Effect of red tide dinoflagellate diet and cannibalism on the bioluminescence of the heterotrophic dinoflagellates Protoperidinium spp.

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ABSTRACT. The effects of diet and cannibalism were assessed from changes in the bioluminescence potential of 2 species of the heterotrophic dinoflagellate Protoperidinium fed 4 species of red tide dinoflagellate prey and also maintained without added prey. The use of bioluminescence as a sensitive indicator of nutritional status and feeding was explored. The bioluminescence of Protoperidinium cf. divergens and P. crassipes was significantly affected by dinoflagellate diet. Total mechanically stimulable luminescence (TMSL) of P. cf. divergens fed different dinoflagellate diets was significantly correlated with feeding frequency (the percent of feeding P. cf. divergens cells) rather than with population growth rate. P. cf. divergens displayed high levels of TMSL and feeding frequency on a diet of Scripsiosiella trochoidea which did not support population growth. Diet did not affect the total number of flashes produced per cell; therefore, changes in TMSL with dinoflagellate diet were related to the amount of chemical substrate available for luminescence, rather than changes in the excitation/transduction process. Individually isolated cells remained viable for only 3 to 5 d without food and exhibited reduced bioluminescence. However, cells maintained in groups survived at least 16 d without added prey and maintained levels of bioluminescence similar to those during favorable prey conditions. Cannibalism observed during this time may have enabled cells of P. cf. divergens to feed and therefore produce high levels of bioluminescence in the absence of added prey. Changes in swimming speed were less than changes in bioluminescence. The results of the present study suggest that energy utilization may be prioritized in the following order: swimming (for grazing) > bioluminescence (for reducing predation) > reproduction (for increasing the population).

KEY WORDS: Bioluminescence Cannibalism Dinoflagellate Energetics Microzooplankton Plankton Predation Red tide

INTRODUCTION

The high abundance and diverse ecological roles of heterotrophic dinoflagellates (Beers et al. 1982, Lessard 1984) attest to their importance in the marine environment. In particular, the ubiquitous heterotrophic dinoflagellate genus Protoperidinium can be an important component in plankton dynamics for the following reasons. First, it is among the most abundant of the >20 μm heterotrophic dinoflagellates found in coastal and oceanic waters (Lessard 1984, Hallegraeff & Reid 1986, Lessard & Rivkin 1986, Jacobson 1987), particularly during some dinoflagellate red tides (Jeong 1995) or diatom blooms (Jacobson 1987). Second, it is not only an important prey for copepods (Giford & Dagg 1991, Jeong 1994a), but also a predator of copepod eggs and early naupliar stages (Jeong 1994b). Third, the diet of Protoperidinium spp. includes a broad range of prey species (Jacobson & Anderson 1986, Buskey et al. 1994, Jeong 1994b, Jeong & Latz 1994). Fourth, in some regions,
they have recently been recognized as major sources of bioluminescence in the mixed layer (Lapota et al. 1989, 1992a, b, Swift et al. 1995).

Bioluminescence in dinoflagellates is believed to serve an anti-predation function by reducing predation pressure (Essias & Curl 1972, White 1979) and by acting as a 'burglar alarm' which increases the vulnerability of the grazer to secondary predation (Meninger & Case 1992, Abrahams & Townsend 1993). Therefore, bioluminescence may be an important factor in predator-prey interactions and the structuring of biological communities.

The bioluminescence capacity of heterotrophic dinoflagellates appears to be dependent on short-term nutritional status, i.e. the total energy resources available for all physiological processes. Nutritional status and bioluminescence are affected by prey concentration (Buskey et al. 1992, 1994), photosynthesis by symbiotic algae if present (Sweeney 1971), and starvation time (Buskey et al. 1992, 1994). Because a luminescent predator usually experiences a variety of prey sources and conditions, bioluminescence must be measured for different nutritional conditions, as well as for a unialgal diet, in order to fully understand the relationships between bioluminescence and prey availability.

In the case of a luminescent predator such as Protoperidinium spp., the energy obtained from grazing is needed not only for metabolism, swimming, and reproduction, but also for light production. Energy is required for synthesis of the luminescent chemistry, its transport through the cytoplasm, and packaging into membrane-bound vesicles called scintillons which are associated with the vacuolar membrane (Nicolas et al. 1991). The energy requirement for producing bioluminescence is unknown, as is the priority of the luminescent system in terms of energy utilization.

The population growth of a predator can be indicative of its nutritional status when feeding on different prey species and concentrations (e.g. Buskey et al. 1994, Jeong & Latz 1994). However, the energy available for bioluminescence and swimming may be similar even when different population growth rates indicate that different amounts of energy are available for reproduction. For example, Protoperidinium cf. divergens has similar maximum feeding frequencies for both the larger prey Gonyaulax polyedra and the smaller prey Scrippsilla trochoidea, even though its population growth rate for a S. trochoidea diet is zero and that for a G. polyedra diet is high (Jeong & Latz 1994). More information is needed on whether a particular diet provides sufficient energy resources for maintenance and bioluminescence, even when insufficient for reproduction. The nutritional status of a predator may be strongly related to survivorship and thus population dynamics. In the present study, the use of bioluminescence of a predator as an indicator for nutritional status and feeding is explored and compared with other indicators such as population growth rate and swimming speed.

