Effects of water-borne gametes on the aggregation behavior of *Lytechinus variegatus*

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ABSTRACT: Water-borne gametes are known to cue mass spawning in externally fertilizing marine invertebrates. In motile animals, such as sea urchins, they might also cue behavior to enhance fertilization success by either (1) increasing movement and thus the number of intraspecific encounters for numerous small-scale events or (2) causing the formation of small-sized aggregations for single spawning events. This study investigated the ability of water-borne gametes to induce spawning and aggregation behavior in the echinoid *Lytechinus variegatus*. In the field, 1 individual from each of 40 pairs of urchins was induced to spawn to test whether spawning would be induced in the other urchin. Additionally, 5 urchins were induced to spawn upstream from 20 individuals placed in a group. Thirteen replicates of this experiment were conducted to test whether the larger volume of gametes combined with more urchins would induce spawning. Ripe and nonripe urchins were also videotaped in a 2 m diameter pool with and without sperm in the water (n = 3). All urchins were found to move rapidly (50 cm min⁻¹), although the amount of movement, distance apart, and number and duration of encounters was the same regardless of their reproductive status or the presence/absence of sperm in the water. Urchins never spawned in response to water-borne gametes although most were found to be ripe.

KEY WORDS: *Lytechinus variegatus* · Movement · Behavior · Spawning · Aggregations · Echinoid

INTRODUCTION

In free-spawning marine invertebrates, synchronization of gamete release is necessary to enhance fertilization success. If gametes are shed asynchronously, they could lose viability and/or become too dilute for successful fertilization (Pennington 1985, Levitan et al. 1992, Sewell & Levitan 1992, Levitan 1995). Phytoplankton (Smith & Strickland 1983, Starr et al. 1990, 1992), water temperature (Minchin 1992, Selvakumaraswamy & Byrne 2000), day length (Minchin 1992, Selvakumaraswamy & Byrne 2000) and water-borne gametes have been shown to cue spawning (Beach et al. 1975, Miller 1989, Unger & Lott 1994, Hardege & Bentley 1997). These stimuli presumably result in mass spawning which should enhance fertilization success if conspecifics are in close proximity (Levitan 1995).

Quantitative observations of mass spawning events by motile marine invertebrates are relatively rare and generally observed with temperate species. Several temperate species of echinoids (Minchin 1992, D. R. Levitan pers. comm.), crinoids (Meyer et al. 1984), sea-stars [Minchin 1987, Run et al. 1988] and holothurians (McEuen 1988, Sewell & Levitan 1992) have been observed to mass spawn. In contrast, observations of spawning in tropical species have mostly been with animals living on the Great Barrier Reef (Babcock & Mundy 1992, Babcock et al. 1992, Olsen et al. 1993). In most of these studies, only a few individuals of a species were observed to spawn at any one time (Pearse et al. 1988, Minchin 1992, Babcock et al. 1992, Olsen et al. 1993). Research on seasonal aggregations of some echinoderms also suggests that discrete spawning events can occur among individuals of subgroups of a population. Several species of temperate or deep-sea
Echinoderms have been observed to form aggregations during the reproductive season (Tyler et al. 1992, Young et al. 1992, Unger & Lott 1994). Members of the aggregations were always more ripe than solitary individuals in the same population. Natural spawning was not observed in any of these studies, so it was unknown whether individuals in the aggregations spawned once or several times before they separated from their partner.

Some tropical species may spawn numerous times during intraspecific encounters that occur in their reproductive season. Populations of the echinoid *Lyttechinus variegatus* reproduce from February through June and again from October through November off SE Florida (McCarthy & Young 2002). Reproductive periods are characterized by females that are all ripe yet have different egg size distributions. This observation suggests that asynchronous spawning occurs within the population (McCarthy & Young 2002). High (1 to 3 m d⁻¹) urchin movement is likely to result in numerous intraspecific encounters during a single reproductive season. Such encounters may facilitate small-scale spawning events involving numerous individuals of the population. It is unknown whether cues that result in mass spawning by some species affect gamete release or reproductive behavior of species such as *L. variegatus* that spawn numerous times during a reproductive season.

This study investigated the role of water-borne gametes as cues for spawning and reproductive behavior in the echinoid *Lyttechinus variegatus*. Field experiments tested (1) whether urchins in naturally occurring aggregations were more ripe than solitary urchins and (2) whether spawning in response to water-borne gametes is affected by aggregation size. Laboratory experiments tested whether ripe and non-ripe urchins respond to sperm by (1) spawning, (2) moving faster to increase intraspecific encounters or (3) aggregating.

**MATERIALS AND METHODS**

**Study areas.** Field experiments were conducted in May 1994 with *Lyttechinus variegatus* (55 to 65 mm in diameter) found off the SE shore of Virginia Key in Biscayne Bay, Florida (Fig. 1). Urchins collected from this area were also used for laboratory experiments during March and July 1994. Additional replicates of the field experiments were run in May 2002 with similar-sized *L. variegatus* found in the Intracoastal Waterway near Jupiter Inlet, Florida (Fig. 1). The bottom at both locations was level, ca. 1.5 m deep and covered homogeneously by a mixture of the seagrasses *Thalassia testudinum, Syringodium filiforme* and *Halodule wrightii*.

