




## Article

# Effects of Different Nutrient and Trace Metal Concentrations on Growth of the Toxic Dinoflagellate *Gymnodinium catenatum* Isolated from Korean Coastal Waters

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Received: 29 May 2020; Accepted: 17 June 2020; Published: 18 June 2020



**Abstract:** The effects of the addition of nutrients (nitrate: N; phosphate: P; and vitamin B<sub>1</sub>) and trace metals (iron: Fe; Copper: Cu; and selenium: Se) on the growth of *Gymnodinium catenatum*, which was isolated from Korean coastal waters, were investigated. The Korean isolate of *G. catenatum* grew under a wide range of concentrations of N and P. Whilst high concentrations of N (> N: P ratio of 23.5) did not stimulate the growth rate, an enhanced growth rate and cell density were observed with the addition of P. The experimental addition of vitamin B<sub>1</sub> revealed that *G. catenatum* is not dependent on vitamin B<sub>1</sub> for growth. Moreover, the addition of Fe and Cu resulted in no significant differences in the growth patterns and rates of *G. catenatum* between the controls and treatments. It is thus possible that growth of the Korean isolate of *G. catenatum* does not require high concentrations of Fe and Cu. However, the cell densities were enhanced in the stationary phases of treatments upon addition of Se, and the maximum cell densities were higher than those in the culture experiments upon additions of other nutrient and trace metals. Our findings indicate that *G. catenatum* prefers P and Se for proliferation, rather than other nutritional sources.

**Keywords:** growth rate; nitrate; phosphate; vitamin B<sub>1</sub>; copper; iron; selenium

## 1. Introduction

Paralytic shellfish toxins, which are produced by some marine toxic dinoflagellates and freshwater cyanobacteria [1–3], can be accumulated in a wide variety of marine organisms through the food web, and this accumulation can lead to outbreaks of paralytic shellfish poisoning (PSP) [2,4–6]. PSP outbreaks have been reported globally and caused human intoxications and death, serious economic

losses in fisheries industries, and negative impacts on marine ecosystems (e.g., [2,4,7,8]). Due to such outbreaks, the toxicity, geological distribution, nutrient ecophysiology and classification of the causative organisms have been studied intensively (e.g., [2,3]).

To date, about 12 species of *Alexandrium*, *Pyrodinium bahamense* and *Gymnodinium catenatum* have been reported as the main toxic dinoflagellates responsible for PSP [3,9–16]. Of these, the PSP outbreaks caused by *G. catenatum* have been frequently reported in Australia [17], Portugal [18], Spain [19], Morocco [20], Japan [21] and Latin America [3]. According to Band-Schmidt et al. [3], blooms of *G. catenatum* have been associated with outbreaks of PSP. The blooms of *G. catenatum* and related PSP incidents have not been recorded to date in Korean coastal area; however, toxicity was reported from isolates of *G. catenatum* collected from the southern coast of Korea [22]. Nevertheless, little is known about the effects of environmental factors related to the growth of *G. catenatum* isolates in Korean coastal waters.