Cannibalism in the genus Protoperidinium occurs mainly when its abundance is high and prey abundance is low (Jeong & Latz 1994), a situation which is expected to occur just after some dinoflagellate red tides. Cannibalism may affect the bioluminescence of Protoperidinium under unfavorable prey conditions; changes in bioluminescence under these conditions may possibly provide an indirect measure of the rate of cannibalism. Cannibalism by Noctiluca scintillans is suspected to explain the increased levels of bioluminescence of cells held in groups compared to that of individually maintained cells (Buskey et al. 1992).

The objective of the present study was to test the following hypotheses regarding the relationship between bioluminescence and nutritional status:

$H_0$: The bioluminescence of Protoperidinium cf. divergens is similar for all unialgal dinoflagellate diets.

$H_2$: If there are differences in population growth rates for Protoperidinium cf. divergens feeding on various unialgal red tide dinoflagellate diets, then bioluminescence will be different.

$H_3$: Prey concentration does not significantly affect the bioluminescence of Protoperidinium cf. divergens.

$H_4$: The bioluminescence of Protoperidinium cf. divergens without added prey is not significantly different from that of P. cf. divergens feeding on optimal prey.

**MATERIALS AND METHODS**

**Culture of experimental organisms.** Cultures of Protoperidinium cf. divergens and P. crassipes, which are abundant during some red tides (Jeong 1993), were established from cells collected from the Scripps Pier (La Jolla, CA, USA) during October, 1992. Details of the culturing of these species are described by Jeong & Latz (1994). Cultures were maintained on a unialgal diet of Gonyaulax polyedra in polycarbonate (PC) bottles. Every 3 to 4 d the P. cf. divergens culture was first sieved through 64 µm Nitex mesh to remove debris, then re-sieved so that cells were retained on 53 µm mesh. P. crassipes cells were first sieved through 100 µm mesh to remove debris, then re-sieved through 64 µm mesh to retain cells. The sieving procedure maintained a homogeneous population size distribution and minimized cell size effects on bioluminescence emission.

Prey cultures of the autotrophic or mixotrophic dinoflagellates Gonyaulax polyedra, Gymnodinium sanguineum, Heterocapsa triqueta, Prorocentrum cf. 

...
*Protoperidinium* *divergens* and *Scrippsiella trochoidea*, were grown in 1/4 enriched seawater media (Guillard & Ryther 1962), minus silicate, at room temperature (20 to 23°C) under continuous cool white fluorescent illumination of 5 x 10^13 quanta cm^{-2} s^{-1}. Cultures in exponential growth phase were used for feeding experiments. Cell concentrations were determined from total cell counts of 1 ml aliquots.

**Experimental design.** The initial concentrations of *Protoperidinium cf. divergens* and its prey are given in Table 1. For all experiments, cells between 53 and 64 μm in diameter were chosen in order to minimize effects due to cell size. Expt 1 was designed to test H01 and H02 (effect of prey species). The initial concentrations of algal prey, based on the results of Jeong & Latz (1994), were obtained by volume dilution, and were chosen for the following reasons: a *Gonyaulax polyedra* concentration of 2000 cells ml^{-1} results in a maximum growth rate of *P. cf. divergens*; a *Gymnodinium sanguineum* concentration of 700 cells ml^{-1} results in a positive growth rate, while that of 2000 cells ml^{-1} results in a negative growth rate. A *Scrippsiella trochoidea* diet results in zero growth for cell concentrations between 1000 and 6000 cells ml^{-1}, although feeding still occurs; therefore, an intermediate concentration of 5000 cells ml^{-1} was used. There is neither feeding by *P. cf. divergens* on *Prorocentrum cf. balticum* nor a positive population growth rate for all prey concentrations tested (Jeong & Latz 1994); therefore, an intermediate prey concentration of 5000 cells ml^{-1} was used.

Dense cultures of *Prorocentrum cf. divergens* and *P. crassipes* were maintained on particular prey for 1 wk, then sieved though 53 and 64 μm mesh, respectively. Retained cells were first transferred to multiwell chambers, then initial concentrations of 1 *Protoperidinium* ml^{-1} were obtained by individually transferring actively swimming cells by a Pasteur pipette into 32 ml polycarbonate (PC) bottles filled with freshly filtered seawater, 1 to 2 ml 1/4 medium, and target prey. For the unfed control condition no prey were added. Bottles were rotated at 0.9 rpm and maintained on a 12 h light:12 h dark (LD) cycle for a 4 d incubation period according to the methods of Jeong & Latz (1994).

Expt 2 was designed to test H03 (effect of prey concentration). Based on feeding studies with 5 dinoflagellate prey species, *Gonyaulax polyedra* was chosen for this experiment because it is the optimal dinoflagellate prey for *Protoperidinium cf. divergens*, with positive population growth for prey concentrations ≥1000 cells ml^{-1} (Jeong & Latz 1994). The initial prey concentrations of 70 to 2500 *G. polyedra* cells ml^{-1} were obtained by volume dilution and the grazer concentration by individually transferring cells as described above. Bottles were rotated as for Expt 1. Because 53 to 64 μm diameter cells were selected for testing, *P. cf. divergens* cell size was similar for each prey concentration treatment (ANOVA, p > 0.05).

