

Reviewer #1

The manuscript is much improved by the changes made. It is concise, logical, and clearly written. Well done! A few minor suggestions and changes:

Title: The title would be more direct as, 'Why is Trichodesmium abundant...'

We have changed as suggested.

Text:

Line 52: Change 'generally' to 'thought to be'

We have changed as suggested. (L51)

Lines 59-61: What is limited? I think a word is missing.

We have revised the sentence as follows. (L58-60)

“As for phosphorus limitation, iron-enhanced nitrogen fixation causes phosphorus depletion, and the nitrogen fixation is consequently limited by phosphorus (Mather et al., 2008).”

Line 65: Change 'condition' to 'nutrient and trace metal concentrations'

We have changed as suggested. (L64-65)

Line 69: Please give the island names, so that it is clearer how this study differs from previous work.

We have added the island names in the revised manuscript. (L69-70)

Line 88-89: Change 'as summer' to 'summer as'

We have changed as suggested. (L91)

Line 183: Change 'is known' to 'is thought'

We have changed as suggested. (L185)

Line 187: Change 'would' to 'could be higher'

We have changed as suggested. (L189)

Line 194: Please give the incubation time for the tests.

In the experiment1, $^{15}\text{N}_2$ gas was injected as bubble form, and the bottle was incubated for at least 12 h to reach equilibrium. Meanwhile, in the experiment2, $^{15}\text{N}_2$ dissolved water was made, and thus, the bottle was not incubated. We wrote the detail in the Supplementary Material.

Fig. 3: The horizontal axis labels should be below the graphs

We have changed as suggested.

Fig. 6b: Typo in horizontal axis label.

We have rewritten the label in horizontal axis.

SI:

Line 22: The units and Dabundo reference are missing.

We have added the units. (L22) The Dabundo et al's paper was referred at L7.

Line 28: If it is significant, please add the stats/P-value.

We have added the result of statistics at L29.

Fig. S1: Please describe error bars.

The Error bars indicate standard deviations of the mean (n=5). We have added this sentence in the figure legend.

Fig. S2: Please add the region this figure shows.

We have added the region in this figure.

Fig S3: Would this be clearer on a log scale?

We have changed to a log scale as suggested.

Fig. S4: Please label the x-axis.

We have labeled the x-axis.

Fig. S5: Please label or identify area K.

We have labeled area K in the revised figure.

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Reviewer #2

I think the manuscript reads more clearly now and the authors have done a good job of addressing reviewers' comments. I suggest a few minor edits:

*line 19: The genus *Trichodesmium* is recognized as an abundant (add the word "an")*

We have changed as suggested. (L18)

line 43: oligotrophic conditions (add "s")

We have changed as suggested. (L42)

line 557: conditions (add "s") and phosphate at the nanomolar level (add "the")

We have changed the sentence as follows. (L63-67)

“Therefore, the distinct high abundance of *Trichodesmium* in the Kuroshio is probably not explained by nutrient and trace metal concentrations; however, distributions of dissolved iron and phosphate at the nanomolar level have not been well studied in this region (Obata et al., 1997; Shiozaki et al., 2010; Kodama et al., 2011).”

section 3.2: Perhaps the authors could report the "r squared" value for all of the correlations you report, not just the one on lines 837-838.

We have added the r squared values at L295 and 307.

1 | **Why ~~does is~~ *Trichodesmium* ~~become~~ abundant in the**
2 **Kuroshio?**

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18 **Abstract**

19 | The genus *Trichodesmium* is recognized as an abundant and major diazotroph in the
20 Kuroshio, but the reason for this remains unclear. The present study investigated the
21 abundance of *Trichodesmium* spp. and nitrogen fixation together with concentrations
22 of dissolved iron and phosphate in the Kuroshio and its marginal seas. We performed
23 the observations near the Miyako Islands, which form part of the Ryukyu Islands,
24 situated along the Kuroshio, since our satellite analysis suggested that material
25 transport could occur from the islands to the Kuroshio. *Trichodesmium* spp. bloomed
26 ($>20,000$ filaments L^{-1}) near the Miyako Islands, abundance was high in the Kuroshio
27 and the Kuroshio bifurcation region of the East China Sea, but was low in the
28 Philippine Sea. The abundance of *Trichodesmium* spp. was significantly correlated
29 with the total nitrogen fixation activity. The surface concentrations of dissolved iron
30 (0.19–0.89 nM) and phosphate (<3–36 nM) were similar for all of the study areas,
31 indicating that the nutrient distribution could not explain the spatial differences in
32 *Trichodesmium* spp. abundance and nitrogen fixation. Numerical particle-tracking
33 experiments simulated the transportation of water around the Ryukyu Islands to the
34 Kuroshio. Our results indicate that *Trichodesmium* growing around the Ryukyu
35 Islands could be advected into the Kuroshio.

37 1. Introduction

38 The Kuroshio is a western boundary current in the North Pacific Ocean that
39 originates in the North Equatorial Current and bifurcates to the east of the Philippines.
40 The main stream of the Kuroshio enters the East China Sea (ECS) northeast of Taiwan,
41 flows out through the Tokara Strait, and runs along the Japanese islands of Shikoku
42 and Honshu. While the Kuroshio and its adjacent waters are characterized by highly
43 oligotrophic conditions, phytoplankton and zooplankton communities in the Kuroshio
44 are distinct compared to those from adjacent waters (McGowan, 1971). McGowan
45 (1971) suggested that some plankton species are delivered by the Kuroshio to the
46 north from the equatorial region.

47 The abundance of the cyanobacterial genus *Trichodesmium* in the Kuroshio is
48 much higher than that in neighboring seas (Marumo and Asaoka, 1974). Because
49 *Trichodesmium* is a major nitrogen fixer in the Kuroshio, it is believed to be the key
50 genus for understanding the Kuroshio ecosystem (Chen et al., 2008, 2014; Shiozaki et
51 al., 2014a). Nevertheless, the factors controlling the distribution of *Trichodesmium* in
52 this region are poorly understood. Marine nitrogen fixation is generally thought to be
53 regulated by the supply of iron and phosphorus (Mahaffey et al., 2005), and
54 *Trichodesmium* thrives in iron-rich oligotrophic regions (Moore et al., 2009; Shiozaki

55 et al., 2010, 2014b). A major source of iron in the ocean is atmospheric dust
56 deposition (Jickells et al., 2005; Mahowald et al., 2009). Modeling studies indicate
57 that dust deposition in the western North Pacific decreases exponentially from the
58 continental shelf to the Philippine Sea (Jickells et al., 2005; Mahowald et al., 2009),
59 and hence, deposition is not as high in the Kuroshio as in the adjacent waters. As for
60 phosphorus limitation, iron-enhanced nitrogen fixation causes phosphorus depletion,
61 and the nitrogen fixation is consequently limited by phosphorus (Mather et al., 2008).

62 The phosphate distribution has been examined in this study region using a
63 conventional colorimetric method, and the surface phosphate concentration in the
64 Kuroshio has been reported to be as low as that in the Philippine Sea (Chen, 2008).

65 Therefore, the distinct high abundance of *Trichodesmium* in the Kuroshio is probably
66 not explained by nutrient ~~condition~~ and trace metal concentrations; however,
67 distributions of dissolved iron and phosphate at the nanomolar level have not been
68 well studied in this region (Obata et al., 1997; Shiozaki et al., 2010; Kodama et al.,
69 2011).

70 Nitrogen fixation by *Trichodesmium* has recently also been found to be active
71 around oceanic islands; New Caledonia Island, Efate Island, Fiji Island, Tahiti Island,
72 and Northern Mariana Islands (Shiozaki et al., 2010, 2013, 2014c; Lin et al., 2011).

73 Furthermore, these studies demonstrated that abundant *Trichodesmium* is delivered by
74 the current to areas that are remote from the islands. Although this phenomenon was
75 noted in the western Pacific warm pool and western South Pacific, it can also occur in
76 and around the Kuroshio and may contribute to the distribution of *Trichodesmium* in
77 this region.

78 In the present study, we simultaneously determined *Trichodesmium* abundance
79 and bulk water nitrogen fixation together with concentrations of dissolved iron and
80 phosphate at the nanomolar level in the Kuroshio and its marginal seas. In addition,
81 we conducted intensive observations around the Miyako Islands section of the
82 Ryukyu Islands located close to the main stream of the Kuroshio.

83

84 **2. Materials and Methods**

85 **2.1. Oceanographic database**

86 Algal blooms in an oligotrophic region may indicate a nitrogen fixation hotspot
87 (Wilson and Qiu, 2008; Shiozaki et al., 2014c). To identify the locations of intensive
88 algal blooms, we used a dataset of chlorophyll (chl) *a* observed by satellite. According
89 to Wilson and Qiu (2008), an algal bloom in an oligotrophic region can be defined as
90 a chl *a* value $>0.15 \text{ mg m}^{-3}$ in summer. In the present study, we used 8-day,

91 moderate-resolution imaging spectroradiometer (MODIS) level 3 chl *a* with 9 km
92 resolution during summer between July 2003 and September 2009. We defined ~~as~~
93 summer as July through September. The bloom frequency for each pixel was
94 calculated from the ratio of counts in which chl *a* was $>0.15 \text{ mg m}^{-3}$ to the total
95 counts in which chl *a* was detected.

