

Growth responses of *Chondrus ocellatus* Holmes (Gigartinales, Rhodophyta) to two endophytes, *Mikrosyphar zosterae* Kuckuck (Ectocarpales, Ochrophyta) and *Ulvella ramosa* (N. L. Gardner) R. Nielsen (Ulvales, Chlorophyta) in culture

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To examine the effects of two endophytic algae, *Mikrosyphar zosterae* (brown alga) and *Ulvella ramosa* (green alga), on the host *Chondrus ocellatus* (red alga), culture experiments were conducted. Four treatments were made: endophyte-free (*Chondrus* only), endophyte-*M* (*Chondrus* + *Mikrosyphar*), endophyte-*U* (*Chondrus* + *Ulvella*), and endophytes-*M·U* (*Chondrus* + *Mikrosyphar* + *Ulvella*). After 3 weeks, the relative growth rates (RGRs) of frond lengths and the number of newly formed bladelets were examined. *M. zosterae* formed wart-like dots on *C. ocellatus* fronds, whereas *U. ramosa* made dark spots. The RGRs of frond lengths of *C. ocellatus* were significantly greater in the endophyte-free and endophyte-*M* treatment groups than in the endophyte-*U* and endophytes-*M·U* treatment groups, indicating that the growth of host *C. ocellatus* was inhibited more by the green endophyte *U. ramosa* than the brown endophyte *M. zosterae*. The number of newly produced bladelets was greater in the endophyte-*U* and endophytes-*M·U* groups than in the endophyte-free and endophyte-*M* treatment groups. These results indicate that the two endophytes inhibit growth of the host *C. ocellatus*. The negative effects of *U. ramosa* on *C. ocellatus* growth were more severe than those caused by *M. zosterae*. Furthermore, *U. ramosa* destroyed the apical meristems of *C. ocellatus*, whereas *M. zosterae* did not. On the other hand, *C. ocellatus* showed compensatory growth in the form of lateral branch production as *U. ramosa* attacked its apical meristems.

Key words: *Chondrus ocellatus*; growth; infection; *Mikrosyphar zosterae*; *Ulvella ramosa*

INTRODUCTION

Chondrus, a red algal genus, is an important primary producer providing food and habitat for various marine organisms as many seaweeds do (Lindstrom 2009, Choi et al. 2015a). This alga is also economically important as a major source of carrageenan (Necas and Bartosikova 2013, Zhou et al. 2014). However, natural populations of

Chondrus have declined due to a variety of abiotic (e.g., pollution, ocean acidification, and typhoons) and biotic environmental factors, including predation by herbivores and pathogens (Correa and McLachlan 1992, 1994). Production of natural or cultivated seaweeds has been diminished by seaweed diseases caused by fungi, bacteria,



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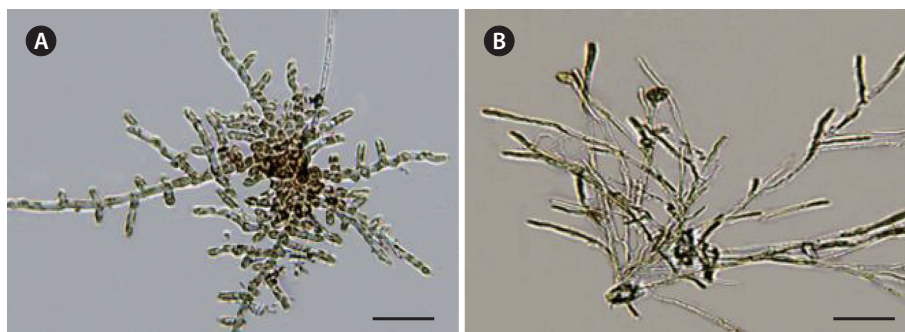


Fig. 1. The two endophytes isolated in laboratory cultures. (A) *Mikrosyphar zosterae*. (B) *Ulvella ramosa*. Scale bars represent: A & B, 50 μ m.

viruses, and endophytic algae (Gachon et al. 2010, Potin 2012, West et al. 2013). Endophytic algae can be infectious and can negatively affect the performance of host seaweeds (Apt 1988, Correa and McLachlan 1992, 1994, Correa et al. 1997). The morphological symptoms of infection with endophytes in host seaweeds include thallus thickening, degradative lesions, cellular damage, gall formation, and morphological deformations (Yoshida and Akiyama 1979, Apt 1988, Correa et al. 1988, 1997, Garbary et al. 2014, Preuss and Zuccarello 2014). The endophytic infection may even reduce the survival, growth, and fertility of host seaweeds (Faugeron et al. 2000, Schoenrock et al. 2013).