Expts 3 and 4 were designed to test H04 (effect of cannibalism). Two methods were used for single-cell starvation studies (Expt 3). All cells of *Protoperidinium cf. divergens* were isolated from cultures maintained on a *Gonyaulax polyedra* diet as described above. Single cells were transferred to individual 10 ml PC bottles which were filled with filtered seawater, tightly capped, and rotated (method 1). Cells were tested for bioluminescence after 0, 3, and 5 d. Other single cells were placed in 7 ml glass scintillation vials partially filled with 4 ml of filtered seawater, loosely capped, and not rotated (method 2). Cells were tested for bioluminescence after 1, 2, and 3 d. For both methods cells were maintained on a 12 h light:12 h dark cycle at room temperature; each cell was tested once.

For group maintained cells without added prey (Expt 4), cells of *Protoperidinium cf. divergens* maintained on a *Gonyaulax polyedra* diet as described above were completely separated from prey cells and debris by sieving through 53 and 64 μm Nitex mesh, respectively. They were then resuspended in filtered seawater in 270 ml PC bottles. Bottles were rotated as for Expt 1. Cells 53 to 64 μm in diameter were tested for bioluminescence after 0, 1, 3, 5, 7, 9, 12, and 16 d.

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**Table 1. Initial concentrations of *Protoperidinium cf. divergens* and prey and incubation times used in each experiment. See text for experimental details.**

<table>
<thead>
<tr>
<th>Expt</th>
<th>Prey species</th>
<th>Prey conc. (cell ml^{-1})</th>
<th>Grazer conc. (cell ml^{-1})</th>
<th>Incub. time (d)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. Effect of unialgal diet</td>
<td><em>Gonyaulax polyedra</em></td>
<td>2000</td>
<td>1</td>
<td>4</td>
</tr>
<tr>
<td></td>
<td><em>Gymnodinium sanguineum</em></td>
<td>700, 2000</td>
<td>1</td>
<td>4</td>
</tr>
<tr>
<td></td>
<td><em>Scrippsiella trochoidea</em></td>
<td>5000</td>
<td>1</td>
<td>4</td>
</tr>
<tr>
<td></td>
<td><em>Heterocapsa triquetra</em></td>
<td>5000</td>
<td>1</td>
<td>4</td>
</tr>
<tr>
<td></td>
<td><em>Prorocentrum cf. balticum</em></td>
<td>5000</td>
<td>1</td>
<td>4</td>
</tr>
<tr>
<td>2. Effect of prey concentration</td>
<td><em>G. polyedra</em></td>
<td>70, 300, 700, 1400, 2000, 3000</td>
<td>1</td>
<td>4</td>
</tr>
<tr>
<td>3. Single-cell starvation</td>
<td>None</td>
<td>1 cell</td>
<td>1, 2, 3</td>
<td></td>
</tr>
<tr>
<td>4. Group maintenance without added prey</td>
<td>None</td>
<td>40</td>
<td>1, 3, 5, 7, 9, 12, 16</td>
<td></td>
</tr>
<tr>
<td>5. Long-term culturing</td>
<td><em>G. polyedra</em></td>
<td>2000</td>
<td>1</td>
<td>130</td>
</tr>
<tr>
<td></td>
<td><em>G. sanguineum</em></td>
<td>750</td>
<td>1</td>
<td>53</td>
</tr>
</tbody>
</table>
Cell size in aliquots maintained under the same conditions was measured using an Elzone model 280PC particle counter with a calibrated 120 μm orifice (Jeong & Latz 1994).

The long-term effect of diet (Expt 5) was investigated to determine if there is a dietary requirement for bioluminescence. Cells of *Protoperidinium cf. divergens* were incubated in 43 ml PC bottles on a diet of luminescent *Gonyaulax polyedra* or a diet of nonluminescent *Gymnodinium sanguineum*. At 4 d intervals, cells were sieved and new prey culture added (Jeong & Latz 1994). The bioluminescence of *P. cf. divergens* cells at Day 0 and Day 50 was measured.

The bioluminescence of freshly collected *Protoperidinium* *cf. divergens* was measured from cells collected in the afternoon by plankton net tows from the Scripps Pier as described above, then individually isolated by Pasteur micropipette. Cells were placed in the dark at local sunset (17:00 h) for bioluminescence measurements.

The bioluminescence of *Protoperidinium crassipes* was tested for a *Gonyaulax polyedra* diet at an initial prey concentration of 2000 cells ml⁻¹, where high population growth occurs (Jeong & Latz 1994).

**Bioluminescence measurements.** Cells were prepared for testing toward the end of the light phase, when bioluminescence is minimally excitable (Biggley et al. 1969, Lapota et al. 1992a). Cells were rinsed by micropipette transfer into filtered seawater, and individually placed into 7 ml glass scintillation vials containing 2 ml filtered seawater. Unless otherwise stated, 20 cells were tested at a room temperature of 20 ± 1°C.

Bioluminescence was measured in a calibrated detection apparatus consisting of a 15 cm diameter integrating sphere collector coupled to a photon-counting photomultiplier (Latz & Lee 1995). Testing was performed during Hours 2 to 4 of the dark phase when high levels of stimulated dinoflagellate bioluminescence occur (Biggley et al. 1969, Lapota et al. 1992a). Single flashes were elicited during continuous stirring of the contents of the vial. Stirring was maintained for 160 s, the length of a data acquisition record, and was repeated until total depletion of luminescence to determine total mechanically stimulable luminescence (TMSL), a measure of bioluminescence capacity.