96 To examine the current field, geoelectrokinetograph and ship-mounted acoustic
97 Doppler current profiler (ADCP) data from the uppermost layer for the summers
98 between 1953 and 2008 were obtained from the Japan Oceanographic Data Center
99 (<http://www.jodc.go.jp>). Regridding, removal of anomalous values, and smoothing of
100 the dataset were performed as described by Isobe (2008).

101

102 2.2. Cruise observations

103 Experiments were conducted during summer on-board the R/V *Tansei-maru*
104 (KT-06-21, September 9–17, 2006; KT-07-22, September 5–13, 2007; KT-09-17,
105 September 8–13, 2009; KT-10-19, September 4–12, 2010) and the T/V
106 *Nagasaki-maru* (242, July 19–28, 2007) (Fig. 1a, Table S1). The stations during the
107 KT-06-21, KT-07-22, and *Nagasaki-maru* 242 cruises were divided into three areas
108 based on the temperature-salinity diagram (see Fig.2 of Shiozaki et al., 2011): the

109 ECS, Kuroshio, and Philippine Sea. During the KT-09-17 cruise, we conducted
110 experiments around the Miyako Islands which were distinguished from the other three
111 areas. During the KT-10-19 cruise, we performed observations in the ECS, the
112 Kuroshio, and around the Miyako Islands (Liu et al., 2013).

113

114 **2.2.1. Light intensity, hydrography, nutrients, and chl a**

115 Water samples for all of the experiments, with the exception of determination of
116 the dissolved iron concentration, were collected using an acid-cleaned bucket and
117 Niskin-X bottles. The depth profile of light intensity was determined immediately
118 before the water sampling using a light sensor (during the KT-06-22, KT-07-21,
119 KT-09-17, and KT-10-19 cruises) or an empirical equation (during the *Nagasaki-maru*
120 242 cruise) (Shiozaki et al., 2011). Temperature and salinity profiles to a depth of 200
121 m were obtained using a conductivity, temperature, and depth (CTD) sensor. Mixed
122 layer depth (MLD) was defined as the depth at which the sigma-t increased by 0.125
123 from its value at a depth of 10 m. Water samples for nitrate+nitrite (N+N) and
124 phosphate were collected from 0, 10, 30, 40, 50, 60, 70, 80, 90, 100, 125, 150, and
125 200 m, and from depths at given light intensities. At all of the stations, the N+N and
126 phosphate concentrations were determined at the nanomolar level using a

127 supersensitive colorimetric system consisting of an AutoAnalyzer II (Technicon) and
128 Liquid Waveguide Capillary Cells (World Precision Instruments, USA) (Hashihama et
129 al., 2009). The detection limits of N+N and phosphate were both 3 nM. When the
130 concentration was greater than 0.1 μM , it was determined by conventional methods
131 using a TRAACS 2000 autoanalyzer (Bran:Luebbe, UK). In addition to the
132 observations at the stations, temperature, salinity, and the *in vivo* chl fluorescence of
133 the surface water were monitored continuously during the cruises by a
134 thermosalinograph (Ocean Seven, Idronaut, Italy) and a fluorometer (Minitracka,
135 Chelsea, UK).

136

137 2.2.2. Dissolved iron

138 Water was sampled to estimate the dissolved iron concentration from 0.5-m depth
139 during the KT-06-21 and KT-07-22 cruises and from 10-m depth during the KT-09-17
140 cruise using an acid-cleaned Teflon bellows pump (AstiPure PFD2; Saint-Gobain)
141 with Teflon tubing (inner diameter = 12 mm). The water was filtered through an
142 acid-cleaned 0.22 μm pore filter (Millipak100; Millipore) connected to the in-line of
143 the Teflon tubing with a Teflon connector. Filtered seawater was collected in a 125
144 mL low-density polyethylene (LDPE) bottle (Nalgene, Nalge Nunc International),

145 which had been washed using following technique: the sample bottles were
146 sequentially cleaned by soaking in 5% alkali detergent for at least 2 days, in 4 N HCl
147 for at least 1 day, in 0.3 N metal analysis-grade HNO₃ at 60°C overnight, and finally,
148 in Milli-Q water at 60°C overnight. After rinsing with Milli-Q water, the bottles were
149 dried in a laminar flow space and stored in double plastic bags. The filtrate samples
150 were acidified to a pH <1.7 with trace-metal-grade HCl (Tamapure AA-100; Tama
151 Chemicals) in a Class-100 clean-air bench, and stored at room temperature for more
152 than 1 year.

153 The dissolved iron concentration was determined using an automatic Fe(III) flow
154 injection analytical system (Kimoto Electric Co., Ltd.) using a chelating resin
155 pre-concentration and chemiluminescence detection method (Obata et al., 1993). A
156 buffer solution of 10 M formic acid and 2.4 M ammonium formate was added to the
157 samples. The sample pH was adjusted to 3.0 with 20% ammonium hydroxide
158 (NH₄OH; Tamapure AA-10; Tama Chemicals) immediately prior to analysis. The
159 detection limit of this method was 0.05 nM. The SAFe reference standards S1 and D2
160 were measured during the course of sample analysis, and the results were within the
161 range of the published consensus values: S1 = 0.097 ± 0.043 nM and D2 = 0.91 ±
162 0.17 nM (Johnson et al., 2007).

163

164 **2.2.3. Nitrogen fixation and abundance of *Trichodesmium* spp.**

165 Samples for the incubation experiments were collected vertically at all of the
166 stations, except at Sts. T0621, GN-3, and T0905, where samples were only collected
167 from the surface. All samples were collected in duplicate in acid-cleaned 4.5-L
168 polycarbonate bottles. During the *Nagasaki-maru* 242 cruise, water samples were
169 collected from four different depths corresponding to 100%, 25%, 10%, and 1% of the
170 surface light intensity. During the other cruises, samples were collected from a depth
171 of 50% surface light intensity. Samples at 100% surface light intensity were collected
172 from 0 m during all of the cruises, except during the KT-10-19 cruise in which the
173 samples were collected from a depth of 5 m. The bulk water nitrogen fixation activity
174 was determined based on primary production using a dual isotopic ($^{15}\text{N}_2$ and ^{13}C)
175 technique (Shiozaki et al., 2009). After ^{13}C -labeled sodium bicarbonate (99 atom%
176 ^{13}C ; Cambridge Isotope Laboratories) was added to each bottle, 2 mL of $^{15}\text{N}_2$ gas (98
177 + atom% ^{15}N ; SI Science Co. Japan) was injected directly into the incubation bottles
178 through a septum using a gastight syringe. The bottles were covered with
179 neutral-density screens to adjust the light level and incubated for 24 h in an on-deck
180 incubator cooled by flowing surface seawater for 24 h. We determined the nitrogen

181 fixation activity using the $^{15}\text{N}_2$ gas bubble addition method (Montoya et al., 1996).
182 This method is believed to underestimate the nitrogen fixation rate relative to the $^{15}\text{N}_2$
183 gas dissolution method (Mohr et al., 2010). The start time of incubation in this study
184 varied at each station (Table S1). Considering daily periodicity of nitrogen fixation in
185 each diazotroph (Zehr, 2011) and the time to reach equilibration of the $^{15}\text{N}_2$ gas
186 bubble with seawater (>12 h, Mohr et al., 2010), the level of underestimation could
187 vary at each station. Meanwhile, the level of underestimation is ~~known~~thought to be
188 low in *Trichodesmium* dominant water because *Trichodesmium* can float to the top of
189 the bottle and directly use the added $^{15}\text{N}_2$ in the bubble method (Großkopf et al.,
190 2012). Although the bias of underestimation could not be estimated from the results in
191 this study, the actual nitrogen fixation rate ~~would~~could be higher than the obtained
192 rate.

193 A recent study demonstrated that commercial $^{15}\text{N}_2$ gas could be contaminated by
194 ^{15}N -labeled nitrate and ammonium (Dabundo et al., 2014). We tested the
195 contamination in $^{15}\text{N}_2$ gas produced by SI Science Co., Ltd., which was used (from
196 different batch numbers) in the present study (see Supporting Information). Briefly,
197 the $^{15}\text{N}_2$ gas was dissolved in aged subtropical surface water, and concentrations of
198 nitrate, nitrite, and ammonium at the nanomolar levels were determined using

199 supersensitive colorimetric systems. The results showed that there were no significant
200 differences between the control and samples to which $^{15}\text{N}_2$ had been added (Fig. S1),
201 suggesting that the contamination of nitrate, nitrite, and ammonium in the $^{15}\text{N}_2$ gas
202 was insignificant (Supporting Information).

203 Water samples were collected for microscopic analysis at all light depths during
204 the *Nagasaki-maru* 242 and KT-07-21 cruises, and only from the surface during the
205 KT-06-22, KT-09-17, and KT-10-19 cruises. The samples were fixed using acidified
206 Lugol's solution. *Trichodesmium* spp. were counted using the Utermöhl method under
207 inverted microscope observation. *Trichodesmium* greater than ca. 300 μm in length
208 were counted as 1 filament and shorter lengths were counted as 0.5 filaments. In
209 addition, phytoplankton other than *Trichodesmium* spp. were identified from the
210 samples obtained during the KT-09-17 cruise.