Correa and his colleagues conducted intensive studies on endophytes in a red alga, *C. crispus* (Correa et al. 1988, 1994, Correa and McLachlan 1992, 1994). They found four endophytic species, including three green endophytes, *Phaeophila dendroides*, *Endophyton ramosum* (currently regarded as *Ulvella ramosa*) and *U. operculata* (previously known as *Acrochaete operculata*), and one brown endophyte, *Streblonema* species. Among these endophytes, *U. operculata* caused depigmentation, cellular disorganization, and wart-like protrusion in *C. crispus* (Correa et al. 1994). Interestingly, in the presence of *U. operculata*, bacteria accelerated cell wall degradation, resulting in the fragmentation and death of *C. crispus* (Correa et al. 1988, Correa and McLachlan 1992, 1994). As a protective mechanism, *C. crispus* gametophytes produced kappa-carrageenan oligosaccharides to inhibit infection of endophytic *U. operculata* (Bouarab et al. 2001, Weinberger 2007).

Chondrus ocellatus, which grows on semi-exposed rocky shores, is a bioengineer enhancing species diversity as primary producer and food resources of various marine organisms, and is a potential carrageenophyte in Korea (Kim et al. 2006, Zhou et al. 2014). Lee et al. (2013) reported green endophytes in *C. ocellatus*, without iden-

tifying the species and a green endophytic *U. ramosa* was found in *C. ocellatus* fronds by Choi et al. (2015b). We also isolated two endophytes, *U. ramosa* and *Mikrosyphar zosterae* from *C. ocellatus* fronds. This study aimed to examine the effects of endophytic *U. ramosa* and *M. zosterae* on the growth of their host *C. ocellatus*, morphological changes of the host seaweed following infection with two infectious endophytes, and the importance of apical growth tips in *C. ocellatus*.

MATERIALS AND METHODS

Sampling

Chondrus ocellatus samples were randomly collected from the low intertidal zone of Manripo, Taean, Korea (36°78' N, 126°8' E) in July 2015. The samples were put in labelled plastic bottles, placed in a cooler, and transported to the laboratory. All samples were classified into abnormal fronds with black spots (or wound, and whitening, etc.) as an indication of endophyte infection and healthy fronds with no endophyte infection.

Endophyte isolation, uni-culture, and identification

Disc-shaped samples of *C. ocellatus* fronds with black spots were excised using a cork borer (1 cm in diameter). The discs were rinsed for 30 s with running tap water and immersed in a 0.5% hypochlorite sodium (NaClO) solution for 30 s in order to remove microorganisms. The discs were then rinsed several times with autoclaved seawater (Gauna et al. 2009). The discs with black spots were placed in petri dishes (size: 15 mm \times 60 mm) containing 10 mL of Provasoli's enriched seawater (PES) medium (Provasoli 1968) and cultured in an incubator at 20°C and with 20

$\mu\text{mol photons m}^{-2} \text{s}^{-1}$, and a photoperiod of 12 : 12 h LD cycle. Light was provided by 40 W cool-white fluorescent tubes and the level of irradiance was measured using a digital illumination meter (DX-200; INS Enterprise Co., Taipei, Taiwan). The growth of diatoms was inhibited by adding GeO_2 ($3 \mu\text{g L}^{-1}$) in culture medium during the first week of cultivation to reduce any negative effects of GeO_2 (Fernandes et al. 2011). Filamentous brown *M. zosteræ* and green *U. ramosa* endophytes appeared on the surface of frond discs within 3 to 4 weeks of cultivation. The endophytes were isolated, stock-cultured, and identified as *M. zosteræ* and *U. ramosa* based on molecular analysis (using the RuBisCo spacer and tufA for the *M. zosteræ* and *U. ramosa* analyses, respectively) and morphological features (Fig. 1A & B).