**Analysis.** Bioluminescence capacity as a function of dinoflagellate diet was determined based on several parameters analyzed as described in Latz & Lee (1995). These parameters provide information on how the packaging and activation of the luminescent system may be affected by nutritional status. Total light emission per cell was expressed as TMSL, which is a function of the total number of flashes produced and the quantum emission per flash. Flash quantum emission is proportional to flash maximum flux (intensity), the intensity of the brightest flash from each cell. The total duration of individual flashes is the sum of rise time (period from initial to maximum flux) and flash decay (period from maximum flux to 3% of maximum) (Latz & Lee 1995). The instantaneous decay rate is the inverse of the e-fold time (duration from maximum flux to e⁻¹ of maximum). Unless otherwise stated, values represent mean ± 1 standard error (SE) of the mean, and statistical differences were tested at the 0.05 significance level with 1-way Analysis of Variance (ANOVA) using Statview software (Abacus Concepts, Inc.). Multiple comparisons were based on the Scheffe test (Winer 1971).

Surface area (SA) of individual cells was calculated according to the formula $SA = 4\pi(D/2)^2$, where $D$ is the equivalent spherical diameter based on the formula $D = (L \times W \times T)^{1/3}$ (Kamykowski et al. 1992), where $L$, $W$, and $T$ are mean cell length, width, and thickness, respectively, measured using an ocular micrometer. Cell volume ($V$) was calculated as $V = \frac{4}{3}\pi(D/2)^3$. Values are expressed as the mean ± SE.

Table 2. Growth rate, maximum feeding frequency and total mechanically stimulable luminescence (TMSL) of *Protoperidinium* of *divergens* results from different heterotrophic dinoflagellate diets in Expt 1. Diets are listed in order of those which resulted in most bioluminescence. ESD: equivalent spherical diameter of cells measured with an electronic cell counter; maximum feeding frequency: instantaneous measure of the percent of the population feeding (based on the presence of a pallium); Growth and feeding data from Jeong & Latz (1994). *TMSL significantly different from that of cells maintained without added prey (Fisher’s PLSD, $p < 0.05$). – no data.

<table>
<thead>
<tr>
<th>Prey species</th>
<th>Prey size (ESD, μm)</th>
<th>Growth rate (d⁻¹)</th>
<th>Maximum feeding frequency (%)</th>
<th>TMSL (quanta cell⁻¹)</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Gonyaulax polyedra</em></td>
<td>37</td>
<td>0.5</td>
<td>27</td>
<td>$6.2 \times 10^9$*</td>
</tr>
<tr>
<td><em>Scripsiella trochoidea</em></td>
<td>19</td>
<td>0</td>
<td>25</td>
<td>$6.0 \times 10^9$*</td>
</tr>
<tr>
<td><em>Gymnodinium sanguineum</em> 700 cells ml⁻¹</td>
<td>32</td>
<td>0.2</td>
<td>20</td>
<td>$5.9 \times 10^9$*</td>
</tr>
<tr>
<td><em>G. sanguineum</em> 2000 cells ml⁻¹</td>
<td>32</td>
<td>-0.2</td>
<td>0</td>
<td>$2.9 \times 10^9$</td>
</tr>
<tr>
<td><em>Heterocapsa triquetra</em></td>
<td>17</td>
<td>0</td>
<td>–</td>
<td>$2.2 \times 10^9$</td>
</tr>
<tr>
<td><em>Prorocentrum</em> <em>cf. balticum</em></td>
<td>11</td>
<td>0</td>
<td>0</td>
<td>$1.3 \times 10^9$</td>
</tr>
<tr>
<td>No prey added</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>$1.9 \times 10^9$</td>
</tr>
</tbody>
</table>
Swimming speed studies. Swimming speeds were measured for Protoperidinium cf. divergens under conditions of no added prey. Measurements were performed at 19 ± 0.5°C using a video microscope setup (Jeong 1994a). Speeds were measured, during single frame playback, from linear displacements of actively swimming cells.

RESULTS

Test of H01 and H02 (effect of prey species)

Expt 1 tested the effect of 5 different unialgal red tide dinoflagellate diets on the bioluminescence of Protoperidinium cf. divergens (Table 2). Because TMSL was significantly different between diets (ANOVA, p < 0.001), H01 can be rejected. However, the total number of flashes produced by each P. cf. divergens cell was similar (ANOVA, p > 0.05), and averaged 11.5 ± 0.9 flashes cell⁻¹ (n = 113) (Fig. 1). Even though all single flash variables exhibited statistical differences between prey conditions (ANOVA, p < 0.05), there were few significant differences based on multiple comparisons (Table 3). Overall the greatest difference was in maximum flux (intensity), with few differences in flash kinetics, indicating that the activation of light emission was not affected by nutritional status.

A Gonyaulax polyedra diet yielded maximum bioluminescence, with a mean TMSL of 6.2 x 10⁸ quanta cell⁻¹ based on an average of 9.7 flashes cell⁻¹, and a maximum flux (i.e. intensity of the brightest flash) of 9.6 x 10⁶ quanta s⁻¹ cell⁻¹.

For a Gymnodinium sanguineum diet at a prey concentration of 700 cells ml⁻¹, considered optimum for Protoperidinium cf. divergens population growth (Jeong & Latz 1994), the TMSL, flash flux, and number of flashes cell⁻¹ of P. cf. divergens were not significantly different from those for a Gonyaulax polyedra diet (ANOVA, p > 0.05). However, at a concentration of 2000 G. sanguineum ml⁻¹, where negative growth of P. cf. divergens occurs (Jeong & Latz 1994), its bioluminescence was significantly reduced (ANOVA, p < 0.05).