211

212 2.3. Statistical analysis of environmental variables

213 We used non-metric multi-dimensional scaling (nMDS) to investigate the spatial
214 differences in the environmental variables that could influence *Trichodesmium* growth
215 and bulk water nitrogen fixation; temperature, mixed layer depth, nitrate, dissolved
216 iron, and phosphate. The environmental variables were transformed by $\log_{10}(x + 1)$

217 prior to analysis. A dissimilarity/similarity matrix between stations was constructed
218 using the Bray-Curtis index. The nMDS was used to visualize similarities in the
219 environmental variables among the stations. An Analysis of Similarity (ANOSIM)
220 was used to test the differences in the environmental variables among the stations.

221

222 2.4. Numerical experiments

223 Numerical particle-tracking experiments were conducted to investigate the
224 transport of water masses at the surface from areas around the Miyako Islands in the
225 summer season from 2003 to 2009. Surface velocity data were derived from the
226 FRA-JCOPE2 reanalysis product (Miyazawa et al., 2009), which is an eddy-resolving
227 ($1/12^\circ$) ocean model combined with three-dimensional variational data assimilation
228 (satellites, ARGO floats, and shipboard observations), and is one of the most reliable
229 models for the region around Japan for the above time period. The method of tracking
230 particles was basically the same as in Itoh et al. (2009), but we did not include the
231 random walk for simplicity. The release points of particles were selected at the surface
232 of the model grid points around the coastal waters of the Miyako Islands. We assumed
233 that the particles did not increase, die, or sink from the surface during the experiments.
234 To focus on transport during the summer season (July–September), particles were

235 released one month before the summer (June 1) and were tracked until September 30.

236 To examine differences in the output depending on the start time within the same
237 year, we also performed experiments starting on June 1, 11, and 21, and July 1 in
238 2009. The ratio of particles that reached areas downstream of the Tokara Strait
239 (hereafter Area K) (Fig. 7), including the particles' entrainment to the Kuroshio, to
240 total particles released from the Miyako Islands was computed in all experiments. It
241 should be noted that these experiments contained the following two uncertainties.
242 First, the distribution of *Trichodesmium* around the islands, which strongly influences
243 the destinations of particles, was not able to be determined in advance.
244 *Trichodesmium* is known to aggregate and not to occur uniformly in the ocean
245 (Capone et al., 1997). Second, the model cannot reproduce the current very close to
246 the islands. If a water mass very near the islands was delivered to the open ocean by
247 tide and/or river plumes that were not considered in the model, seaward dispersion of
248 particles was likely underestimated.

249

250 **3. RESULTS**

251 **3.1. The Kuroshio path and bloom frequency**

252 The average surface current field indicated that the main stream of the Kuroshio

253 flowed along the continental shelf in the ECS, and then passed to the south of the
254 Kyushu and Shikoku Islands (Fig. 1b). In addition, the Kuroshio branch bifurcated
255 northward at 25°N and 30°N at the continental shelf. Hence, all of the stations in the
256 ECS were subject to the influence of the Kuroshio. While the northeastward stream of
257 the Kuroshio was prominent in this region, smaller-scale flows and circulations were
258 observed in the areas around and to the southeast of the Ryukyu Islands. In the west of
259 the main stream of the Kuroshio, because the average chl *a* was over 0.15 mg m⁻³ (Fig.
260 S2), the frequency of chl *a* values >0.15 mg m⁻³ was high (Fig. 1b). In contrast, the
261 bloom frequency in the east of the main stream of the Kuroshio differed from the
262 distribution of the average chl *a*; algal blooms occurred frequently in the Ryukyu
263 Islands. Around the Miyako Islands, water of high bloom frequency was located to the
264 west of the islands, extending to the north.

265

266 3.2. Region-wide environmental conditions, *Trichodesmium* spp., and 267 nitrogen fixation

268 The sea surface temperature (SST) ranged from 25.1–30.5°C at all of the stations
269 (Table S1), and there were no significant differences among the areas ($p>0.05$,
270 Tukey's honestly significant difference [HSD] test). The MLD varied from 12–60 m

271 at all of the stations, and was relatively deep around the Miyako Islands compared to
272 the other areas (Table S1). The surface N+N concentration varied between <3 and 42
273 nM, except around the Miyako Islands (Shiozaki et al., 2010, 2011) (Table S1). The
274 highest surface N+N concentration (374 nM) was observed at St. T0904 where
275 upwelling occurred (see below). No significant difference in the surface N+N was
276 observed among the four areas ($p > 0.05$, Tukey's HSD test). The surface phosphate
277 concentration varied between <3 and 36 nM at all of the stations (Fig. 2a). The
278 phosphate concentration at the surface and within the MLD was not significantly
279 different among the four areas ($p > 0.05$, Tukey's HSD test). There was a greater
280 increase in the phosphate concentrations below 40–50 m in the ECS compared to the
281 other areas (Fig. 3a–d). Furthermore, the phosphate concentrations below 40–50 m
282 near the Miyako Islands were higher than those in the Kuroshio and the Philippine
283 Sea, which were depleted down to 100 m, except at St. T1004 located near the
284 continental shelf. The N/P (= N+N/phosphate) ratio at the surface varied from 0.28 to
285 6.40 except at St. T0904 (N/P = 16.3) (Table S1), and no significant differences were
286 observed among the four areas ($p > 0.05$, Tukey's HSD test). The surface dissolved
287 iron concentration ranged from 0.19 to 0.89 nM at all of the stations (Fig 2b), with no
288 significant spatial differences among the four areas ($p > 0.05$, Tukey's HSD test). The

289 surface dissolved iron concentration at Sts. T0622 and T0907 was elevated to 0.83 nM
290 and 0.89 nM, respectively, with lower salinity water than in the adjacent waters
291 (salinity data are shown in Fig. 4a and Kodama et al., 2011). The nMDS showed that
292 the environmental variables at all stations were the same at the >80% similarity level
293 and were >90 % similar excepting station T0904 (Fig. 5). The ANOSIM indicated no
294 significant differences among the stations ($p > 0.05$).

295 The abundance of *Trichodesmium* spp. was highest at the surface at almost all of
296 the stations during the *Nagasaki-maru* 242 and KT-07-21 cruises (Fig. S3). The
297 surface *Trichodesmium* spp. abundances were positively correlated with the
298 depth-integrated abundances ($r^2 = 0.51$, $p < 0.05$, ~~t -test~~) (Fig. 6a). Thus, the surface
299 abundance was used to discuss the geographical distribution of *Trichodesmium* spp.
300 The *Trichodesmium* spp. abundance at the surface varied widely, and there was no
301 significant difference among the four areas ($p > 0.05$, Tukey's HSD test).
302 *Trichodesmium* spp. were observed at all of the stations in the Kuroshio and around
303 the Miyako Islands, whereas they were not always observed in the ECS and the
304 Philippine Sea (Fig. 2c). The average surface abundance in the Philippine Sea was the
305 lowest among all of the areas (Table 1). The highest abundance of *Trichodesmium* spp.
306 (>20000 filaments L⁻¹) was observed near the Miyako Islands at St. T0906, where

307 they bloomed (see below). Tuft-shaped colonies were found at Sts. T0706, T0723,
308 CK-10, and T0906. The nitrogen fixation rate was highest in the upper 25% light
309 depth, and decreased with increasing depth at all of the stations (Fig. 3e–h). The
310 surface rates were positively correlated with the depth-integrated rates ($r^2 = 0.79$, $p <$
311 0.05 , ~~t -test~~) (Fig. 6b), suggesting that the distribution of nitrogen fixation was indexed
312 by the surface activity. Surface and depth-integrated nitrogen fixation ranged from
313 0.54 to 62 nmol N L⁻¹ d⁻¹ and from 29.5 to 753 μmol N m⁻² d⁻¹, respectively (Fig. 2d
314 and Table S1). Surface nitrogen fixation in the Philippine Sea was significantly lower
315 than that in the Kuroshio ($p < 0.05$, t -test).

316 The surface abundance of *Trichodesmium* spp. in the entire study area was
317 positively correlated with the nitrogen fixation rate at the surface ($r^2 = 0.80$; $p < 0.05$
318 [$r^2 = 0.525$; $p < 0.05$ if the datum taken at the *Trichodesmium*-bloom station T0906 is
319 excluded]) (Fig. 6c), suggesting that they significantly contributed to nitrogen fixation
320 in the study region. However, active nitrogen fixation occurred in the ECS where
321 *Trichodesmium* abundance was low, and hence, the other diazotrophs could also be
322 important for nitrogen fixation.

323

324 **3.3. Observation around the Miyako Islands during the KT-09-17 cruise**

325 The SST was lower to the northwest of the Miyako Islands than in adjacent
326 waters, and chl *a* was enriched in the same location (Fig. 4b,c). Therefore, the
327 enhanced productivity was probably due to nutrient supply by upwelling. This
328 upwelling generally occurs in the lee of islands (Hasegawa et al., 2009), suggesting
329 that there was a northward current during the cruise. The surface salinity was lower
330 east of the Miyako Islands than in the surrounding waters (Fig. 4a). The absence of
331 any large river on the east side of Miyako-jima Island and the separation of low
332 salinity water from the island suggest that the low salinity was caused by rainfall.
333 St. T0904 was located near the upwelling water; its SST of 29.0°C was lowest and its
334 surface N+N concentration of 374 nM was highest among all of the stations. However,
335 the N+N concentration at St. T0904 at the surface was higher than that at the
336 subsurface (an approximate depth of 50 m; Fig. S4), indicating that St. T0904 was not
337 located in the middle of the upwelling. At St. T0904, the surface phosphate
338 concentration was also highest (23 nM) and the N/P ratio (=16.3) was higher than the
339 Redfield ratio. With the exception of the surface at St. T0904, the phosphate
340 concentration was low (<3–9 nM) in the upper 50 m, with no noticeable variation
341 among the stations (Fig. 2a). The dissolved iron concentration varied between 0.19

342 and 0.89 nM at the surface (Fig. 2b). The highest dissolved iron concentration was
343 observed at St. T0907.