Preparation of *Chondrus ocellatus* juveniles

Healthy and fertile carposporophytes (>10 cm in length) of *C. ocellatus* were cleaned in autoclaved seawater and immersed in 1.5% KI solution for 20 min to remove surface microorganisms without releasing damaged spores (Wang et al. 2006, Li et al. 2010). The fronds were rinsed again several times in the autoclaved seawater, and several hundred frond discs bearing cystocarps were obtained using a cork borer. The discs were dried at room temperature (20°C) for 1 h and submerged in autoclaved seawater to induce carpospore release.

Carpospores were inoculated into Petri dishes containing six cut microscope slides (25×20 mm) and PES medium (40 mL), and then were left overnight to settle. The following day, the slides bearing sporelings were transferred to a culture dish (size: 40 mm \times 100 mm) containing 100 mL of PES medium and GeO_2 ($3 \mu\text{g L}^{-1}$). Sporelings were cultured at 20°C, $60 \mu\text{mol photons m}^{-2} \text{s}^{-1}$ and 12 : 12 h LD cycle. The culture medium was replaced weekly.

Endophyte effect on the growth of *Chondrus* juveniles

To examine the effects of the endophytes on the growth of *C. ocellatus* juveniles, 48 *C. ocellatus* juveniles (9-11 mm in length) were prepared. They were allocated into four endophyte treatments, with each endophyte treatment having 12 *C. ocellatus* juveniles that were placed in plant culture dishes containing 50 mL of autoclaved seawater each. The endophyte-free treatment (control) had *Chondrus* juveniles only. Three mixed cultures of endophyte-*M* (*C. ocellatus* + *M. zosteræ*), endophyte-*U* (*C. ocellatus* + *U. ramosa*), and endophytes-*M-U* (*C. ocellatus* + *M. zos-*

teræ + *U. ramosa*) were also prepared. Endophyte suspensions were made by mixing the specified endophytes grown in an area of 4 cm² and blended for 20 s with 100 mL of autoclaved seawater. Endophytic species grown in Petri dishes were carefully detached by a blade prior to make endophyte suspension. The *Mikrosyphar-Ulvella* endophyte suspension was prepared with *U. ramosa* (2 cm²) and *M. zosteræ* (2 cm²). For each treatment, an endophyte suspension (100 mL) of *M. zosteræ*, *U. ramosa*, or *Mikrosyphar-Ulvella* was inoculated into each plant culture dish containing 12 *C. ocellatus* juveniles. The control dishes with no endophyte (*C. ocellatus* juveniles only) were inoculated with 100 mL autoclaved seawater.

During the 1-week culture period, endophyte-*M*, endophyte-*U*, and endophytes-*M-U* grew and released zoospores into their respective plant culture dishes. Zoospore release from the experimental endophytes was confirmed under a light microscope using the cut glass slides that had been inserted into the cultures at the beginning of the experiment. After 1 week, 4 *C. ocellatus* juveniles from the 12 juveniles in each endophyte treatment group were selected and placed into three replicate beakers, each containing 100 mL of PES medium. The control dishes having 12 *C. ocellatus* juveniles were also treated like the other endophytic treatments and absence of endophytes was confirmed with the cut glass slides placed on the bottom of the culture dishes. After this, the samples were cultured for 3 weeks at 20°C and with $60 \mu\text{mol photons m}^{-2} \text{s}^{-1}$ and a 12 : 12 h LD cycle. The culture medium was replaced weekly.

All the experimental *C. ocellatus* juveniles were weekly cleaned by using a paint brush, photographed, and measured in frond length using Image J ver. 1.48 software (National Institutes of Health, Bethesda, MD, USA). We supposed that cleaning activity of the juveniles was influenced with same intensity for endophyte infection and proliferation. New proliferations were counted after 3 weeks. Relative growth rates (RGRs) of fronds (lengths) were estimated by the following formula (Kim et al. 2006):

$$\text{RGR (\% d}^{-1}\text{)} = 100 \ln (L_t / L_0) / t$$

, Where L_0 is the initial frond length, L_t is the final length after t days of cultivation.