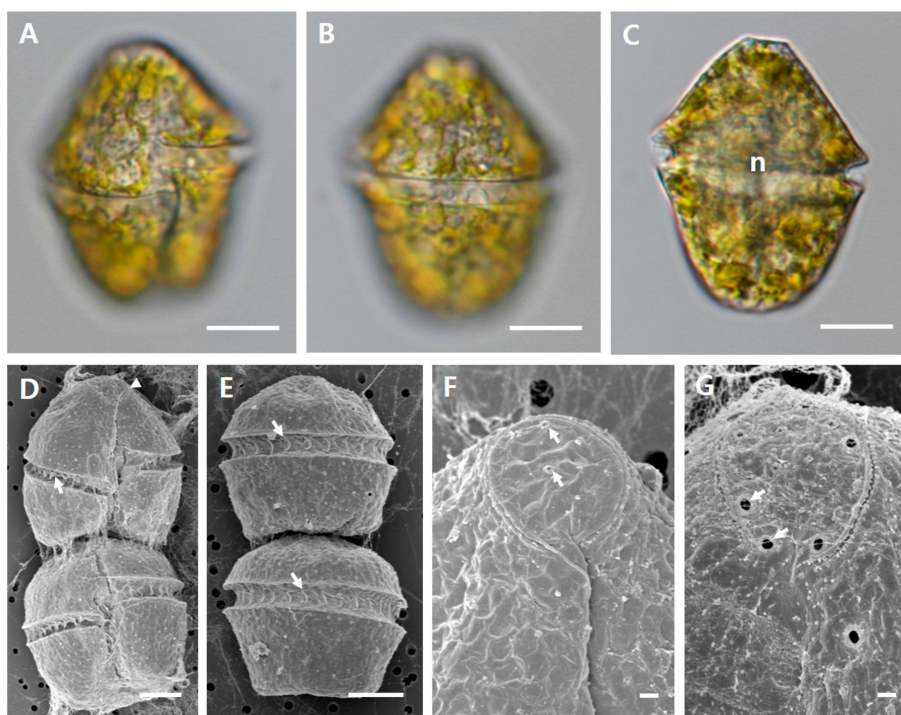
Nutrients, together with light and temperature, are important factors in phytoplankton growth and distribution in marine ecosystems [23,24]. In particular, there is evidence of strong relationships between anthropogenic nutrient loading and harmful algal blooms [2,4,25,26]. An understanding of environmental factors, such as nutrients, that can lead to the proliferation of harmful dinoflagellates is important in predicting harmful algal blooms, as well as for developing management strategies to deal with this threat. Despite extensive study of the effects of nutrients on the growth of some harmful dinoflagellates through laboratory and field experiments, our knowledge of the effects of nutrients on harmful dinoflagellates is quite poor (e.g., [27]). In addition, as most of the studies of harmful algal blooms focusing on nutrients have primarily investigated the importance of macronutrients, such as nitrate and phosphate [28,29], the effects of micronutrients, such as vitamins and trace metals, in relation to the growth of harmful species has rarely been considered.

Early studies revealed that trace metals such as iron [30,31] and selenium [32] are important factors in the development of some harmful algal blooms, and selenium, in particular, has been considered to stimulate the growth of *G. catenatum* [33,34]. Couet et al. [35] also reported the over-production of toxins of harmful dinoflagellates exposed to copper stress, and Tang et al. [36] discussed the significant ecological role of B-vitamins in regulating the dynamics of harmful algal blooms. However, in Korean coastal waters, the effects of nutritional factors including macronutrients on harmful species have rarely been investigated, and most studies have focused on *Alexandrium* species. According to Lee et al. [37], *G. catenatum* is distributed widely along the south coast of Korea, indicating that this species can be regarded as potentially harmful species in Korean coastal areas [22]. In this study, to improve our understanding of the ecophysiology of *G. catenatum* in Korean coastal waters, we aim to investigate the growth of the Korean isolate of *G. catenatum* under different concentrations of nutrients and trace metals.

## 2. Materials and Methods

### 2.1. Culture of *Gymnodinium catenatum*

A strain of *Gymnodinium catenatum* (LIMS-PS-2604) was obtained from the Library of Marine Samples, Korea Institute of Ocean Science and Technology. This strain was established from phytoplankton samples collected in the South Sea of Korea (34°29′7.68″ N, 128°28′54.54″ E) and has been maintained at 20 °C and ca 100  $\mu\text{mol photons m}^{-2} \text{s}^{-1}$  cool-white illumination (fluorescent lamp) under a 12L:12D photo-cycle. The morphological features of *G. catenatum* were photographed using an AxiCam MRc digital camera on an upright microscope (Axio Imager 2, Zeiss, Germany) and a field emission scanning electron microscope (JSM 7600F, JEOL, Tokyo, Japan), which are shown in Figure 1. The phylogeny of the strain was reported in a previous study [38].



**Figure 1.** Light and scanning electron micrographs of a Korean isolate of *Gymnodinium catenatum* (culture strain LIMS-PS-2604). (A) Surface focus of ventral view showing the cingulum. (B) Surface focus of dorsal view showing the cingulum. (C) Deeper focus of dorsal view showing the nucleus (n). (D) ventral view of a two-celled chain showing apical groove (arrowhead) and transverse flagella (arrow). (E) dorsal view of a two-celled chain showing cingulum and transverse flagella (arrows). (F,G) Details of apical groove, with small pores (arrows). Scale bars: A–E = 10  $\mu\text{m}$ ; F–G = 1  $\mu\text{m}$ .