344 During the same cruise, we encountered a *Trichodesmium* spp.-bloom at St.
345 T0906 (Fig. 2c), which had colored water at the surface. The abundance of
346 *Trichodesmium* spp. at St. T0906 was >20,000 filaments L⁻¹, which was far higher
347 than that at other stations (2–102 filament L⁻¹). The nitrogen fixation rate at the
348 surface (61.9 nmol N L⁻¹ d⁻¹) of this station was more than 30-fold that just below the
349 surface, and was the highest among all of the stations (Fig. 3h). The diatom
350 abundance was markedly higher at St. T0904 than that at the other stations.
351 *Cylindrotheca closterium* was the most numerically dominant diatom (59%), followed
352 by *Navicula* spp. (23%) and *Nitzschia* spp. (13%). *C. closterium* was not detected at
353 the other stations, indicating that the high chl *a* induced by the island wake effect
354 mainly consisted of diatoms.

355

356 3.4. Numerical simulation

357 As the Kuroshio generally flows along the continental slope north of the Miyako
358 Islands (Fig. 1b), particles around the Miyako Islands were not transported along the
359 typical path of the Kuroshio to the northeast, especially at their initial stages (Fig. 7a).

360 Some particles migrated around the Miyako Islands, or turned south after they passed
361 the Tokara Strait. Nevertheless, the particles delivered to Area K east of the Tokara
362 Strait increased as time elapsed, and the ratio of particles delivered to Area K to the
363 total released particles ranged from 13–56% ($30 \pm 16\%$) by day 120 in 2003–2009
364 (Fig. 7b). The year-to-year variations in the ratio are mainly due to influences of
365 mesoscale eddies as partly seen in the particle trajectories in Fig. 7a, and likely
366 occurred over relatively short time scales (shorter than the seasonal time scale). This
367 is supported by another series of experiments in which particles were released on June
368 1, 11, and 21, and July 1 in 2009, which yielded ratios of 6.2–38% ($22 \pm 13\%$) by day
369 120 (Fig. S5).

370

371 **4. DISCUSSION**

372 **4.1. Distribution of phosphate and dissolved iron concentrations**

373 Phosphate concentrations were consistently low within the MLD in all of the
374 studied areas, and the maximum abundance of *Trichodesmium* spp. and total nitrogen
375 fixation activity generally occurred near the surface, suggesting that the phosphate
376 conditions for surface *Trichodesmium* spp. and other diazotrophs were similar among
377 all of the areas. Furthermore, with the exception of St. T1004 located near the

378 continental shelf, the vertical distribution of phosphate in the Kuroshio was analogous
379 to that in the Philippine Sea. Therefore, at least in the oceanic region of the two areas,
380 phosphate availability for *Trichodesmium* spp. and the other diazotrophs was similar
381 throughout the water column.

382 The surface distribution of the dissolved iron concentration demonstrated no
383 significant variation among the areas. The dissolved iron concentration (0.19–0.89
384 nM) was higher than that in the western North Pacific subtropical region (0.15–0.4
385 nM) (Brown et al., 2005). Obata et al. (1997) demonstrated that the vertical
386 distribution of the dissolved iron concentration in the ECS showed two peaks (at the
387 surface and in the deep water), suggesting that aerial dust significantly contributes to
388 the high dissolved iron concentration at the surface in all of our study areas. In
389 accordance with our results, previous modeling studies estimated the amount of dust
390 deposition to be similar in all four areas (Jickells et al., 2005; Mahowald et al., 2009).
391 Therefore, iron availability for *Trichodesmium* spp. and the other diazotrophs was also
392 likely similar across all of the study areas. Iron can be supplied from deep water to the
393 surface by mixing processes (Johnson et al., 1999). However, if this were the case, the
394 nitrate concentration would be expected to increase simultaneously at the surface
395 (Johnson et al., 1999), and we observed no noticeable elevation in N+N in any of the

396 areas, except at St. T0904. High concentrations of dissolved iron (>0.8 nM)
397 corresponded with low salinity at Sts. T0622 and T0907, suggesting that wet
398 deposition was an important process for iron supply. Dry deposition could also be
399 important since the iron-enriched water at Sts. T0601 and T0715 did not correspond
400 with low salinity.

401 Satellite data analysis indicated that there was a “pipeline” of material transport
402 from the Miyako Islands to the Kuroshio, and this was supported by numerical
403 simulations. According to the hypothesis of Marumo and Asaoka (1974), the growth
404 of *Trichodesmium* in the Kuroshio could be maintained by the supply of iron and
405 phosphorus from the islands situated along the Kuroshio, and the Miyako Islands
406 were considered a possible nutrient source to the Kuroshio. Hence, assuming this
407 hypothesis to be valid, the iron and phosphate concentrations near the Miyako Islands
408 (especially in our observed area) would be expected to be higher than those in the
409 other areas. However, we observed no significant difference in the iron and phosphate
410 concentrations among the four areas. This suggested that there was no detectable
411 washout of iron and phosphorus from the Miyako Islands during our observations, or
412 that diazotrophs and other phytoplankton exhausted the nutrient supply close to the
413 islands.

414

415 **4.2. Factors controlling the distributions of *Trichodesmium* spp. and**
416 **nitrogen fixation**

417 Although there was no statistically significant difference in *Trichodesmium* spp.
418 abundance among the study areas probably because the data were limited and the
419 variation was large, *Trichodesmium* spp. were always observed in the Kuroshio and
420 were abundant at most stations. Furthermore, at St.CK-10 in the East China Sea which
421 is in the Kuroshio branch current, a high abundance of *Trichodesmium* spp. was
422 observed. On the other hand, *Trichodesmium* spp. abundance in the Philippine Sea
423 tended to be lower than in the other areas. Such *Trichodesmium* distribution was also
424 reported in the previous study (Marumo and Asaoka, 1974). The present study also
425 showed lower surface nitrogen fixation in the Philippine Sea compared to that in the
426 Kuroshio ($p < 0.05$, *t*-test). Previous studies demonstrated that *Trichodesmium* spp.
427 flourished in some regions of the subtropical ocean where the iron levels were high
428 (Moore et al., 2009; Shiozaki et al., 2014b), which can be attributed to the high iron
429 requirement of *Trichodesmium* spp. for their growth compared to other diazotrophs
430 and non-diazotrophs (Kustka et al., 2003; Saito et al., 2011). Therefore, the
431 distribution of *Trichodesmium* spp. in the study area was expected to be associated

432 with the dissolved iron concentration at the surface. Furthermore, the iron-enhanced
433 active nitrogen fixation causes phosphorus depletion, and is consequently limited by
434 phosphorus (Mather et al., 2008). No significant differences in surface iron and
435 phosphate were observed among the study areas, which cannot explain the
436 distribution of *Trichodesmium* spp. and nitrogen fixation in the study region.

437 Johnson et al. (1999) reported that the iron supply increased around the
438 continental shelf because re-suspension from the bottom to the euphotic zone
439 becomes significant. However, in the continental shelf of the ECS, the abundance of
440 *Trichodesmium* spp. and nitrogen fixation were low (Marumo and Asaoka, 1974;
441 Zhang et al., 2012). Zhang et al. (2012) suggested that the low nitrogen fixation in the
442 continental shelf was attributable to mixing processes and the influence of the
443 Changjiang River. Turbulence near the sea floor influences the surface water in the
444 shallower bottom region (Matsuno et al., 2006), and Zhang et al. (2012) suggested
445 that the physical disturbance reduces diazotrophy since diazotrophs including
446 *Trichodesmium* favor calm seas. Furthermore, the water in the continental shelf of the
447 ECS is strongly influenced by the Changjiang River. The N/P ratio of the Changjiang
448 River plume is significantly higher than the Redfield ratio, which results in
449 phosphorus limitation, and can contribute to the low nitrogen fixation (Zhang et al.,

450 2012). In the present study, despite the fact that the surface phosphate concentration
451 was low throughout the study areas, the N/P ratio was generally lower than the
452 Redfield ratio, suggesting that biological production was limited by the availability of
453 nitrogen compared to phosphate (Moore et al., 2008, 2013). Furthermore, the
454 insignificant difference in MLD among the ECS, the Kuroshio, and the Philippine Sea
455 ($p>0.05$; Tukey HSD test) indicated similar vertical mixing conditions. Therefore, the
456 environmental variables related to nitrogen fixation only slightly differed as
457 demonstrated by the nMDS plot..

