | 76        | 157   | 250  | 186       | 122        |       | 54    |
| Protoperidinium leonis           | 81    | 3850  | 1966  | 1333   | 1600  | 2147  | 1342  | 1323  | 1314   | 1432  | 2628  | 2000   | 4114   | 2323   | 209   | 5 1470  | 1666   | 1500   | 3000   | 4705  | 1694  | 1648   | 952   | 644     | 409   | 510   | 680   | 937   | 520   | 600   | 469   | 448   | 527   | 172      | 346       | 275   | 196  | 271       | 81         | 132   | 148   |
| Protoperidinium obtusum          |       |       | 103   | 121    | 133   | 58    |       |       |        | 27    | 85    |        |        | 176    | 5 19  | 3 205   |        |        | 58     | 29    | 111   | 81     | 47    | 7       |       |       |       |       | 20    | 40    |       | 20    | 36    | 51       | 19        | 39    | 17   | $ \frown$ |            | 147   | 18    |
| Protoperidinium latissinum       |       |       | 34    |        |       |       |       | 29    |        |       | 28    |        |        |        |       | 29      | )      |        |        |       |       |        |       |         |       |       | 21    |       | 20    |       |       |       |       |          | 19        |       |  | $ \neg $  |            |       |       |
| Protoperidinium claudicans       | 40    | 400   | 172   | 91     | 100   | 88    | 142   | 147   | 143    | 216   | 285   | 187    | 57     | 205    | 5 129 | 9 29    | 138    | 138    | 147    | 117   | 83    | 27     | 71    | 200     | 204   | 244   | 170   | 291   | 100   | 200   | 387   | 489   | 273   | 155      | 153       | 255   | 178  | 135       | 41         |       | 129   |
| Votadinium calvum                |       |       | 34    |        |       | 29    | 1     |       |        |       | 28    |        |        | 58     | 3 6   | 4       | 27     | 27     |        | 88    | 27    |        | 47    | 7       | 68    | 20    | 42    | 62    | 40    | 60    | 20    | 20    | ) 73  | 34       | 19        | 20    |  | 16        |            | 102   | 31    |
| Protoperidinium latidorsale      |       |       |       |        |       |       |       |       |        |       |       |        |        |        |       |         |        |        |        |       |       |        |       |         |       |       |       |       |       |       |       |       |       | 1        |           |       |  | (         |            | 29    |       |
| Protoperidinium inaequale        | 13    |       |       | 30     |       |       |       |       | 114    | 27    |       |        | 28     |        |       |         | 55     |        | 29     | 29    | 27    | 54     | 23    | 5       | 22    |       | 21    |       |       | 40    |       |       |       | 1        |           | 20    |  | 33        | 14         |       |       |
| Protoperidinium steidingerae     |       |       |       |        |       |       |       | 88    |        |       |       |        |        |        |       |         |        |        |        |       |       |        |       |         |       |       |       |       |       |       |       |       |       | 1        |           |       |  | (         |            |       |       |
| Protoperidinium symmetricum      | 13    | 200   | 103   | 121    | 66    |       | 57    | 58    | 29     | 27    | 171   | 62     |        |        |       | 29      | )      |        | 29     |       | 55    |        | 23    | 5       |       | 81    | 85    |       |       | 80    | 40    | )     | 36    | 86       | 19        | 39    |  | 16        | 14         |       |       |
| Protoperidinium subinerme        | 27    | 100   |       | 30     |       | 58    |       | 58    | 57     | 27    |       |        | 28     | 58     | 3 32  | 2 58    | 55     |        | 88     |       | 27    |        |       | 111     | 68    | 40    | 42    | 20    | 60    | 20    | 20    | )     | 90    | 17       | 19        | 59    | 71   | 33        | 27         | 14    | 31    |
| Protoperidinium americatum       |       | 150   | 138   | 91     | 133   | 117   | 28    | 88    | 57     | 27    | 171   | 62     | 57     | 29     | 6     | 4 88    | 3 111  | 83     | 29     | 117   | 166   | 27     | 47    | 44      | 113   | 102   | 85    | 41    | 60    | 120   | 40    | 40    | 145   | 17       | 57        | 59    | 35   | 67        |            | 44    |       |