When offered dinoflagellates not known to be grazed by Protoperidinium cf. divergens, including Heterocapsa triquetra and Prorocentrum cf. balticum (Jeong & Latz 1994), P. cf. divergens exhibited bioluminescence not significantly different from unfed controls (Fig. 1; Table 2; ANOVA, p > 0.05).

Protoperidinium cf. divergens fed a Scrippsiella trochoidea diet had similar TMSL, maximum flux, and total flashes to those of cells maintained on Gonyaulax polyedra [Fisher’s PLSD, p > 0.1], even though a S. trochoidea diet is not sufficient to support population growth (Jeong & Latz 1994). Therefore, H02 can be rejected.

Test of H03 (effect of prey concentration)

With increasing Gonyaulax polyedra concentration, TMSL of Protoperidinium cf. divergens increased up to...
Table 3. Effects of different autotrophic and mixotrophic dinoflagellate diets on mechanically stimulated flashes of cultured cells of Protoperidiniun cf. divergens during Expt 1. Only first and second flashes from each mechanically stimulated cell were included in the analyses. Values represent means ± 1 SE. n: number of flashes analyzed. Other experimental details are given in Table 1 and text. All flash variables exhibited significant differences between values (ANOVA, p < 0.05). Note that Gymnodinium sanguineum was given at 2 different concentrations, 700 cells ml⁻¹ and 2000 cells ml⁻¹.

<table>
<thead>
<tr>
<th>Diet</th>
<th>Maximum flux (quanta s⁻¹)</th>
<th>Rise time (ms)</th>
<th>Decay rate (% s⁻¹)</th>
<th>Total decay time (ms)</th>
<th>Total duration (ms)</th>
<th>Quantum emission (quanta flash⁻¹)</th>
<th>%TMSL flash⁻¹</th>
<th>n</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gonjaulax polyedra</td>
<td>8.8 ± 0.3 x 10⁸</td>
<td>46 ± 5</td>
<td>2.3 ± 0.1</td>
<td>149 ± 20</td>
<td>198 ± 22</td>
<td>6.7 ± 2.4 x 10⁷</td>
<td>16 ± 4</td>
<td>40</td>
</tr>
<tr>
<td>Scripsiella trochoidea</td>
<td>8.9 ± 0.3 x 10⁸</td>
<td>39 ± 4</td>
<td>2.4 ± 0.2</td>
<td>197 ± 24</td>
<td>238 ± 28</td>
<td>9.6 ± 2.4 x 10⁷</td>
<td>23 ± 4</td>
<td>37</td>
</tr>
<tr>
<td>Gymnodinium sanguineum (700)</td>
<td>7.9 ± 0.4 x 10⁸</td>
<td>34 ± 2</td>
<td>3.5 ± 0.2</td>
<td>124 ± 12</td>
<td>156 ± 9</td>
<td>4.0 ± 0.3 x 10⁷</td>
<td>11 ± 2</td>
<td>34</td>
</tr>
<tr>
<td>Gymnodinium sanguineum (2000)</td>
<td>5.8 ± 0.3 x 10⁸</td>
<td>32 ± 3</td>
<td>3.0 ± 0.2</td>
<td>124 ± 12</td>
<td>157 ± 13</td>
<td>2.9 ± 0.3 x 10⁷</td>
<td>46 ± 16</td>
<td>30</td>
</tr>
<tr>
<td>Heterocapsa triquetra</td>
<td>4.3 ± 0.5 x 10⁸</td>
<td>35 ± 5</td>
<td>2.4 ± 0.2</td>
<td>165 ± 30</td>
<td>198 ± 33</td>
<td>4.2 ± 1.2 x 10⁷</td>
<td>27 ± 3</td>
<td>21</td>
</tr>
<tr>
<td>Prorocentrum cf. balticum</td>
<td>2.2 ± 0.5 x 10⁸</td>
<td>26 ± 2</td>
<td>4.3 ± 0.6</td>
<td>114 ± 13</td>
<td>143 ± 16</td>
<td>1.2 ± 0.4 x 10⁷</td>
<td>26 ± 12</td>
<td>20</td>
</tr>
<tr>
<td>No prey added</td>
<td>4.4 ± 0.7 x 10⁸</td>
<td>28 ± 2</td>
<td>2.8 ± 0.3</td>
<td>111 ± 11</td>
<td>139 ± 12</td>
<td>2.5 ± 0.7 x 10⁷</td>
<td>21 ± 4</td>
<td>16</td>
</tr>
</tbody>
</table>

*Maximum flux for the S. trochoidea, G. polyedra, and G. sanguineum (700) diets was significantly different from that of cells maintained without added prey (Scheffe test, p < 0.05).

**Rise time for the G. polyedra diet was significantly different from that for the P. cf. balticum diet (Scheffe test, p < 0.05).

***Instantaneous decay rate for the P. cf. balticum diet was significantly different from that of all other diets except for the G. sanguineum diet (700) (Scheffe test, p < 0.05).

****Emission for the G. polyedra diet was significantly different from those of the P. cf. balticum and G. sanguineum (2000) diets (Scheffe test, p < 0.05).
quanta cell$^{-1}$ on Day 0 to $8.5 \times 10^7$ quanta cell$^{-1}$ after 3 d, a value similar to that of single cells without added prey; however, TMSL of the cells maintained in groups increased to $4.6 \times 10^8$ quanta cell$^{-1}$ after 16 d. The Day 16 TMSL was not significantly different from that measured on Day 0 for cultures well fed on a diet of *Gonyaulax polyedra* (initial condition) (ANOVA, $p > 0.05$; Fig. 3). Therefore, $H_0$ cannot be rejected.