Experimental cultures were established in 2 L culture bottles (SPL, Pocheon, Korea) containing f/2 culture medium (Marine Water Enrichment Solution, Sigma Aldrich, St. Louis, MO, USA) without silicate, prepared with sterile sea water at 20 °C and ca 100  $\mu\text{mol photons m}^{-2} \text{s}^{-1}$  cool-white illumination (fluorescent lamp) under a 24L:0D photo-cycle on a roller apparatus (Wheaton, IL, USA). Cultures from the exponential growth phase were used for the experiments.

## 2.2. Growth Experiment

Surface seawater collected from the East China Sea (32°00' N, 127°00' E) was used to prepare the culture medium with controlled concentrations of nitrate (N), phosphate (P), vitamin B<sub>1</sub>, Iron (Fe) and Copper (Cu). The concentrations of dissolved inorganic nitrogen (sum of  $\text{NH}_4^+$ ,  $\text{NO}_2^-$ ,  $\text{NO}_3^-$ ) and phosphorus ( $\text{PO}_4^{3-}$ ) in the seawater were 7.4 and 0.6  $\mu\text{M}$ , respectively, as measured by a nutrient auto analyzer (QuickChem 8000, Lachat, Loveland, CO, USA). For the selenium (Se) addition, seawater was collected from a location off the Korean coast (37°32'26.62" N, 130°50'43.31" E). The seawaters were filtered through a 47 mm membrane filter and autoclaved for all experiments.

In this study, we carried out two major experiments involving nutrients (N, P and vitamin B<sub>1</sub>) and trace metals (Fe, Cu and Se). The basal culture medium used to control the concentrations of nutritional factors was the f/2 culture medium without silicate (f/2-Si culture medium), and the N:P ratio in the stock solution with this culture medium was 24.2 (the values of nitrate and phosphate calculated from the collected seawater and the culture medium). Treatments were made by adding or reducing the concentrations of the nutrients and trace metals, and the treatment without addition of nutritional factors was used as a control (Table 1). However, as the original recipe of the f/2 culture medium does not include Se, the Se concentration was varied through the addition of selenous acid to a solution of the basal culture medium (Table 1).

**Table 1.** Nutrient and trace metal concentrations ( $\mu\text{M}$ ) used in the experimental design. N:P ratio in parentheses.

	Control	Treatment 1	Treatment 2	Treatment 3	Treatment 4	Treatment 5
<b>Nutrient</b>						
N	None added	$8.82 \times 10^1$ (2.6)	$8.82 \times 10^2$ (24.2)	$1.76 \times 10^3$ (48.1)	-	-
P	None added	$3.62 \times 1$ (210.9)	$3.62 \times 10^1$ (24.2)	$7.24 \times 10^1$ (12.2)	-	-
Vitamin B <sub>1</sub>	None added	$2.96 \times 10^{-2}$	$2.96 \times 10^{-1}$	$5.92 \times 10^{-1}$	-	-
<b>Trace Metal</b>						
Fe	None added	$1.17 \times 1$	$1.17 \times 10^1$	$2.34 \times 10^1$	-	-
Cu	None added	$3.93 \times 10^{-3}$	$3.93 \times 10^{-2}$	$7.86 \times 10^{-2}$	-	-
Se	None added	$10^{-5}$	$10^{-4}$	$10^{-3}$	$10^{-2}$	$10^{-1}$

Growths of *Gymnodinium catenatum* were tested in stock solutions supplied with different concentrations of nutrients and trace metals and filtered and autoclaved seawater was used to make the stock solutions. Twenty-six stock solutions with target concentrations of nutrients and trace metals were made (Table 1), and 30 mL of each solution was added to separate 50 mL Pyrex tubes. The subcultures of *G. catenatum* for the experiments were established in 2 L culture bottles (SPL, Pocheon, Korea) containing individual nutritional source-poor media, and these were used for inoculation into experimental tubes. *G. catenatum* in concentrations ranging from 100 to 200 cells/mL was inoculated into the 50 mL Pyrex tubes. The tubes were placed in incubators and incubated at 20 °C and ca 100  $\mu\text{mol photons m}^{-2} \text{s}^{-1}$  cool-white illumination (fluorescent lamp) under a 14L:10D photo-cycle. All experiments were conducted in triplicate.