458 In our study, we found a *Trichodesmium* spp. bloom near the Miyako Islands.
459 Recent studies demonstrated that *Trichodesmium* spp. thrived near oceanic islands
460 (Shiozaki et al., 2010, 2014c; Dupouy et al., 2011). Given that some aspect of the
461 environment around the islands increases *Trichodesmium* spp. abundance and that
462 they are transported from the islands to the Kuroshio, this can explain why the
463 *Trichodesmium* distribution was not estimated from environmental variables.
464 Accordingly, the low abundance of *Trichodesmium* spp. in the Philippine Sea was
465 likely due to the low density of islands. Furthermore, higher nitrogen fixation in the
466 Kuroshio than in the Philippine Sea might be explained in the same manner.
467 *Trichodesmium* is a major nitrogen fixer in the Kuroshio (Chen et al., 2008, 2014;

468 Shiozaki et al., 2014a), and our results showed that the bulk water nitrogen fixation
469 was positively correlated with *Trichodesmium* abundance.

470 The numerical simulation demonstrated that released particles from the Miyako
471 Islands were generally transported to the northeast and flowed along the Kuroshio
472 during summer between 2003 and 2009. Thus, if *Trichodesmium* increases and active
473 nitrogen fixation usually occurs around the Miyako Islands, the water would be
474 delivered to the Kuroshio. Furthermore, we performed additional particle tracking
475 experiments whose particle release points were set at major islands in the Ryukyu
476 Islands (Amami Islands, Okinawa Main Island, and the Ishigaki Islands) (Figs. S6 and
477 S7). The results demonstrated that the particles released from the other islands of the
478 Miyako Islands were also delivered to the Kuroshio, with some exceptions. Based on
479 the calculations for 2003–2009, 13–56% ($30 \pm 16\%$) of particles released from the
480 islands reached Area K by day 120 (Fig. S7).

481 Studies on nitrogen fixation around islands in the study region are fairly limited
482 (Liu et al., 2013), and the present study is the first report of a *Trichodesmium* bloom
483 around islands in the area. The Miyako Islands are surrounded by reefs, and studies
484 have shown that *Trichodesmium* blooms can be associated with reef environments
485 (Bell et al., 1999; McKinna et al., 2011). However, the factors causing the

486 *Trichodesmium* blooms around islands are not well understood (Shiozaki et al.,
487 2014c). Further studies are required to identify which characteristics of the near island
488 environment are important for the growth and/or accumulation of *Trichodesmium* and
489 other diazotrophs.

490

491 5. CONCLUSIONS

492 We hypothesize that the high abundance of *Trichodesmium* spp. and active
493 nitrogen fixation in the Kuroshio were ascribable not to the unique nutrient
494 environment, but rather to the supply of *Trichodesmium* spp. and other diazotrophs
495 from the surrounding islands. The Ryukyu Islands would not be the only islands with
496 abundant *Trichodesmium* spp., as *Trichodesmium* spp. also flourish in the upstream
497 Kuroshio near Luzon Island (Chen et al., 2008). Therefore, the abundance of
498 *Trichodesmium* spp. would be generally increased around islands situated along the
499 Kuroshio, and the abundant *Trichodesmium* spp. would likely be transported to the
500 mainstream of the Kuroshio. *Trichodesmium* is a major diazotroph in the Kuroshio
501 (Chen et al., 2008, 2014; Shiozaki et al., 2014a), and diazotrophy in the Kuroshio is
502 considered to influence the nutrient stoichiometry in the North Pacific (Shiozaki et al.,
503 2010). Thus, our results indicate that phenomena around the islands located along the

504 Kuroshio are important for determining the partial nitrogen inventory in the North

505 Pacific.

506

507 **Author Contributions**

508 T.S., S.T., S.I., and K.F. designed the experiment and T.S., S.T., T.K., X.L., F.H., and
509 K.F. collected the samples at sea. T.S. determined nitrogen fixation and abundance of
510 *Trichodesmium* spp. during the KT-06-21, KT-07-21, KT-09-17, and *Nagasaki-maru*
511 242 cruises, and X.L. did during the KT-10-19 cruise. T.S. analyzed datasets of
512 satellite and climatological current field. S.T. analyzed concentration of dissolved iron.
513 S.I. performed numerical experiments. T.K. and F.H. determined nutrient
514 concentration. T.S. prepared the manuscript with contributions from all co-authors.

515

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528

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- 677

678 Table 1 Summary of *Trichodesmium* at the surface, and depth-integrated nitrogen

679 fixation and its related parameters in the four representative study areas.

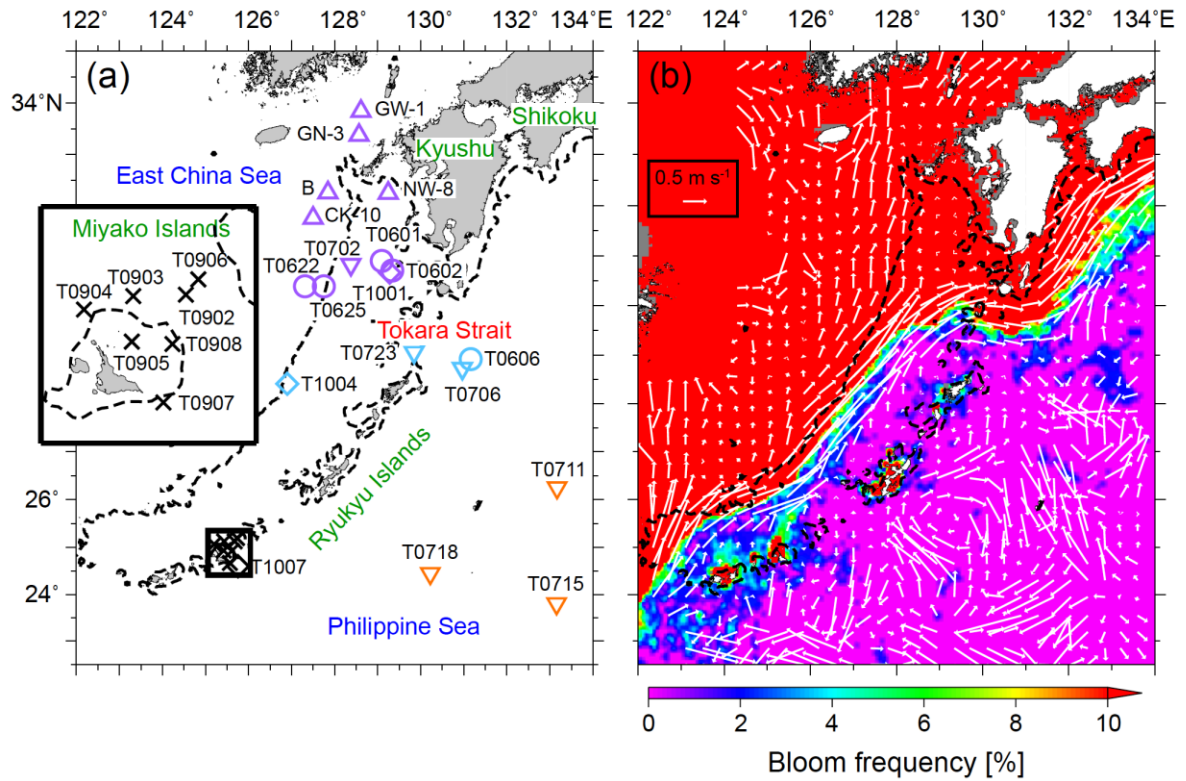
Area	<i>Trichodesmium</i> * [filaments l ⁻¹]	N ₂ fixation [μmolN L ⁻¹ d ⁻¹]	Temperature* [°C]	MLD [m]	NO ₃ ⁻ +NO ₂ ^{-*} † [nM]	PO ₄ ^{3-*} † [nM]	DFe* [nM]
East China Sea	21±58	170±140	28.5±1.2	24±12	19±11	15±9	0.76±0.18
Kuroshio	43±33	199±142	29.4±0.81	27±8	9±8	15±7	0.45±0.13
Philippine Sea	8±8	58.3±25.1	29.4±0.1	23±3	8±3	14±19	0.51±0.25
Miyako Islands	3019±8478	201±274	29.3±0.3	40±12	61±128	8±7	0.38±0.24

680 * values in surface water

681 †When the concentration was below the detection limit (3 nM), we assumed a concentration of 3 nM to

682 calculate the mean.

683



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685

686 Figure 1. (a) Sampling stations during the KT-06-21 (circles), KT-07-22 (inverted

687 triangles), KT-09-17 (crosses), KT-10-19 (diamonds), and 242 (triangles) cruises.