The maximum flux from each cell also decreased after 3 d without added prey for both group maintained and single isolated cells from an initial value of $9.2 \times 10^8$ quanta s$^{-1}$ to $2.3 \times 10^8$ quanta s$^{-1}$. However, for group maintained cells the maximum quantum flux cell$^{-1}$ subsequently increased to a maximum of $7.2 \times 10^8$ quanta s$^{-1}$ after 16 d. There was no significant temporal difference in the total number of flashes (ANOVA, $p > 0.05$), which averaged 10 flashes cell$^{-1}$.

Cannibalism, observed in *Protoperidinium* cf. *divergens* populations maintained without added prey, was based on the presence of a *P.* cf. *divergens* cell within the pallium of a feeding cell. Cannibalism is believed to be responsible for the maintenance of high levels of bioluminescence.

For single isolated cells, in addition to the decrease in TMSL with starvation, the proportion of luminescing cells also decreased. After 1 d of starvation, 16 of 16 cells were luminescent, and after 2 d of starvation, 14 of 20 cells (70%) were luminescent (method 2). For method 1, with 15 cells originally incubated for each test, after 3 d of starvation, 8 of 10 cells were luminescent (80% response, 67% survival), while after 5 d of starvation only 1 of 6 cells produced flashes (17% response, 40% survival) (method 1). Therefore starvation decreased bioluminescence and increased mortality of individually maintained cells.

**Effect of long-term laboratory culturing**

There was minimal effect of long-term culturing of *Protoperidinium* cf. *divergens* on a unicellular diet of *Gonyaulax polyedra*. Over 4.5 mo (from 4 Jan to 21 May 1993) there was no significant change in TMSL of $5.2 \times 10^9$ quanta cell$^{-1}$ and total flash number of 10 flashes cell$^{-1}$ (ANOVA, $p > 0.05$), and only a slight decrease in maximum flux per cell from $9.7 \times 10^8$ to $9.2 \times 10^8$ quanta s$^{-1}$ (ANOVA, $p < 0.05$).

Because of the sieving process used as part of the culturing methods, cultured *Protoperidinium* cf. *divergens* cells were smaller in size than freshly collected (unsieved) cells (ANOVA, $p < 0.01$); cultured and collected cells had volumes of $1.1 \pm 0.09 \times 10^5$ μm$^3$ and $1.7 \pm 0.1 \times 10^5$ μm$^3$, respectively. When adjusted for cell size, the bioluminescence capacity per unit surface area of cultured cells was similar to that of freshly collected cells (1-way ANOVA, $p > 0.05$), and there was no significant difference in either maximum quantum (flux cell$^{-1}$) or total number of flashes cell$^{-1}$ (Table 4) (Fisher's PLSD, $p > 0.05$).

The bioluminescence of *Protoperidinium* cf. *divergens* incubated on *Gymnodinium sanguineum* for 4 d was compared to that of cells maintained on a *G.* san-

Table 4. Total stimulated bioluminescence of *Protoperidinium cf. divergens* and *P. crassipes*. Cultured cells were maintained on a *Gonyaulax polyedra* diet. Total mechanically stimulable luminescence (TMSL) was elicited by maintained stirring until depletion of light emission. Surface area (SA) calculated from mean equivalent cell diameter (see ‘Materials and methods: analysis’). Values represent means ± 1 SE; n: number of cells measured.

<table>
<thead>
<tr>
<th></th>
<th>TMSL (quanta cell(^{-1}))</th>
<th>Cell length (µm)</th>
<th>Cell width (µm)</th>
<th>TMSL SA(^{-1}) (quanta cell(^{-1}) mm(^{-2}))</th>
<th>Total no. of flashes cell(^{-1})</th>
<th>n</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>P. cf. divergens</em></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cultured</td>
<td>6.2 ± 0.9 \times 10^6</td>
<td>75 ± 1</td>
<td>56 ± 2</td>
<td>9.6 ± 1.0 \times 10^{10}</td>
<td>9.7 ± 1.6</td>
<td>19</td>
</tr>
<tr>
<td>Freshly collected</td>
<td>13.9 ± 2.7 \times 10^6</td>
<td>81 ± 2</td>
<td>68 ± 2</td>
<td>10.5 ± 1.8 \times 10^{10}</td>
<td>12.5 ± 2.5</td>
<td>11</td>
</tr>
<tr>
<td><em>P. crassipes</em></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cultured</td>
<td>7.9 ± 1.6 \times 10^6</td>
<td>79 ± 1</td>
<td>78 ± 0.4</td>
<td>6.0 ± 1.3 \times 10^{10}</td>
<td>5.8 ± 0.8</td>
<td>12</td>
</tr>
</tbody>
</table>

Table 5. Paired comparison between first and second mechanically stimulated flashes of *Protoperidinium cf. divergens* and *P. crassipes* cells cultured on a *Gonyaulax polyedra* diet. Flashes were elicited by maintained stirring. Values represent means ± 1 SE; n: number of cells tested. *Significant difference within species between values for first and second flashes (paired t-test, \(p < 0.05\)).