### 2.3. Calculation of Growth Rate

Culture growth was monitored at 2 day intervals for 32 days and determined using an in vivo fluorometer (Turner Designs Model 10-AU, Sunnyvale, USA), and the fluorescence data were used to calculate specific growth rates. The regression equation for in vivo fluorescence values provided a good fit to the observed cell densities; the adjusted  $r^2$  value for *Gymnodinium catenatum* was >0.99 (data not shown). To estimate the specific growth rate ( $\mu$ ) of *G. catenatum*, we used the following equation:

$$\mu = \log_2 (N_t - N_0) / t_1 - t_0 \quad (1)$$

where  $N_0$  and  $N_t$  are the initial ( $t_0$ ) and final ( $t_1$ ) in vivo fluorescence values during the incubation experiments, respectively. The in vivo fluorescence values estimated during the logarithmic growth phase were used to obtain the specific growth rate.

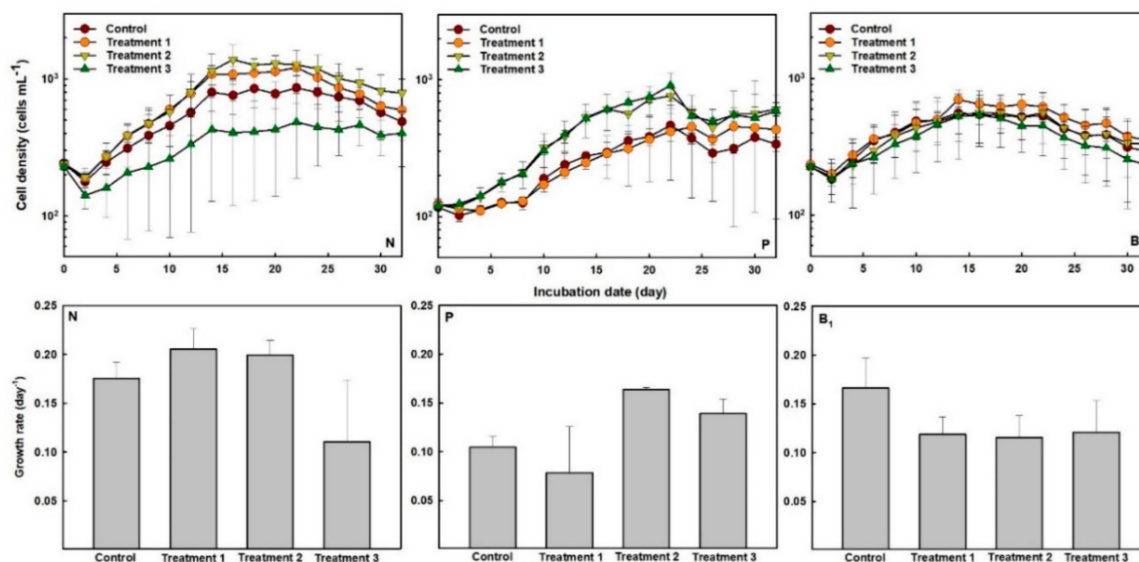
## 3. Results and Discussion

### 3.1. Growth of *Gymnodinium catenatum* under Different Concentrations of N, P and Vitamin B<sub>1</sub>

The growth curves and rates of *Gymnodinium catenatum* cultures exposed to different concentrations of nutrients (N, P and vitamin B<sub>1</sub>) are shown in Figure 2 and Table 2. In all the nutrient treatments, *G. catenatum* culture grew consistently during days 2–20 (Figure 2). In the N control, and the treatments 1 and 2, *G. catenatum* entered the senescence phase from day 24, following an stationary phase for 8 days; however, in treatment 3, with the addition of  $1.76 \times 10^3 \mu\text{M}$  nitrate, the stationary phase continued after an incubation of 14 days (the exponential growth phase). The pattern in treatment 3 was observed with low cell density, and the growth rate ( $0.11 \text{ day}^{-1}$ ) was also the lowest. The maximum cell density ( $1389 \text{ cells mL}^{-1}$ ) was observed in treatment 2 (Table 2), and the control and treatments 1 and 2 had similar growth rates ( $0.18$ – $0.21 \text{ day}^{-1}$ ). Compared with the N control and the treatments, the high growth rates of *G. catenatum* in the P addition experiments were observed in treatments 2 (addition of  $3.62 \times 10^1 \mu\text{M}$  phosphate) and 3 (addition of  $7.24 \times 10^1 \mu\text{M}$  phosphate), with high cell densities ( $764$  and  $909 \text{ cells mL}^{-1}$ ) (Table 2); however, the maximum cell density was lower than that for the N treatments. Low growth rates were observed in the control ( $0.10 \text{ day}^{-1}$ ) and



treatment 1 ( $0.08 \text{ day}^{-1}$ ), with low cell densities (Table 2), and the growth rate in treatment 3 with P addition was higher than that in N treatment 3. *G. catenatum* grew consistently until day 22 and then entered a stationary phase, and the senescence phase was not observed until 32 days of incubation. This growth pattern was similar to that observed in N treatment 3.