Morphological symptoms of infection in host *C. ocellatus* juveniles were also inspected by the naked eye and under an Olympus BX53 microscope (Olympus, Tokyo, Japan). *C. ocellatus* tissues, either infected or healthy, were photographed under microscope after semi-thin cryosection. Prior to sectioning, samples were fixed in

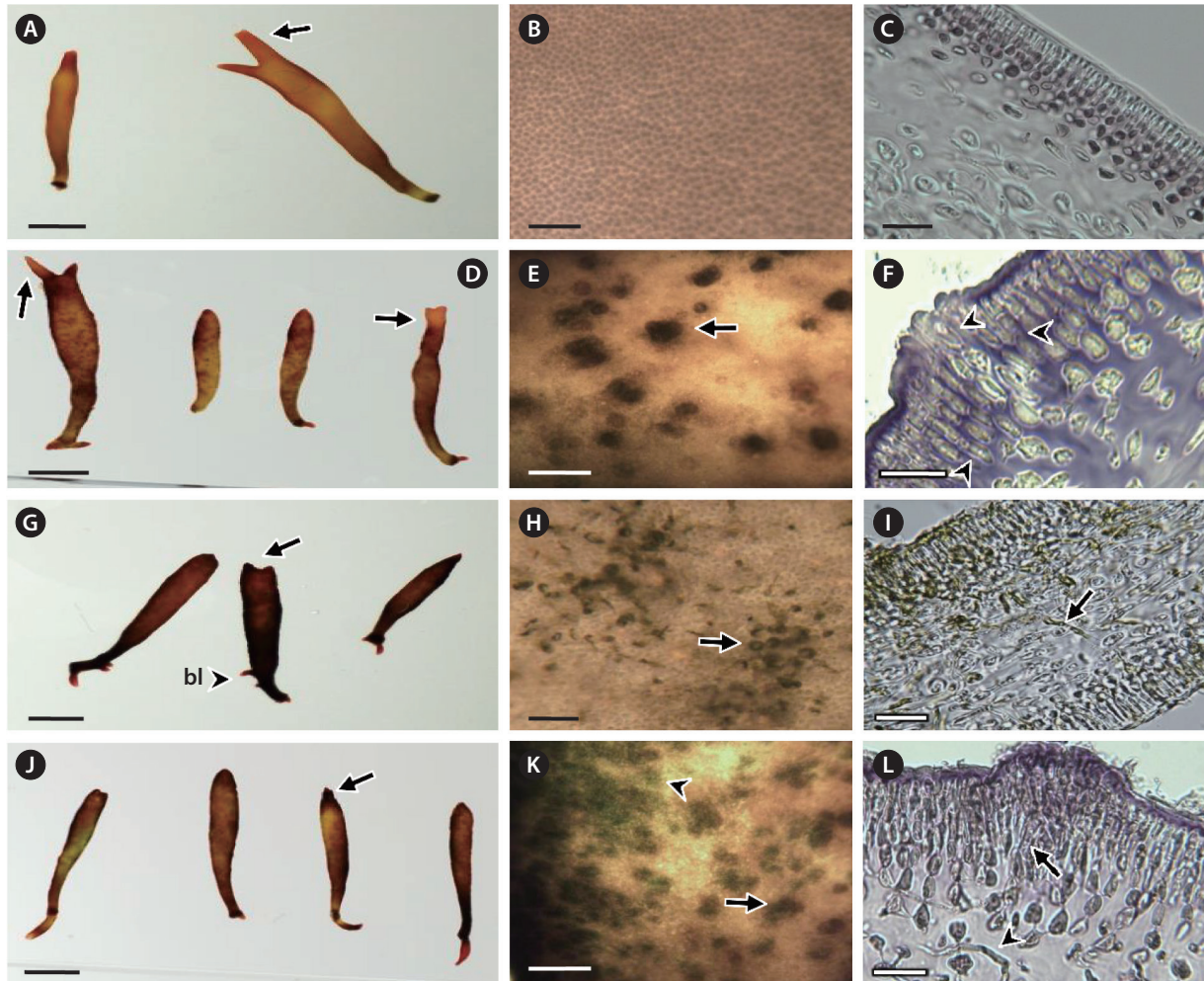


Fig. 2. Morphological symptoms of *Chondrus ocellatus* juveniles infected with the endophytes *Mikrosyphar zosterae* and *Ulvella ramosa*. (A) Non-infected *C. ocellatus* with healthy apices (arrow). (B) Clean host surface tissues. (C) Cross-sectioned tissues of non-infected *C. ocellatus*. (D) *C. ocellatus* juveniles infected with *M. zosterae* and with healthy apices (arrows). (E) Wart-like spot (arrow) on *C. ocellatus* surface tissues caused by *M. zosterae*. (F) *M. zosterae* filamentous thalli (arrowheads) in cross-sectioned host tissue. (G) *C. ocellatus* juveniles infected with *U. ramosa* and with coated apices (arrow) and a bladelet (bl) on the host frond. (H) Green patches (arrow) on host surface tissue caused by *U. ramosa*. (I) *U. ramosa* filamentous thalli (arrow) in a cross-sectioned host frond. (J) *C. ocellatus* juveniles infected by *U. ramosa* and *M. zosterae* with coated apices (arrow). (K) Green patches (arrowhead) and wart-like spots (arrow) on host tissues. (L) Filamentous thalli of *U. ramosa* (arrowhead) and *M. zosterae* (arrow), in cross-sectioned host infected tissues. Scale bars represent: A, D, G & J, 3 mm; B, 50 μ m; C, H & I, 30 μ m; E & K, 300 μ m; F & L, 20 μ m.

a 10% formalin-seawater solution and then embedding medium (Tissue-Tek O.C.T. compound; Sakura Finetek USA Inc., Torrance, CA, USA) in a cryochamber at -30°C . Sections were made with a Shandon Cryotome Cryostat microtome (Shandon-Thermo Fisher Scientific, Waltham, MA, USA) and stained with a 10% hematoxylin solution (Mayer's hematoxylin; Dako North America Inc., Carpinteria, CA, USA).

Importance of apical growth tips of *Chondrus* juveniles

The endophyte infection experiments showed that *U. ramosa* endophytes infected selectively the apical growth tips of host *C. ocellatus* resulting in the reduction of the growth of *C. ocellatus* juveniles. Therefore, an additional experiment on the role of apical growth tips in *Chondrus* juveniles were designed. Twelve *C. ocellatus* juveniles were divided into apices (from tip to middle) and bases (from middle to base). Twelve intact juveniles, with both apices and bases were used as a control. Three replicate