**Aggregation as a function of reproductive state.** To test whether ripe urchins were more likely to be aggregated or solitary, the proportion of reproductive individuals in naturally occurring pairs was compared with that of randomly paired solitary urchins. Twenty pairs and 40 solitary urchins were sampled in 1994 in Virginia Key and again in 2002 at Jupiter Inlet. Aggregated urchins consisted of at least 2 individuals with their spines touching. Solitary urchins were at least 2 m from any conspecific. Sex and gonad maturity were determined by either histological examination of gonads or injection of 0.55 M KCl to induce spawning. For each location, a G-test was used to compare the likelihood that randomly paired and observed pairs will have 0, 1 or 2 reproductive individuals (Sokal & Rohlf 1995). A Kruskal-Wallis test was used to determine where the number of urchins in each category varied between 1994 and 2002 (Sokal & Rohlf 1995)

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![Fig. 1. Study areas in Florida. Bottom panel shows close up of Key Biscayne area. Right panel shows close up of Jupiter area. #: specific study sites within each area](image)

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of gametes would induce spawning in the other urchin of the pair (see Fig. 2). Ten urchin pairs were tested at Virginia Key in 1994 with an additional 30 pairs tested at Jupiter Inlet in 2002. In a second set of experiments, 5 urchins were placed 0.25 m upstream from a group of 20 individuals. The upstream group was injected with 0.55 M KCl to test whether gametes drifting down over the downstream group would induce any of these individuals to spawn. In these experiments, a 15 min acclimation time period was allowed prior to each run to minimize any behavioral artifacts from handling. Normally, urchins would quickly stop moving if they were in an area with some seagrass cover. In both experiments, urchins were observed for 30 min after the induction of spawning. Three runs were done at Virginia Key in 1994 and 10 at Jupiter Inlet in 2002.

**Movement and aggregation responses to sperm.** This experiment tested whether ripe and nonripe urchins responded to the presence of sperm by (1) spawning, (2) moving fast to increase intraspecific encounter rates or (3) remaining aggregated when encounters occurred. Three replicate samples of 9 urchins were videotaped indoors in a 2 m diameter fiberglass tank (20 cm depth) with no outside circulation for 6 h periods using a video camera wired to a time-lapse VCR (see Fig. 3). Water temperature, dissolved oxygen and salinity were monitored during all runs. The first 6 h run started in the early morning. Urchins were initially placed approximately 0.25 m apart. At the end of the first run of each day, the pool was drained, cleaned and the urchins were rinsed with clean seawater. They were then placed back into the newly filled pool for the second run. Sperm from 8 additional male urchins obtained by KCl injection were combined with seawater in a 1000 ml container to create a highly concentrated solution of fresh sperm. The solution was added to the pool at the beginning of the run and a fresh sperm suspension was added 3 h later. During successive days of the experiment, the order of the sperm treatment and the non-sperm treatment was alternated to block for potential artifactual differences between the morning and afternoon. Three replicates of these experiments were completed during March (reproductive season) and July (nonreproductive period) to investigate potential seasonal behavioral differences.

The videotape data were analyzed on a video monitor by tracing the path of each urchin and recording (1) the distance traveled for each hour of the experiment, (2) the mean distance from each urchin to its nearest neighbor and (3) the number of encounters and duration of each encounter during the course of the experiment. Each of these data sets was analyzed with a 2-factor (reproductive state, sperm in the water) repeated measures (each successive hour) ANOVA (SYSTAT 1992).

**RESULTS**

**Aggregation as a function of reproductive state**

A G-test revealed no significant difference in the proportion of reproductive individuals between naturally occurring pairs and single individuals paired at random (Fig. 2). In 1994, there were 8 male/male, 5 male/female, 3 female/female, 3 male nonripe and 1 nonripe/nonripe combinations (Fig. 2a). There were 20 male, 15 female and 5 nonripe solitary urchins. In 2002, there were 4 male/male, 14 male/female and 2 female/female combinations (Fig. 2b). In this year, there were 23 male, 15 female and 2 nonripe solitary urchins. The number of urchin pairs in each of the 3 categories (0, 1 or 2 ripe individuals) did not vary significantly between 1994 and 2002 (Kruskal-Wallis statistic = 0.5162, P < 0.915).