**Figure 2.** Growth curves and rates of *Gymnodinium catenatum* culture exposed to different nitrate (N), phosphate (P) and vitamin B<sub>1</sub> concentrations.

**Table 2.** Growth rates ( $\text{day}^{-1}$ ) and maximum cell densities ( $\text{cells mL}^{-1}$ ) of *Gymnodinium catenatum* in the controls and treatments with nutrients and trace metals. Maximum cell density in parentheses.

	Control	Treatment 1	Treatment 2	Treatment 3	Treatment 4	Treatment 5
<b>Nutrient</b>						
N	0.18 (862)	0.21 (1214)	0.20 (1389)	0.11 (485)	-	-
P	0.10 (464)	0.08 (455)	0.16 (764)	0.14 (909)	-	-
Vitamin B <sub>1</sub>	0.17 (545)	0.12 (699)	0.12 (561)	0.12 (540)	-	-
<b>Trace Metal</b>						
Fe	0.14 (851)	0.15 (1125)	0.16 (1310)	0.14 (852)	-	-
Cu	0.17 (1055)	0.14 (986)	0.15 (1071)	0.17 (1058)	-	-
Se	0.17 (993)	0.16 (1391)	0.16 (1388)	0.15 (1811)	0.16 (2021)	0.15 (1862)

Band-Schmidt et al. [39] investigated the growth rates of *G. catenatum* strains under different growth conditions, and reported the growth rates of a strain (established from Bahía Concepción Gulf of California, Mexico) tested in f/2 medium at  $20^\circ\text{C}$  and ca  $100 \mu\text{mol photons m}^{-2} \text{ s}^{-1}$  cool-white illumination (fluorescent lamp) under a 12L:10D photo-cycle, which is similar to the conditions described here, ranging from  $0.28$  to  $0.32 \text{ day}^{-1}$ . These growth rates are higher than those obtained for f/2-Si culture medium (referred to as treatment 2) in the present study. However, Bustillos-Guzmán et al. [40] reported that the growth rate of *G. catenatum* culture established from Concepción Bay, Gulf of California is  $0.24 \text{ day}^{-1}$  (the N:P ratio of 23.5), which is similar to own observations. This finding indicates that the growth rate of *G. catenatum* can vary within strains, or can depend on different geographical strains. Bustillos-Guzmán et al. [40] also found that the maximum cell density for treatment with a N:P ratio of 23.5 is higher than those for treatments with low N:P ratios (5.4 and 9.2), and that higher N:P ratios than 23.5 can produce intermediate densities. This result accords with our observation that treatment 2 with a N:P ratio of 24.2 produce a maximum cell density. However, in this case, treatment 3, with the addition of a relatively high N concentration (a N:P ratio of 48.1), had the lowest maximum cell density

(485 cells mL<sup>-1</sup>) among the control and all treatments, indicating that higher N:P ratios than 24.2 do not increase the growth rate and cell density of Korean isolates of *G. catenatum*.

Interestingly, although the cell density in treatment 3 was lower than those in the control and other treatments, no senescence phase was observed in treatment 3. The senescence phase is observed when nutritional sources such as nitrate and phosphate in the culture medium are lower. Consequently, while the addition of a high N concentration (or >N:P ratio of 24.2) did not stimulate growth rate or cell density in the Korean isolate of *G. catenatum*, it can nevertheless have a positive effect on the maintenance of cell density. According to Bustillos-Guzmán et al. [40], *G. catenatum* may have a high storage capacity for phosphorus and nitrogen. This indicates that under N sufficient condition, *G. catenatum* utilizes a N storage strategy for maintaining cell density.