beakers for each treatment, containing four *C. ocellatus* juveniles and 100 mL of PES medium each, were prepared. The parts of *C. ocellatus* juveniles (apex, base, or intact) were cultured at 20°C. The irradiance and photoperiod were 60 $\mu\text{mol photons m}^{-2} \text{s}^{-1}$ and 12 : 12 h LD cycle, and the culture medium was replaced weekly. Because the *C. ocellatus* juveniles were so small, total frond weights of each replicate (comprising four juveniles) were measured at the beginning of the experiment and after 3 weeks of cultivation. Frond RGRs (weights) of *C. ocellatus* juveniles were calculated using the equation shown above. The total number of new proliferations from the four juvenile fronds was also counted.

Data analysis

A one-way ANOVA was applied to evaluate the effect of endophytes on the RGRs of fronds (lengths) and the number of new proliferations in *C. ocellatus* juveniles. A one-way ANOVA was also used to examine the effect of apical meristems on the RGRs (frond weight) and the number of new proliferations in *C. ocellatus* juveniles. Homogeneity of variances was tested using Cochran's test. Tukey's honest significant difference (HSD) test was applied when significant differences between means were detected (Sokal and Rohlf 1995). Statistical analyses were carried out using STATISTICA ver. 7.0 software.

RESULTS

Symptoms in the infected *Chondrus*

C. ocellatus juveniles grew well for 3 weeks, without morphological changes (Fig. 2A-C), in the absence of the brown endophyte *M. zosterae*. When cultured with *M. zosterae*, host *C. ocellatus* formed black spots on their fronds except for apical areas (Fig. 2D). *C. ocellatus* also developed many wart-like spots (Fig. 2E) where endophytic filaments penetrated the cortex of *C. ocellatus* to the upper medulla zone (Fig. 2F). The apical tips of *C. ocellatus* juveniles were coated by *U. ramosa* endophytes (Fig. 2G), and dark green patches were found on the surfaces of whole host fronds (Fig. 2H). This green *U. ramosa* endophyte was also found in host tissues (Fig. 2I). In the endophytes-*U-M* treatment, green patches and wart-like spots were observed on the *C. ocellatus* fronds (Fig. 2J & K). Endophytic filaments of both species, *U. ramosa* and *M. zosterae*, were also observed on fronds and in host tissues (Fig. 2K & L).