<table>
<thead>
<tr>
<th></th>
<th>Maximum flux (quanta s(^{-1}))</th>
<th>Rise time (ms)</th>
<th>Decay rate (% s(^{-1}))</th>
<th>Total decay time (ms)</th>
<th>Total duration (ms)</th>
<th>Quantum emission (quanta flash(^{-1}))</th>
<th>%TMSL flash(^{-1})</th>
<th>n</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Protoperidinium cf. divergens</em></td>
<td></td>
<td></td>
<td></td>
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<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>First</td>
<td>8.9 ± 0.5 \times 10^8</td>
<td>45 ± 4</td>
<td>2.6 ± 0.1</td>
<td>104 ± 5</td>
<td>151 ± 7</td>
<td>5.1 ± 0.3 \times 10^7</td>
<td>15.8 ± 2.4</td>
<td>17</td>
</tr>
<tr>
<td>Second</td>
<td>8.4 ± 0.7 \times 10^8</td>
<td>35 ± 4</td>
<td>2.3 ± 0.3</td>
<td>144 ± 24</td>
<td>185 ± 28</td>
<td>5.9 ± 1.0 \times 10^7</td>
<td>16.6 ± 3.5</td>
<td>17</td>
</tr>
<tr>
<td><em>Protoperidinium crassipes</em></td>
<td></td>
<td></td>
<td></td>
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<td></td>
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<td></td>
<td></td>
</tr>
<tr>
<td>First</td>
<td>4.7 ± 0.2 \times 10^8</td>
<td>24 ± 1</td>
<td>2.9 ± 0.2</td>
<td>71 ± 4</td>
<td>96 ± 5</td>
<td>1.9 ± 1.4 \times 10^8</td>
<td>.34 ± 5</td>
<td>16</td>
</tr>
<tr>
<td>Second</td>
<td>3.8 ± 0.3 \times 10^8</td>
<td>24 ± 1</td>
<td>3.1 ± 0.2</td>
<td>76 ± 9</td>
<td>100 ± 9</td>
<td>1.4 ± 1.7 \times 10^8</td>
<td>.24 ± 3</td>
<td>16</td>
</tr>
</tbody>
</table>

guineum diet for 1.8 mo. There was no significant difference in TMSL, total number of flashes cell\(^{-1}\), or cell size (ANOVA, \(p > 0.05\)), and only a barely significant change in maximum flux (ANOVA, \(p = 0.05\)). Therefore diet during the preincubation period did not affect bioluminescence measured after a 4 d incubation, and there was no apparent degradation of the physiological state of *P. cf. divergens* cultures maintained on a *G. sanguineum* diet for more than 50 d. The lack of change in bioluminescence with long-term maintenance on a non-luminescent diet suggests that the luminescent chemistry is synthesized *de novo* and is not obtained through the diet.

Size-dependent interspecific differences in bioluminescence

Even though the TMSL of the larger species *Protoperidinium crassipes* was greater than that of *P. cf. divergens*, when expressed per unit surface area, total bioluminescence per cell was less than that of *P. cf. divergens* (Table 4). Individual flashes were brighter than those of *P. cf. divergens* but had shorter rise and decay times (Table 5). Neither species displayed dramatic differences between first and second flashes. *P. crassipes* flashes averaged 100 ms in duration with a rise time of 24 ms while those of *P. cf. divergens* were approximately 150 to 190 ms in duration with rise times of 35 to 45 ms.

Swimming speed during maintenance without added prey

The speed of actively swimming *Protoperidinium cf. divergens* cells (53 to 64 µm diameter) maintained in group conditions without added prey was measured every 2 to 3 d. Even though the swimming speed of these cells significantly decreased from an initial value of 0.98 ± 0.02 mm s\(^{-1}\) to 0.83 ± 0.02 mm s\(^{-1}\) after 12 d (Fisher’s PLSD, \(p < 0.05\)), this represented a decrease of only 1% d\(^{-1}\). Cells presumably maintained nutrition and swimming ability through cannibalism, because a greater decrease in swimming speed would be expected if no food supply were available. In fact, cells maintained individually without added prey were dead in 3 to 5 d.
DISCUSSION

The hypotheses tested in the present study, except for $H_04$, were rejected. Therefore prey species does affect the bioluminescence of Protoperidinium ($H_01$), bioluminescence is not necessarily related to population growth rate ($H_02$), prey concentration does affect bioluminescence ($H_03$), and cannibalism increases bioluminescence to high levels even when no added prey are present ($H_04$).

Effect of red tide dinoflagellate diet

The results of the present study show that the bioluminescence of Protoperidinium cf. divergens is significantly affected by the species and cell abundance of red tide dinoflagellate diets. For all diets the TMSL of $P$. cf. divergens fed on dinoflagellate prey was not significantly correlated with population growth, but instead with feeding frequency, the percent ratio of $P$. cf. divergens cells with a pallium (containing a prey cell) to total cells at any one time (Fig. 4). Bioluminescence may be an indicator of in situ feeding frequency even when no population growth occurs.

Previously, high levels of bioluminescence in Protoperidinium huberi feeding on mixed and unialgal diatom diets have been shown to be associated with high population growth rates (Buskey et al. 1994). For $P$. cf. divergens, only its optimal dinoflagellate diet of Gonyaulax polyedra supported the relationship between high bioluminescence and population growth rate (Fig. 5).

Based on the results of the present study, the following relationships between bioluminescence and in situ abundance of Protoperidinium cf. divergens are expected: during Gonyaulax polyedra red tides, high population growth rates and high bioluminescence of $P$. cf. divergens would occur. During Prorocentrum cf.