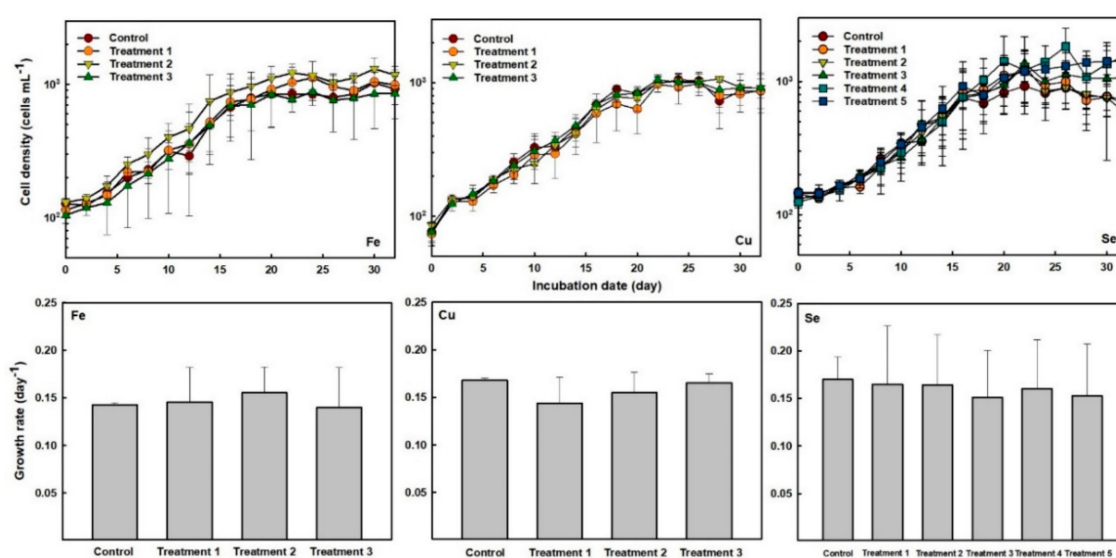
In contrast to the additions of N, additions of P stimulated the growth rate and cell density of *G. catenatum* (Figure 2). According to Oh et al. [41], under P-limited conditions the growth of *G. catenatum* can be stimulated by the addition of dissolved organic phosphorus (DOP). DOP can act as an additional source of P for microorganisms in a marine ecosystem in which dissolved inorganic phosphorus is seasonally depleted [42], and *G. catenatum* can take advantage for the uptake of P from DOP [41]. In addition, Yamamoto et al. [43] concluded that in the P-depleted condition, *G. catenatum* may be able to form blooms in Hiroshima Bay, Japan, with a higher affinity for DOP [43]. These observations indicate that sources of P are an important nutritional factor in enhancing the growth of *G. catenatum*. However, as in this study the additions of both N and P resulted in similar maximum cell densities (Table 2), sources of P alone certainly favor neither the enhancement of cell density of *G. catenatum* nor bloom formation. In the growth curve for the P control and treatments, no senescence phase was observed for the 32-day incubation, which differs from the growth pattern seen for the N control, and for treatments 1 and 2 with moderate concentrations of N. This may be related to the storage strategy for P, as well as for a high N concentration, suggesting that *G. catenatum* may be able to make use of one of two nutritional sources for maintaining cell density, when either N or P is depleted or sufficient.

Tang et al. [36] reported that vitamins are specifically important to the occurrence of harmful algal blooms, and Gobler et al. [44] observed the enhancement of large dinoflagellates by the enrichment of coastal waters with vitamin B<sub>1</sub> and B<sub>12</sub>. However, in this study the addition of vitamin B<sub>1</sub> did not stimulate the growth rate of *G. catenatum*, and the highest growth rate of *G. catenatum* was observed in the control without addition of vitamin B<sub>1</sub> (Figure 2), and the maximum cell density was observed for the treatment 1 (addition of  $2.96 \times 10^{-2}$  μM vitamin B<sub>1</sub>) (Figure 2). Although we did not measure the concentration of vitamin B<sub>1</sub> in the seawater used to make the solutions, neither the increased growth rate nor maximum cell density seem to be related to high concentration of vitamin B<sub>1</sub>. In addition, despite the fact that the culture medium for the controls and treatments in this experiment contained N, P, and vitamin B<sub>12</sub>, significant increases in growth and cell density of *G. catenatum* were not observed, reflecting the fact that *G. catenatum* does not require these vitamins for growth. According to Croft et al. [45], more than 20% of the 306 microalgal species surveyed are auxotrophs for the vitamin B<sub>1</sub>, and Tang et al. [36] reported that, among 45 species of dinoflagellates investigated, the numbers of auxotrophs for the vitamin B<sub>1</sub> are 49%, which is lower than the percentage of auxotrophs for the vitamin B<sub>12</sub> (91%). In addition, according to Gobler et al. [44], vitamin B<sub>1</sub> is present in coastal waters at concentrations greater than vitamin B<sub>12</sub>, suggesting a possible preference of some dinoflagellates for vitamin B<sub>12</sub> over vitamin B<sub>1</sub> for growth. In that case, given that the vitamin B<sub>12</sub>-dependent species must be able to obtain the vitamin from an external source such as bacteria [46], more detailed studies are required to clarify the relationship between vitamin uptake of *G. catenatum* and the presence of bacteria. Nevertheless, in the addition experiments of vitamin B<sub>1</sub>, an interesting observation was that the growth pattern of *G. catenatum* was similar to those of the N control, and, in treatments 1 and 2, that the senescence phase was observed from day 22 following the stationary phase. This indicates that even in a culture in which N and P are both depleted by uptake, *G. catenatum* does not require vitamin B<sub>1</sub> for growth. However, further studies are still needed to clarify the relationship between the