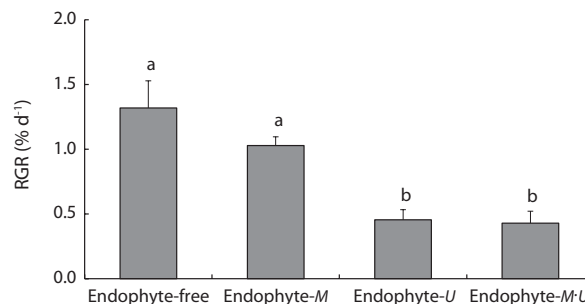


Fig. 3. Relative growth rates (RGRs, frond lengths) of *Chondrus ocellatus* juveniles, which cultured for 3 weeks in different endophytic treatments (endophyte-free, endophyte-*M*, endophyte-*U*, and endophytes-*M-U*). Different letters indicate significant differences between the groups determined by Tukey's honest significant difference test. Bars indicate the mean \pm standard error ($n = 3$). Endophyte-free, *Chondrus* only; endophyte-*M*, *Chondrus* + *Mikrosyphar*; endophyte-*U*, *Chondrus* + *Ulvella*; endophytes-*M-U*, *Chondrus* + *Mikrosyphar* + *Ulvella*.

Effects of two endophytes on *Chondrus* growth

Chondrus ocellatus juveniles grew with infectious *M. zosterae* or *U. ramosa*, or with both. In 3 weeks of culture after infection, the RGR was significantly greater in the endophyte-free and endophyte-*M* treatment groups than in the endophyte-*U* or endophytes-*M-U* treatment groups (Fig. 3). Interestingly, no differences were found between the endophyte-free and endophyte-*M* treatment groups, and between the endophyte-*U* and endophytes-*M-U* treatment groups (Fig. 3), which indicates that the significant negative effects on the growth of host *C. ocellatus* juveniles were attributable to infection with *U. ramosa*.

Three weeks after infection with the endophytes, no *C. ocellatus* bladelets developed in the endophyte-free and endophyte-*M* treatment groups, 1.33 ± 0.33 (mean \pm standard error [SE], $n = 3$) bladelets formed in the endophyte-*U* treatment group, and 1.67 ± 0.66 ($n = 3$) bladelets formed in the endophytes-*M-U* treatment group. The number of newly formed bladelets appeared to be significantly greater in the endophyte-*U* and in the endophytes-*M-U* treatment groups as compared to the control (endophyte-free) or endophyte-*M* treatment groups ($p < 0.05$). However, no significant differences were found between the endophyte-free and endophyte-*M* treatment groups and between the endophyte-*U* and endophytes-*M-U* treatment groups.

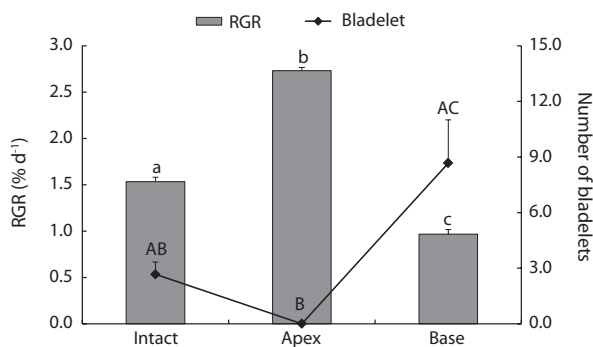


Fig. 4. Relative growth rates (RGRs, frond weights) of *Chondrus ocellatus* juveniles, and the number of bladelets after 3 weeks of cultivation at three different treatments (intact, apex, and base). Different letters indicate significant differences between the groups determined by Tukey's honest significant difference test. Bars indicate the mean \pm standard error ($n = 3$).

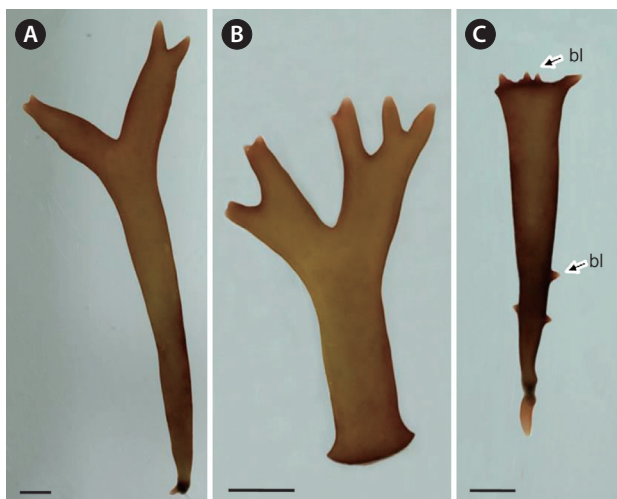


Fig. 5. Growth patterns of *Chondrus ocellatus* juveniles after 3 weeks in laboratory culture. (A) An intact juvenile as a control. (B) An apex frond without bladelets from a juvenile. (C) A basal frond producing many small bladelets (bl) on the excised part and lateral surface (arrows) of a juvenile frond. Scale bars represent: A-C, 1 mm.