| Stelladinium stellatum           |       |       |       |        |       |       |       |       |        | 27    |       | 31     |        |        |       |         | 27     |        |        |       |       |        |       | 1       |       |       |       |       |       |       |       |       |       |          |           |       |  |           |            | 14    |       |
| Trinovantedinium applanatum      | 13    |       |       |        | 33    |       |       |       | 29     |       | 28    |        |        |        |       | 29      | )      |        |        |       |       | 27     | 23    | 5       |       |       | 21    | 20    |       |       | 20    | 40    | 55    | . 17     |           | ,,    | 17   | - T       | 13         |       | 18    |
| Bri gantedinium spp.             | 283   | 3200  | 2069  | 2394   | 3300  | 3382  | 2400  | 2588  | 2000   | 1811  | 2571  | 3250   | 2742   | 1088   | 3 100 | 205     | 972    | 416    | 794    | 794   | 1472  | 648    | 619   | 1266    | 863   | 1040  | 936   | 937   | 720   | 860   | 653   | 632   | 2 709 | 655      | 403       | 588   | 446  | 661       | 284        | 29    | 462   |
| Bri gantedinium simplex          | 635   | 2300  | 1759  | 1636   | 1900  | 1470  | 2342  | 1705  | 1600   | 1541  | 1057  | 1093   | 1028   | 2000   | 296   | 7 3294  | 2083   | 2888   | 2294   | 2088  | 3611  | 3405   | 2214  | 1933    | 1522  | 1693  | 1723  | 1812  | 1100  | 1340  | 918   | 1142  | 927   | 758      | 750       | 1431  | 678  | 949       | 635        | 191   | 444   |
| Brigantedinium asymmetricum      | 324   | 350   | 345   | 242    | 233   | 411   | 314   | 294   | 257    | 270   | 371   | 312    | 257    | 705    | 5 290 | 0 764   | 916    | 1138   | 852    | 1029  | 611   | 918    | 666   | 866     | 1000  | 857   | 1148  | 875   | 660   | 900   | 448   | 755   | 636   | 482      | 557       | 902   | 464  | 474       | 324        | 250   | 333   |
| Brigantedinium majusculum        |       |       | 69    | 61     |       |       |       | 29    | 86     | 54    | 171   | 250    | 28     |        | 354   | 4 147   | 27     | 166    |        | 29    | 27    |        |       |         |       |       |       | 41    | 20    |       | 20    | )     |       | 17       |           | 20    | 17   | - T       |            | 88    | 18    |
| Brigantedinium irregulare        | 37    |       | 34    | 30     |       |       |       | 58    | 29     | 108   | 57    | 125    | 28     |        | 6     | 4 29    | 55     | 27     |        | 29    | 83    | 27     |       |         | 68    | 20    |       | 20    |       | 20    |       |       | 18    |          |           | 20    | 17   |           |            | 29    | 31    |
| Dubridinium caperatum            | 162   | 550   | 207   | 152    | 333   | 470   | 200   | 235   | 286    | 297   | 571   | 343    | 371    | 500    | 54    | 8 529   | 444    | 555    | 352    | 352   | 55    | 81     | 428   | 800     | 522   | 224   | 361   | 354   | 320   | 360   | 346   | 346   | 309   | 241      | 230       | 353   | 267  | 237       | 162        | 117   | 3'    |
| Oblea acanthocysta               | 14    | 650   | 483   | 424    | 466   | 411   | 257   | 676   | 457    | 378   | 485   | 750    | 857    | 235    | 5 774 | 4 1029  | 500    | 888    | 411    | 441   | 1416  | 432    | 380   | 533     | 590   | 551   | 382   | 375   | 180   | 300   | 204   | 224   | 200   | 103      | 115       | 235   | 89   | 101       | 14         | 14    | 18    |
| Scrippsiella spp.                | 27    |       |       |        |       |       | 28    | 6     |        | 27    |       |        |        |        |       |         |        |        | 58     | 29    | 20    | 54     |       | 44      | 363   | 122   | 276   | 83    | 280   | 140   | 224   | 204   | 145   | 120      | 250       | 275   | 196  | 101       | 27         | 14    | 20    |
|                                  | 1790  | 11950 | 7966  | 7000   | 8530  | 8758  | 7338  | 7699  | 6657   | 6568  | 9135  | 8652   | 9966   | 7494   | 8929  | 9 8286  | 7270   | 7909   | 8405   | 10140 | 9596  | 7618   | 5730  | 6752    | 6039  | 5626  | 6184  | 6263  | 4420  | 5380  | 4237  | 5053  | 4435  | 3062     | 3032      | 4745  | 2938   | 3280      | 1757       | 1214  | 1994  |
| Total                            | 2887  | 62250 | 94634 | 126969 | 49849 | 91724 | 47479 | 92254 | 120914 | 50297 | 89388 | 169930 | 124107 | 152932 | 6908  | 7 92724 | 127406 | 113183 | 101519 | 75725 | 91622 | 131509 | 39656 | 33238   | 14443 | 15316 | 9838  | 14924 | 5940  | 7280  | 6701  | 7293  | 6558  | 4300     | 4258      | 6431  | 5593   | 4833      | 2654       | 3960  | 5268  |