vitamin requirement of *G. catenatum* and the concentration of N under conditions that other nutritional sources are depleted.

### 3.2. Growth of *Gymnodinium catenatum* under Different Concentrations of Fe, Cu and Se

The growth curves and rates of *Gymnodinium catenatum* culture under different concentrations of trace metals (Fe, Cu and Se) are shown in Figure 3 and Table 2. On additions of Fe and Cu, no significant differences were observed in the growth patterns and rates between control and treatments. For the addition of Fe, *G. catenatum* grew consistently in the control and all treatments for days 1–22 and then entered a stationary phase. A senescence phase was not observed. The growth rates in this experiment were between 0.14 and 0.16 day<sup>−1</sup>. For Cu addition, both the growth curves and the rates (0.14–0.17 day<sup>−1</sup>) were similar to those seen for Fe addition, although *G. catenatum* grew until day 18 before entering a stationary phase. The senescence phase was not observed in this experiment. The growth patterns for the Fe and Cu addition were similar to that for P addition. The maximum cell densities for Fe and Cu addition were 1310 cells mL<sup>−1</sup> and 1071 cells mL<sup>−1</sup>, respectively (Table 2), and both were observed in treatment 2 with additions of  $1.17 \times 10^1$  μM Fe and of  $3.93 \times 10^{-2}$  Cu.



**Figure 3.** Growth curves and rates of *Gymnodinium catenatum* culture exposed to different iron (Fe), copper (Cu) and selenium (Se) concentrations.

The culture media for the control and treatments in these experiments contained N, P, and vitamins, all of which can affect growth and cell density in *G. catenatum*. In this culture, additions of Fe and Cu did not seem to enhance the growth rates of *G. catenatum*, given that there were no significant differences in growth curves or rates between the treatments and controls without the addition of Fe and Cu (Figure 3). Fe and Cu are essential elements for phytoplankton, and play an important role in many of the biochemical and metabolic processes involved in cell growth [47–49]. Some studies recorded that the growth of *Protoceratium reticulatum* and *Alexandrium tamarense* increased with increasing Fe concentrations [50,51], although in contrast the cell growth of *Scrippsiella trochoidea* was inhibited by an increase in Fe [52]. Other studies reported a significant negative effect of Cu on the growth of dinoflagellates such as *A. catenella*, *Ostreopsis cf. ovata* [35,53]. These findings indicate that Fe and Cu requirements can vary greatly among phytoplankton. The basic knowledge of the Fe and Cu requirements of *G. catenatum* is lacking to date; however, our results indicate that elevated trace metal (Fe and Cu) concentrations do not increase the growth and cell density of *G. catenatum* in nutrient rich waters.

There has been considerable interest in the Se requirements of harmful algal species (e.g., [32,33,50,54]), suggesting that the addition of Se to a culture medium stimulates growth. In

particular, culture experiments show a clear requirement for Se by *G. catenatum* [54]. In our experiment, Korean isolates of *G. catenatum* could grow in an f/2 culture medium without Se addition (referred to as the control), however the growth was more elevated when Se was added (Figure 3). The maximum cell densities in treatments 3, 4 and 5 were twice as high as that of the control, which were also higher than those of other nutrient and trace metal sources in this study (Table 2). There were no significant differences in growth rates ( $0.15\text{--}0.17\text{ day}^{-1}$ ) between the control and the treatments. In both the control and the treatments, *G. catenatum* grew until day 16 or 20 before entering a stationary phase, and a senescence phase was not observed. This growth pattern was similar to those seen in the P, Fe and Cu addition experiments.