688 Symbols of stations located in the East China Sea, the Kuroshio, the Philippine Sea,

689 and near the Miyako Islands are indicated in purple, light blue, orange, and black,

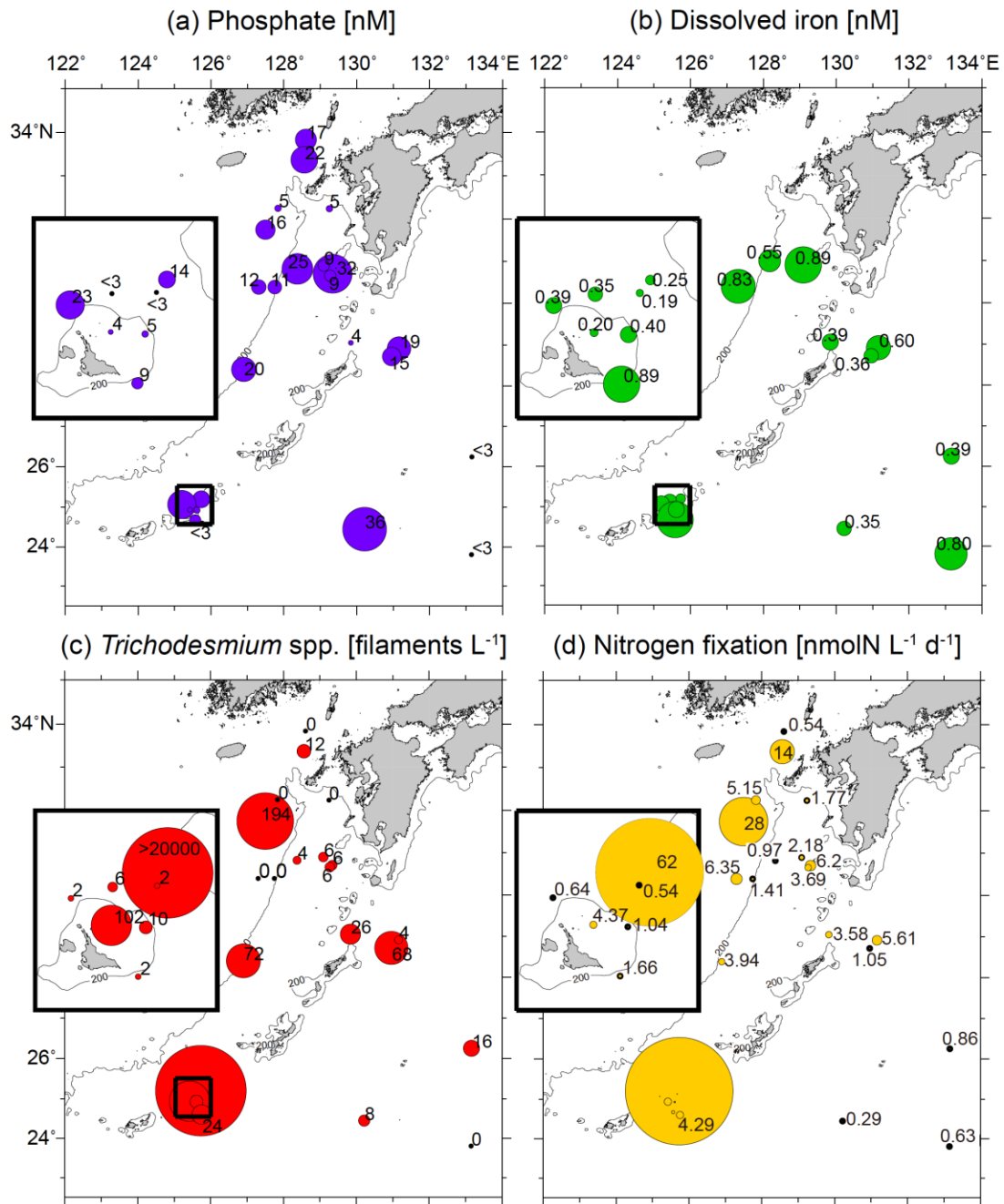
690 respectively. (b) Climatological surface current fields during summer (1953–2008)

691 from geoelectrokinetograph measurements and ship-mounted ADCP data. The

692 background contour represents the percentage of chlorophyll *a* of $>0.15 \text{ mg m}^{-3}$

693 during summer between 2003 and 2009. Dashed lines indicate 200 m isobaths.

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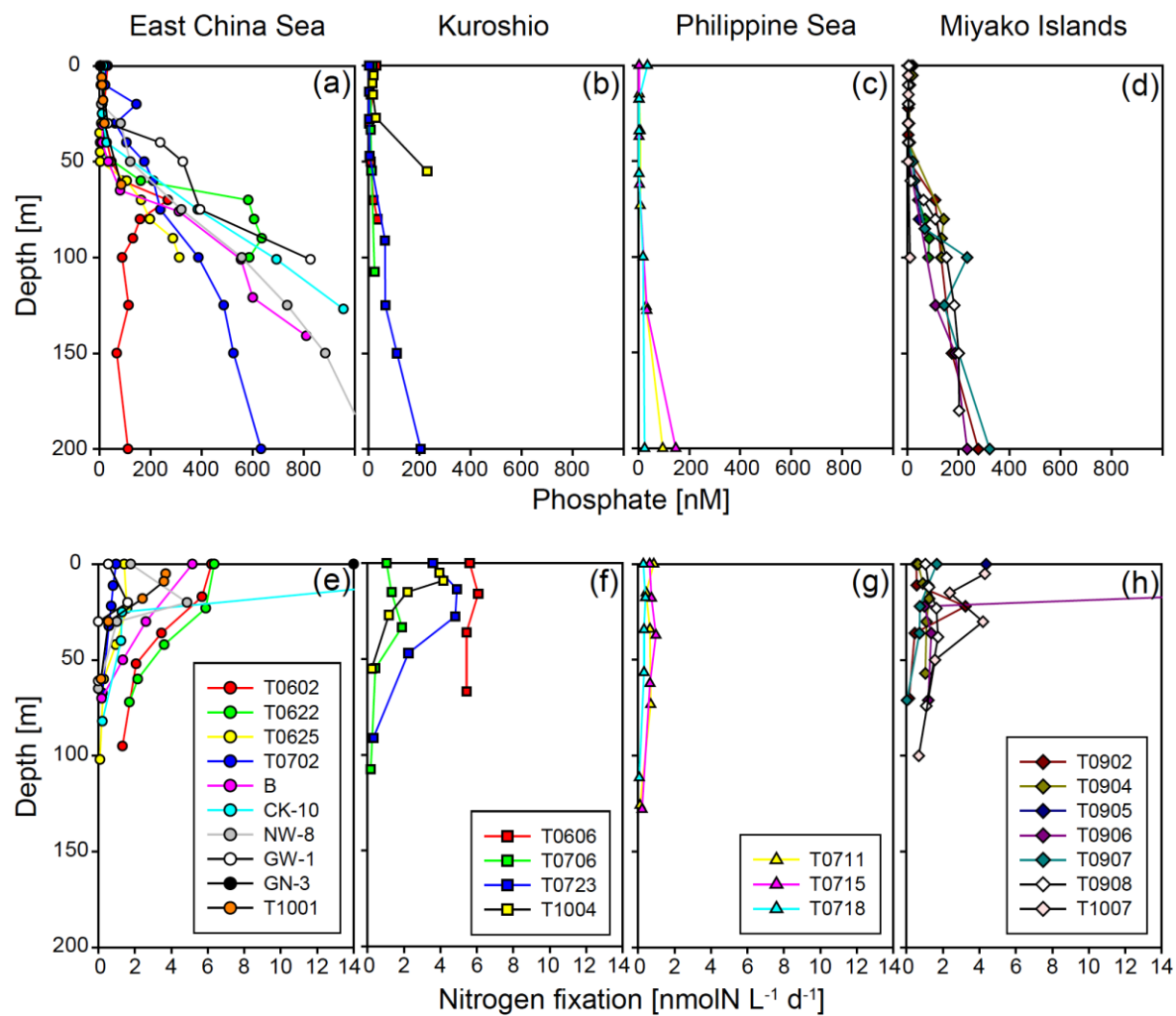
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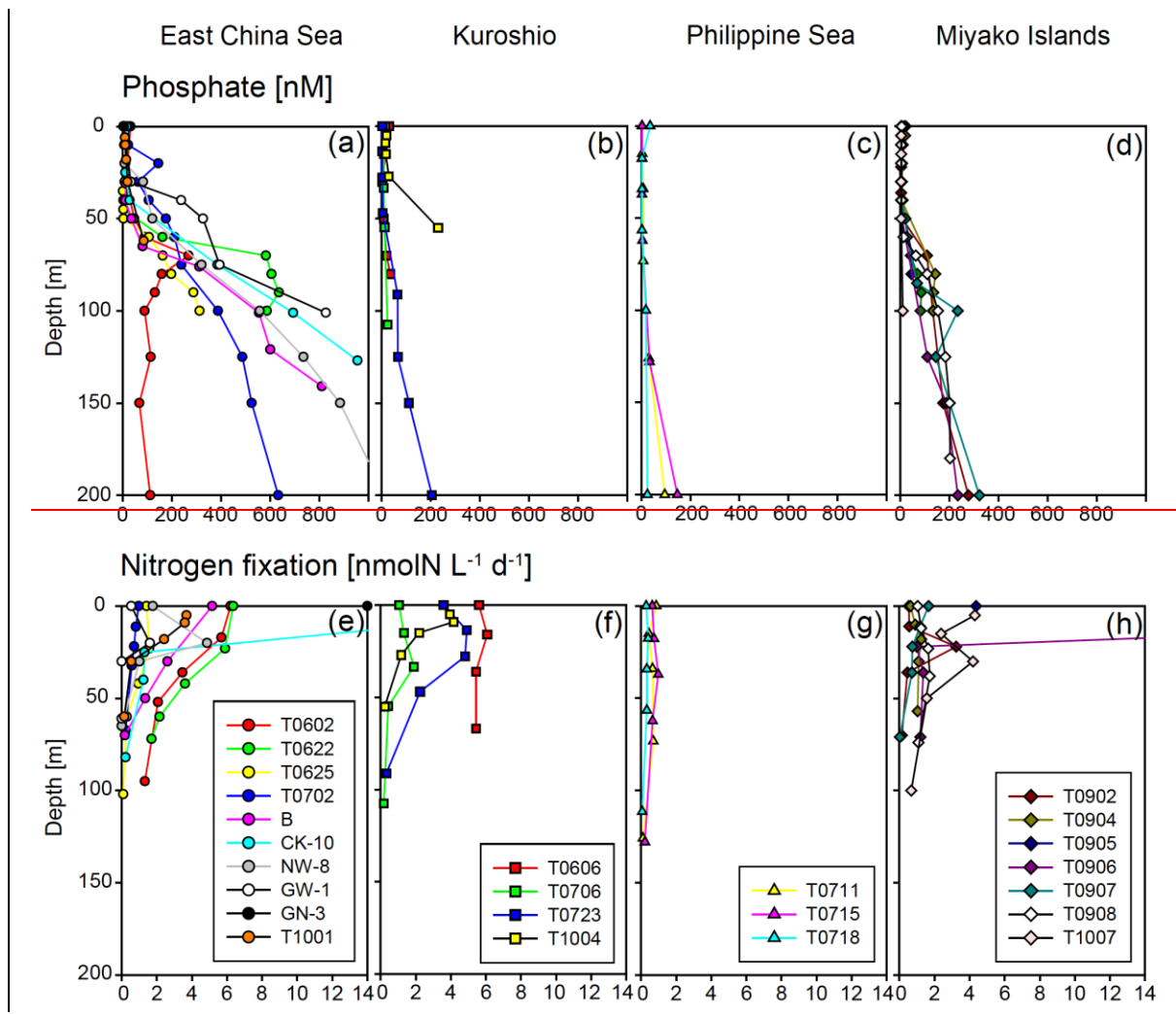
697 Figure 2. Distribution of (a) phosphate, (b) dissolved iron, (c) *Trichodesmium* spp.,