![Graph A](image1.png)

![Graph B](image2.png)

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Fig. 4. Relationship between feeding, growth, and bioluminescence of Protoperidinium cf. divergens for various autotrophic and mixotrophic dinoflagellate diets (see Table 1, Expt 1). (A) TMSL as a function of instantaneous feeding frequency. (B) TMSL as a function of population growth rate. Feeding and population growth data from Jeong & Latz (1994).

Fig. 5. Bioluminescence of Protoperidinium cf. divergens as a function of population growth rate for a Gonyaulax polyedra diet. (A) TMSL. The equation of the linear regression was TMSL = $3.58 \times 10^8 + (9.95 \times 10^8) \times$ (growth rate), $R^2 = 0.83$. (B) Maximum flux per cell. Calculated from the data displayed in Fig. 2 using the growth rate data of Jeong & Latz (1994). Values represent means ± 1 SE.
*balticum* red tides, low bioluminescence and abundance of *P. cf. divergens* are predicted because no feeding on this prey occurs. During *Gymnodinium sanguineum* red tides, low bioluminescence and abundance are predicted at high prey concentrations ≥2000 cells ml⁻¹, while at low prey concentrations ≤700 cells ml⁻¹ high bioluminescence and abundance of *Protoperidinium cf. divergens* would occur. During *Scripsiella trochoidea* red tides, high bioluminescence is predicted due to high feeding rates, even though the abundance of *P. cf. divergens* would be low.

**Effect of cannibalism**

There was a dramatic difference between the bioluminescence of *Protoperidinium cf. divergens* maintained individually without added prey, in which cells became nonluminescent prior to death and/or after 3 d, and group maintained cells which resisted starvation through cannibalism. Cannibalism, which appeared to be most important after 3 d without added prey, resulted in increased survival time and gradually increasing levels of bioluminescence, which eventually reached levels similar to those of cells maintained under optimal prey conditions. Therefore, if feeding occurs, whether due to cannibalism or ingestion of red tide dinoflagellate prey, *P. cf. divergens* cells can produce bright bioluminescence, even when population growth does not occur. Cannibalism might be an important strategy for maintaining high bioluminescence after red tides when *P. cf. divergens* abundance is high and red tide dinoflagellate prey abundance is low.

**Bioluminescence dynamics**

TMSL per unit size of freshly collected *Protoperidinium cf. divergens* cells, as well as that of cultured cells maintained on an optimum *Gonyaulax polyedra* diet, followed the correlation of Seliger et al. (unpubl., cited in Buskey et al. 1992). This correlation states that, for autotrophic dinoflagellates, TMSL scales to approximately 10¹⁷ quanta cell⁻¹ mm⁻² surface area. Therefore cultured cells of *P. cf. divergens* were healthy and expressed maximum bioluminescence capacity.

There were few significant differences between the emission properties of first and second flashes of *Protoperidinium cf. divergens* and *P. crassipes* cells maintained on a *Gonyaulax polyedra* diet. This indicates that there is a single mechanism for synchronization and light production in the luminescent microsources within the cell (Widder & Case 1982). Flash kinetics were similar to those measured for *Protoperidinium spp.* in the north Atlantic (60°N, 20°W) during May 1991, when rise times averaged 29 ms, flash duration was 141 s, maximum flux was 1.1 x 10¹⁰ quanta s⁻¹, and each cell produced approximately 6 flashes with a TMSL of 2 x 10⁹ quanta cell⁻¹ (Latz unpubl. data). The rapid kinetics of *Protoperidinium spp.* flashes are similar to those of other dinoflagellates such as *G. polyedra* and *Noctiluca scintillans* (Eckert 1965, Latz unpubl. data), but are faster than those of *Pyrocystis* species (Widder & Case 1981, Jess 1985).

Changes in bioluminescence capacity were reflected in the number of quanta resulting from each flash, not the total number of flashes produced by a cell. This suggests that the bioluminescence excitation process, which involves an action potential propagated along the vacuole membrane (Eckert 1965, Widder & Case 1981) leading to proton flux across the membrane of the vesicles containing the luminescent chemistry (Hastings & Dunlap 1986), is independent of quantum emission. A reduction in flash quantum flux, presumably due to reduced amount of energy available for the luminescent system, may reduce the effectiveness of bioluminescence as an antipredation behavior (Esaiae & Curr 1972, White 1979).

**Energy utilization**

Based on the results of the present study, the energy requirements for swimming, bioluminescence, and growth can be considered. Two lines of evidence are important: (1) differences in the population growth rate of *Protoperidinium* feeding on different dinoflagellate prey did not necessarily signify differences in bioluminescence (*H₂O₂*), and (2) there was no significant difference in swimming speed even when there was a difference in bioluminescence. These data suggest that energy requirements for bioluminescence are less than those for reproduction. When nutritional status is high, sufficient energy is available for all metabolic needs, including bioluminescence and growth. For medium levels, insufficient energy is available for growth, although bioluminescence may still be high. When the nutritional status is low, available energy is inadequate to support bioluminescence and growth. Therefore energy utilization by *Protoperidinium cf. divergens* may be prioritized in the following order: swimming (for grazing) > bioluminescence (to reduce predation) > reproduction (for population increase).

Bioluminescence may be a sensitive indicator of the nutritional status and feeding history of natural populations of *Protoperidinium*, especially when suboptimal prey conditions result in low population growth rates.
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LITERATURE CITED


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