According to Boisson et al. [55] and Price et al. [56], Se is either essential for some phytoplankton in low concentrations or it cannot enhance growth of phytoplankton. This indicates that Se requirements can vary among phytoplankton, and that phytoplankton can have an optimal range of Se for growth. In Se addition, no obvious differences in cell densities of *G. catenatum* were observed until the end of the exponential growth phase, resulting in similar growth rates between the control and treatments. However, cell densities were increased in the stationary phases of treatments 3, 4 and 5 with addition of Se. This result suggests that concentrations greater than  $10^{-3}\text{ }\mu\text{M}$  of Se can enhance cell densities or biomass yield in culture experiments, although we did not measure the concentration of Se of the seawaters used for making the solutions. According to Doblin et al. [32], the Se requirement of *G. catenatum* varies between strains obtained from Australia, Spain and Japan, with addition of selenium at concentrations in the range  $10^{-9}\text{--}10^{-7}\text{ M}$  causing a variable increase in growth and biomass yield. These Se requirements also differ from that of the Korean isolate of *G. catenatum*. It is thus possible that the growth of *G. catenatum* can be enhanced for a wide range of concentrations of Se.

### 3.3. Proliferation Potential of *Gymnodinium catenatum* and Nutritional Conditions in Korean Coastal Waters

According to Band-Schmidt et al. [57], blooms caused by *G. catenatum* have been associated with an increase in nutrients, mainly by nitrogen compounds in the water column. In our study, N seems to be one of the most important nutritional factors in the growth of the Korean isolate of *G. catenatum*, however the *G. catenatum* prefers high concentrations of P and Se, rather than preference to N. In a previous study, it was reported that the maximum cell density of the Korean isolate of *G. catenatum* in laboratory experiments was observed at  $20\text{ }^{\circ}\text{C}$ , which is commonly recorded in summer in Korea [38]. According to Koo [58], in the southern coastal area of Korea from which the isolate of *G. catenatum* was obtained, average N and P concentrations were  $0.8$  and  $0.1\text{ }\mu\text{M L}^{-1}$  in summer, respectively. This indicates that P can be a limiting factor for phytoplankton growth in Korean coastal waters, and because of this, in summer *G. catenatum* may prefer high concentration of P for the growth.

According to Koo [58], Cho [59] and Jang et al. [60], low levels of N and P in summer could be a result of the uptake of dinoflagellates, among which *Margalefidinium polykrikoides* (= *Cochlodinium polykrikoides*), *Prorocentrum obtusidens* (= *P. donghaiense*), and *Alexandrium* species are well known to dominate in Korean coastal waters in summer, causing dense blooms (e.g., [61–64]). The maximum growth rates of these species in the laboratory are higher than that of *G. catenatum* (e.g., [65–67]), indicating that *G. catenatum* may be at a disadvantage beside the more the dominant species in terms of nutrient competition. Consequently, proliferation of *G. catenatum* is not expected in summer of Korean coastal area. However, as during summer, after rainfall events, riverine input of inorganic Se and its interaction with dissolved organic matter (DOM) can be a critical factor for bloom initiation of *G. catenatum* [34], this species may have a competitive advantage if Se, P and DOM loading after rainfall is increased in Korean coastal waters.

**Author Contributions:** Data curation, formal analysis, writing-original draft and writing-review & editing, K.H.H. and H.H.S.; Data curation and formal analysis, H.J.K.; Data curation and writing-review & editing, Z.L.; Data curation and investigation, J.Y.Y. and K.Y.K.; Formal analysis, M.H.S., J.H., S.D.L. and S.M.Y.; Writing-review & editing, S.J.O., J.W.P. and W.-A.L.; Conceptualization and funding acquisition, H.H.S. All authors have read and agree to the published version of the manuscript.



**Funding:** This work was supported by grants from NIFS (R2020041), KIOST (PE99821) projects, and NNIBR project (202001103).

**Conflicts of Interest:** The authors declare no conflict of interest.

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