Importance of apical meristems in *Chondrus* juveniles

After 3 weeks of culture, the RGR values (frond weights) of *C. ocellatus* juveniles ranged from 0.97 to 2.73% d⁻¹ (Fig. 4). The values were significantly different between the three treatments, with the highest RGR value in the apex treatment and the lowest in the base treatment (Fig. 4). In addition, the number of newly formed bladelets was significantly different between the treatment groups ($p <$

0.05). A Tukey's test revealed a significant difference between the apex and base groups, but no differences were found between the intact and apex groups, and between the intact and base groups (Fig. 4). Intact juveniles of *C. ocellatus* produced a few bladelets (Fig. 5A), whereas the apex alone produced no bladelets (Fig. 5B). However, the base of *C. ocellatus* juveniles produced many bladelets in the excised, lateral parts of fronds (Fig. 5C). The number of newly formed *C. ocellatus* bladelets was 2.67 ± 0.67 (mean \pm SE, $n = 3$) in the intact control and 8.67 ± 2.33 ($n = 3$) in the base treatment group. This result indicates that *C. ocellatus* juveniles grow apically, but that their growth can switch to bladelet production in the absence of apical meristems.

DISCUSSION

Previous studies have shown that *C. crispus* harbors endophytes such as brown *Streblonema* spp. and green *U. ramosa* and *U. operculata* (Correa et al. 1988). Choi et al. (2015b) recently isolated a green endophyte, *U. ramosa*, from *C. ocellatus*. Here, we report that the brown endophyte *M. zosterae* was present in *C. ocellatus*. This is the first report of *M. zosterae* as an endophyte of a carrageenophyte. We also found that morphological symptoms in *C. ocellatus* of infection with the two endophytes, green *U. ramosa* and brown *M. zosterae*, were different. *U. ramosa* infection produced dark green patches, whereas *M. zosterae* infection resulted in wart-like spots. Various green endophytic species in several carrageenophytes induce color changes, form holes, or have other effects on host plants (Table 1).

Although endophytes can cause some morphological changes in seaweeds, the causative organisms forming galls (hyperplasia and hypertrophy in host seaweed cells) on red algal fronds were found to be mostly bacteria, fungi, animals, or viruses (Apt and Gibor 1991, Correa et al. 1993, Goecke et al. 2012). On *C. crispus* fronds, galls formed in response to nematodes and bacteria (Apt 1988). Another red alga, *Iridaea laminarioides* (currently regarded as *Mazzaella laminarioides*) also developed galls on its fronds because of infection with the cyanobacterium *Pleurocapsa* sp. (Correa et al. 1993). The findings in the present study indicate that infection with the endophyte *M. zosterae* can also result in the formation of gall-like structures in the red alga *C. ocellatus*.

In the present study, the infectious endophytes inhibited the growth of host *C. ocellatus* juveniles, but the degrees of negative effects were different between the endo-

phytes, *M. zosteræ* and *U. ramosa*. *C. ocellatus* juveniles grew significantly better in the endophyte-free (1.32% d⁻¹ in RGR) and endophyte-*M* treatment (1.03% d⁻¹) groups than in the endophyte-*U* (0.45% d⁻¹) and endophytes-*M·U* treatment (0.43% d⁻¹) groups. In comparison to the RGR values with endophyte-free treatment, the RGR of the host *C. ocellatus* was reduced by approximately 70% with endophyte-*U* treatment. In this condition, *U. ramosa* densely covered the apical meristem tissues of *C. ocellatus*, perhaps reducing photosynthetic activity.