698 and (d) nitrogen fixation at the surface. The parameters in the small boxes indicate

699 results from the KT-09-17 cruise. The areas of the circles are proportional to the

700 concentration, abundance, or activity.





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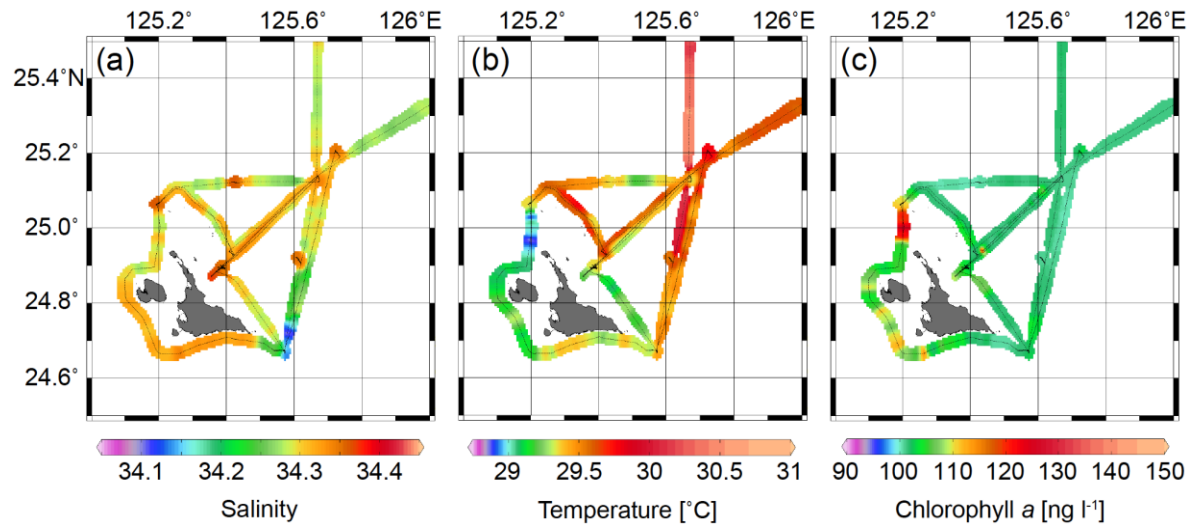
703

704 Figure 3. Vertical profiles of phosphate and nitrogen fixation in the East China Sea (a

705 and e), the Kuroshio (b and f), the Philippine Sea (c and g), and the Miyako Islands (d

706 and h).

707



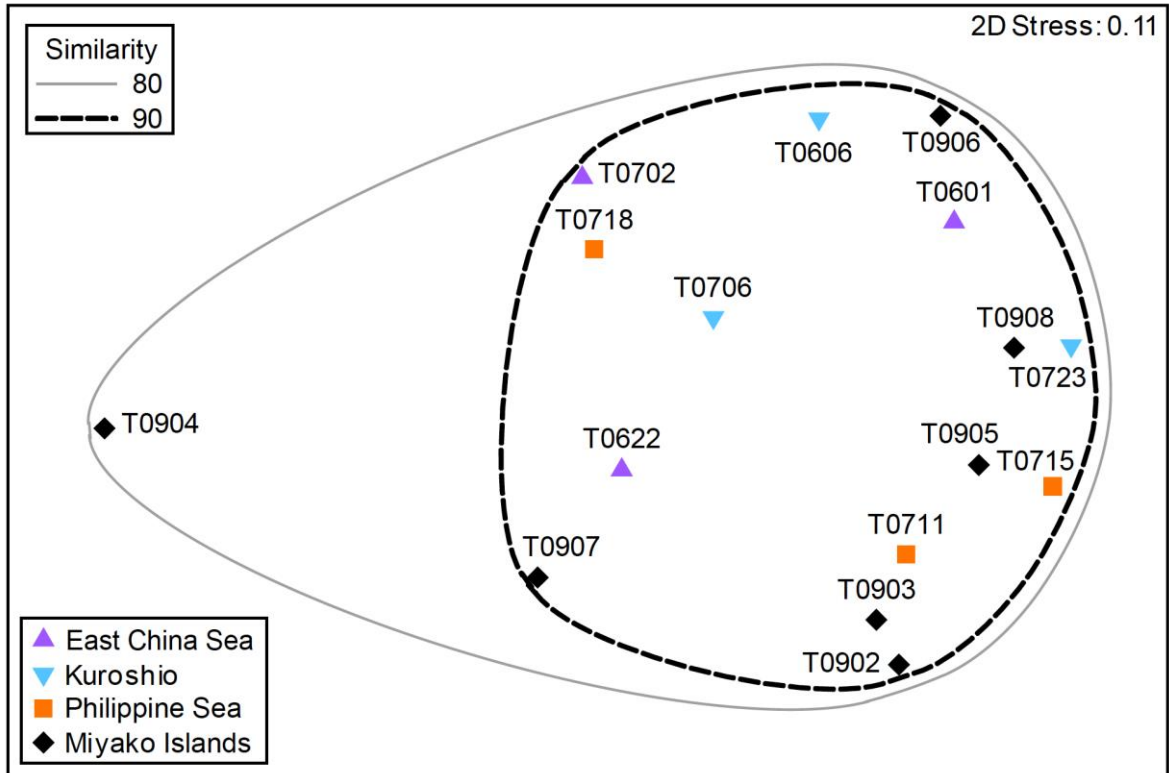
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710 Figure 4. Surface (a) salinity, (b) temperature, and (c) chlorophyll *a* during the

711 KT-09-17 cruise.

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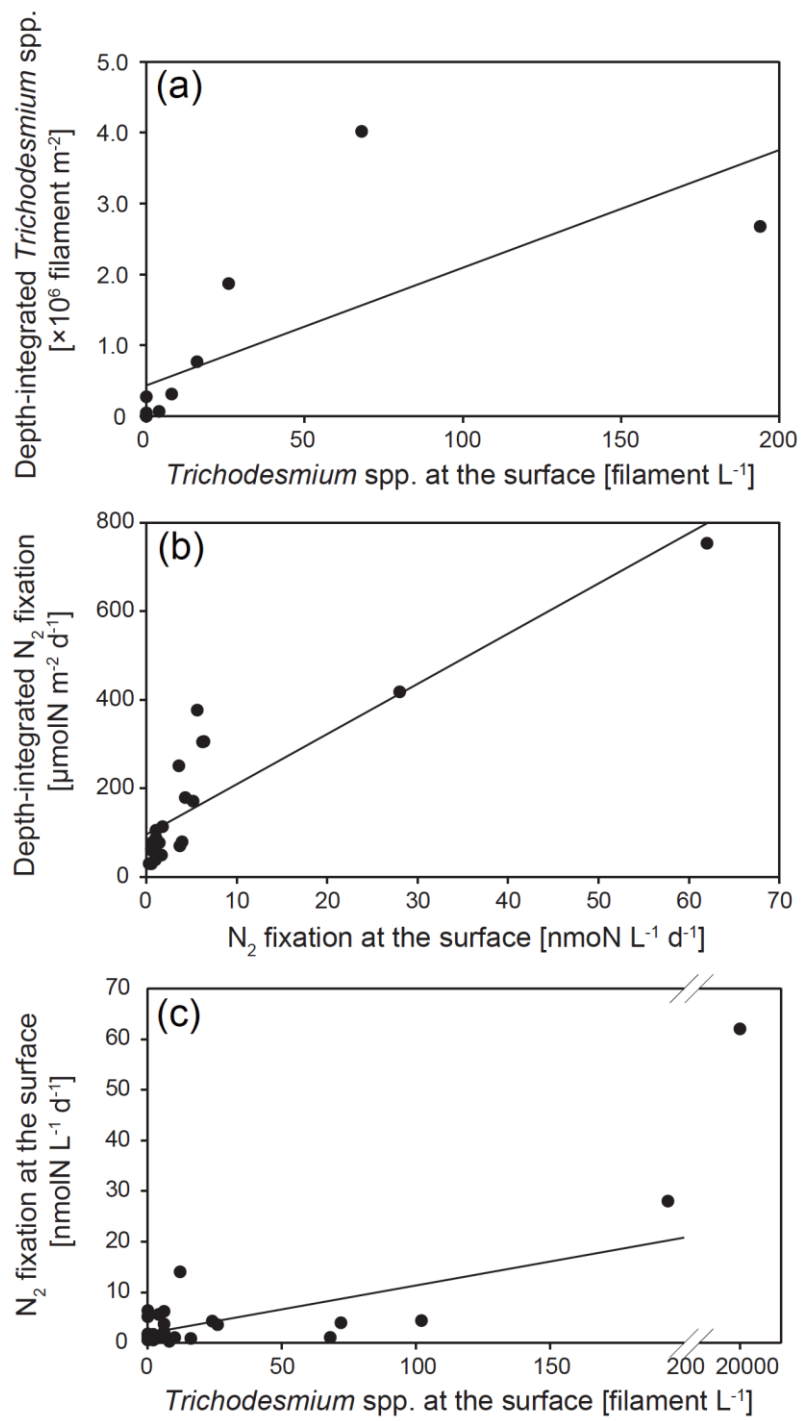