This study is the first report that the endophyte *U. ramosa* damages apical meristem tissues of *C. ocellatus* juveniles. The growth of *C. ocellatus* was mainly apical, and transverse divisions in apical cells are common. Apical division of *C. ocellatus* was not observed in the endophyte-*U* treatment or endophytes-*M·U* treatment groups (Fig. 2). This resulted in a significant reduction in the RGRs in these conditions. Removal of apical meristems from *C. ocellatus* juveniles reduced growth by 37% compared to that in the intact treatment group and by 65% compared to that in the apex treatment group. This result implies the importance of apical meristems to the growth of *C. ocellatus*. Previously, *C. crispus* were shown to grow three times more in apical meristems than in other parts of the organism (Chopin and Floc'h 1992).

Interestingly, defects in the apical meristems of *C. ocellatus* resulted in the formation of bladelets. Surface and marginal proliferations are commonly observed in adult-stage reproductive organs in *Chondrus* species, including

C. nipponicus and *C. ocellatus* (Brodie et al. 1991, 1993). In this study, we found that *C. ocellatus*, even in the juvenile stage, produced small bladelets (≤ 2.1 mm in length) in the absence of apical growth tips or with apical tissue damaged by *U. ramosa*. Formation of many bladelets in the damaged fronds or in the basal fronds of *C. ocellatus* may be a mechanism to compensate for the lack of growth in the removed apical meristems. Similarly, *Fucus vesiculosus* produced many lateral branches that served as new apical meristems when their apices were damaged (Honkanen and Jormalainen 2002). *Chondrus crispus* was also shown to produce many bladelets (Corey et al. 2012). Corey et al. (2012) reported that bladelet development in *C. crispus* was more pronounced at temperatures of 17°C or above, and high temperatures may be optimal for the growth of the green endophyte *U. operculata* or *U. ramosa*, or both (Correa et al. 1988).

Here, we reported *M. zosteræ* as a new endophyte of *C. ocellatus*. What we do not know is the level of host-specificity of *M. zosteræ*. Although *M. zosteræ* formed wart-like spots on *C. ocellatus*, this endophyte did not cause significant negative effects on the growth of the host alga. Therefore, there is a need to analyze more seaweed species, especially carrageenophytes to determine the host-specificity of *M. zosteræ*. It is also important to determine whether different life stages of *C. ocellatus* respond differently to these endophytes, because some endophytes (e.g., *U. operculata*) affect the sporophytic fronds of *C. crispus* more severely than its gametophytes (Correa and

Table 1. Morphological symptoms of carrageenophytes in response to infection with a variety of green endophytic algae

Host seaweed	Endophyte	Symptom	Reference
<i>Grateloupia</i> sp.	<i>Blastophysa rhizopus</i>	Green spot rotting	Iima and Tatewaki (1987)
<i>Chondrus crispus</i>	<i>Acrochaete operculata</i> , <i>Acrochaete heteroclada</i> , <i>Phaeophila dendroides</i>	Dark patches Lesions Tissue softening	Correa et al. (1988)
<i>Iridaea cordata</i>	<i>Acrochaete heteroclada</i> <i>Acrochaete operculata</i>	Dark patches Lesions Tissue softening	Correa and McLachlan (1991)
<i>Iridaea laminarioides</i>	<i>Endophyton</i> sp.	Green patches	Correa et al. (1994)
<i>Mazzaella laminarioides</i>	<i>Endophyton ramosum</i>	Dark green	Sánchez et al. (1996)
<i>Chondrus crispus</i>		Lesions	
<i>Rhodoglossum californicum</i>		Tissue softening	
<i>Erythrodermis traillii</i>		Discoloration	
<i>Phyllophora pseudoceranioides</i>		Cell necrosis	
<i>Ahnfeltiopsis devoniensis</i>			
<i>A. furcellata</i>			
<i>Grateloupia lanceolata</i>	<i>Ulvela eptochaete</i> , <i>Blastophysa rhizopus</i> , <i>Bolbocoleon piliferum</i>	Light-green spots, Hole formation, Discoloration Cell necrosis	Kim et al. (2014)

Acrochaete operculata = *Ulvela operculata*, *Acrochaete heteroclada* = *Ulvela heteroclada*, *Endophyton ramosum* = *Ulvela ramosa*.

McLachlan 1994). The endophytes found in this study, *M. zosterae* and *U. ramosa*, are pathogens of *C. ocellatus* that result in distinctive symptoms, and, in the case of *U. ramosa*, decrease growth of the host seaweed by damaging meristem tissues.

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