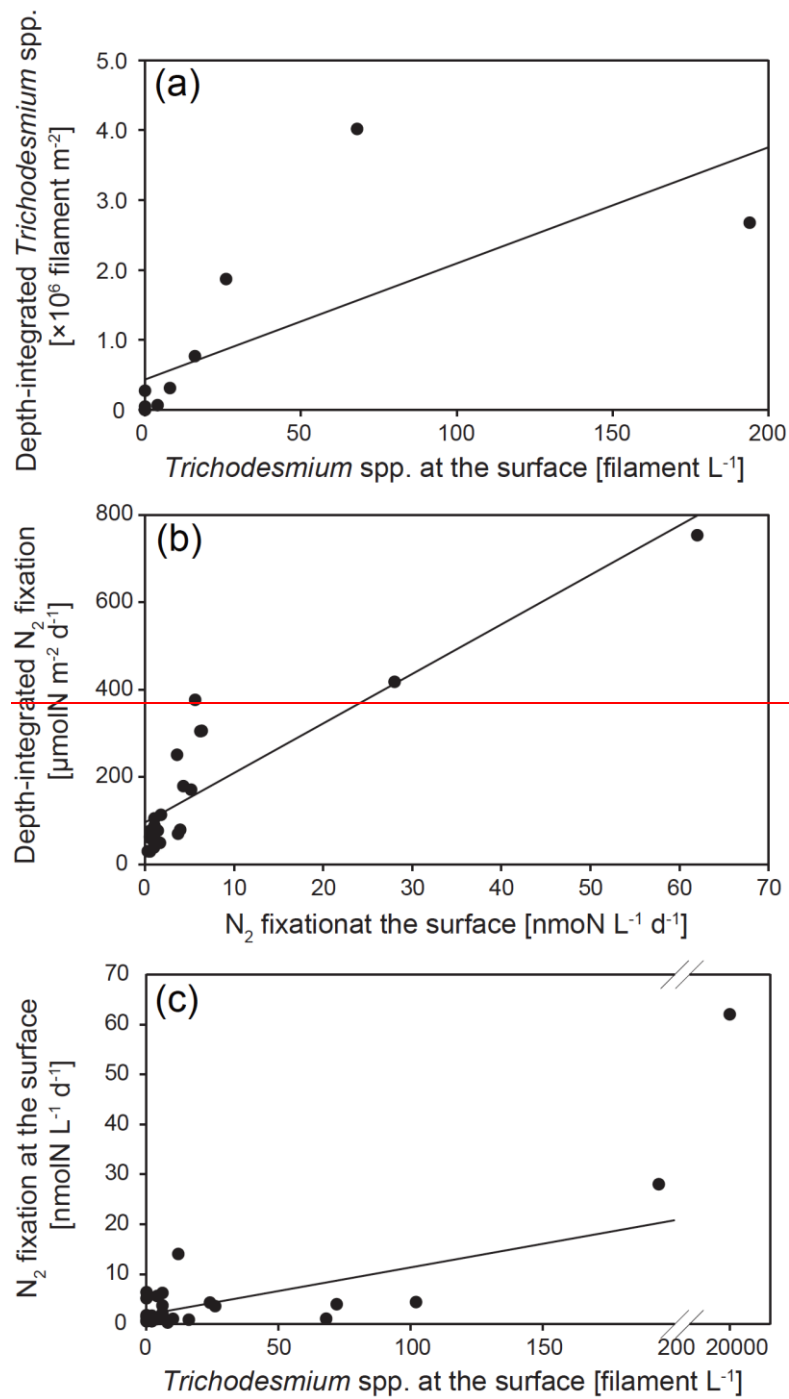
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715 Figure 5. nMDS ordination of sampling stations with environmental variables

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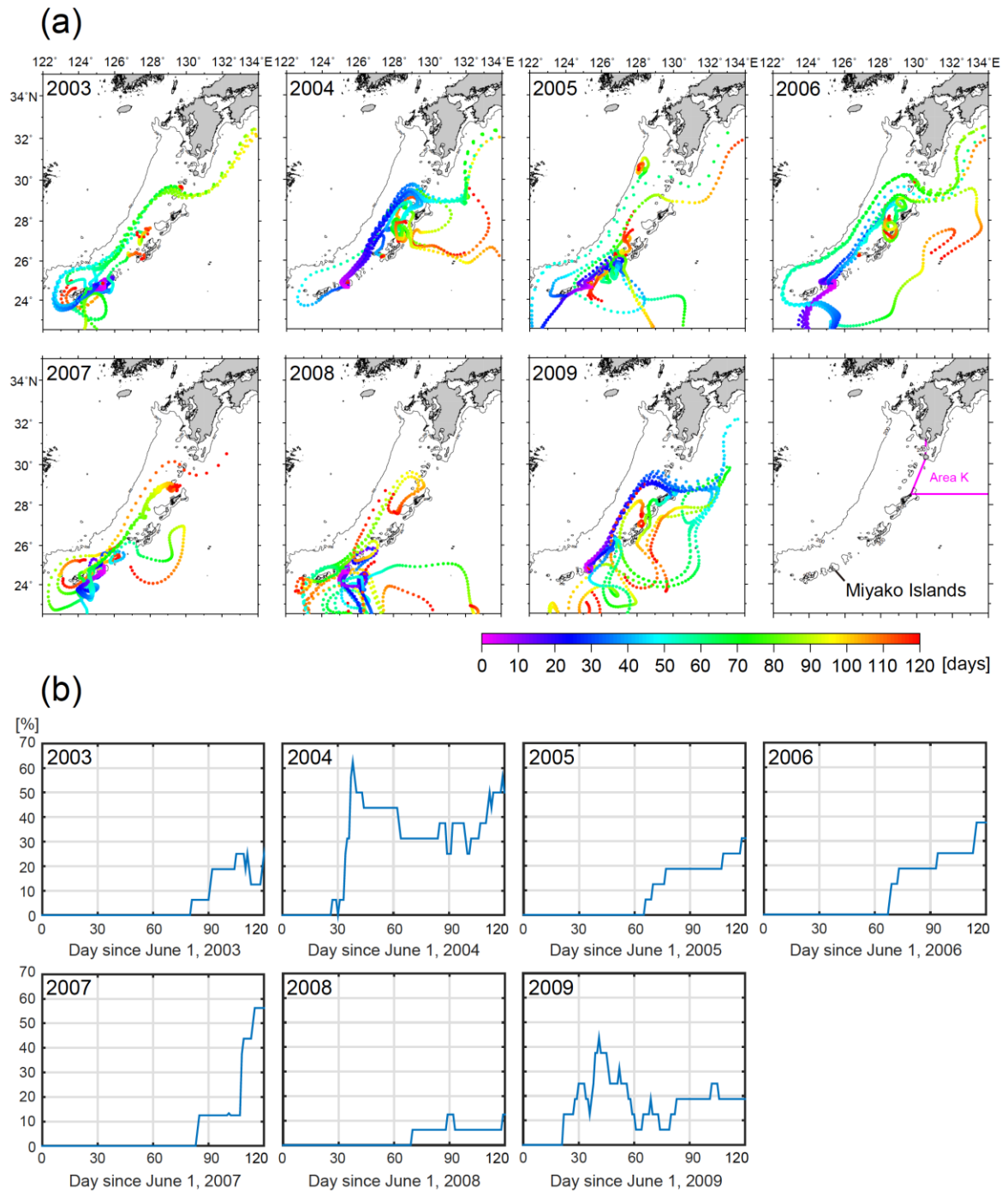
719

720 Figure 6. Relationships (a) between surface and depth-integrated *Trichodesmium* spp.

721 abundance, (b) between surface and depth-integrated nitrogen fixation rates, and (c)

722 between *Trichodesmium* spp. abundance and nitrogen fixation rate at the surface.

723



724

725

726 Figure 7. (a) Trajectories of particles released from points around the Miyako Islands

727 on June 1, 2003–2009. (b) The ratio of particles delivered to Area K to the total

728 released particles.