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**Fig. 2. Lytechinus variegateus.** Number of observed versus randomly paired sea urchins having 0, 1 or 2 reproductive individuals in (a) 1994, (b) 2002 and (c) the 2 yr combined. G_{obs} = observed G value. ns = not significant.
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**Spawning responses in natural and artificial aggregations**

No individual in any of the 40 naturally occurring pairs spawned when its partner was induced to spawn with 0.55 M KCl (Fig. 3). In 1994, there were 5 all male and 1 all female pairs of urchins (Fig. 3a). There was 1 pair of nonripe urchins and 3 pairs with nonripe partners. In the latter case, all ripe individuals were females. In 2002, all individuals in the 30 encountered pairs were ripe (Fig. 3b). In 60% of the pairs in the experiment, KCl injection resulted in a male spawning next to a non-spawning female. A male spawning next to a non-spawning male was encountered 20% of the time, while a female to a male or a female accounted for the remaining 20% of the experimental pairs.

In the 13 artificial aggregates, only 1 female spawned in response to injected individuals that were spawning. Within each group (a total of 260) 90% of all individuals were ripe with 10% being nonreproductive. The sex ratios for the upstream and downstream groups appeared similar for 1994 and 2002 (Fig. 4).

![Fig. 3. Lytechinus variegatus. Sex ratio of observed pairs of sea urchins used in field spawning experiments in (a) 1994, (b) 2002 and (c) the 2 yr combined. No individual in any of the pairs spawned when its partner was induced to spawn.](image)

M = male; F = female; N = nonripe

**Movement and aggregation responses to water-borne gametes**

No spawning was observed by any of the urchins during the laboratory experiments. Histological sections of gonads revealed that, during the nonreproductive season, 100% of the urchins had gonads that were in early gametogenesis. During the reproductive season, 92% of the urchins were ripe, with 8% being in earlier gametogenic stages. Urchin behavior was not observed to be different between ripe and nonripe individuals during the reproductive season.

The mean distance traveled during each hour of the laboratory experiment averaged between 25 and 175 cm (Fig. 5). No significant difference was found between any treatment means although the mean locomotory rate was lowest for fully ripe urchins with sperm in the water than any of the other treatments (Table 1, Fig. 5). A significant time effect (log-transformed data) was found ($F = 8.79; p < 0.001$) in which the mean locomotory rate was fastest in the first hour of the 6 h experiment (Table 1).

The mean distance between urchins varied from 26 to 42 cm (Fig. 6). No significant differences were found among treatment means. However, generally the mean
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among treatment means (Table 3a). The longest duration of encounters occurred in the treatment with ripe urchins and sperm present although it was not found to be significant (Table 3b).

Environmental factors were at fairly consistent levels for experimental runs within each season (reproductive or nonreproductive) of 1994 (Table 4). Most measurements were also fairly similar between runs of both seasons (Table 4). Water temperature was the exception, with values of 24.8°C and 29.0°C for March and July, respectively.

We ran several Pearson correlations to determine whether changes in environmental factors during the course of the day may have affected the experimental results. No significant correlations could be found between any of the measured environmental factors and dependent variables.

**Spawning responses in natural and artificial aggregations**

Gametes spawned over natural and artificial aggregations failed to cue spawning from most members at both locations. At Virginia Key, some non-ripe urchins were encountered in collected pairs whereas all pairs were ripe at Jupiter Inlet. The sex ratio and number of ripe individuals of upstream and downstream groups of urchins was similar at both sites. No differences in response could be related to the size of the aggregation. These results could have occurred because (1) of chemical interference by KCl, (2) natural re-
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**DISCUSSION**

**Aggregation as a function of reproductive state**

Ripe *Lytechinus variegatus* were found alone and also in small aggregations at both Virginia Key and Jupiter Inlet. This suggests that ripe individuals (1) might not be able to find each other easily, (2) randomly find each other and stay together either seasonally or for a short time, (3) are waiting to respond to a localized cue to temporarily aggregate or (4) do not aggregate for reproduction. It appears that *L. variegatus* does not seasonally aggregate as previously investigated (McCarthy & Young 2002). Therefore, if small-scale reproductive aggregations occur, they must be short term and result from either random encounters or a very localized cue.

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<td>Reproductive state (Rs)</td>
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Fig. 6. *Lytechinus variegatus*. Hourly mean (+1.0 SD) distance between urchins with (a) immature and (b) mature gonads in treatments with and without sperm. No significant differences were found among the treatment means.

Fig. 7. *Lytechinus variegatus*. Mean duration of encounters (+1.0 SD) in the laboratory experiment with ripe and nonripe urchins, in the presence and absence of sperm. No significant differences were found among the treatment means.

Responses were obscured by urchin reactions to handling or (3) urchins were not given enough time to respond. However, others have obtained positive results using similar techniques.

Unger & Lott (1994) were able to use water-borne gametes to induce spawning by *Sphaerechinus granularis* found in natural aggregations. Urchins were given a short acclimation, yet still spawned in response to water-borne gametes within 20 min of the start of the experiment.

We believe that if water-borne gametes induce spawning or behavior in *Lytechinus variegatus*, they do so in combination with 1 or several exogenous factors that did not exist in our experiments. For instance, water-borne gametes may only be important when present with phytoplankton, moon phase, tide or darkness. It has been suggested that *L. variegatus* spawning may be linked to the lunar phase (Moore et al. 1963, Moore & Lopez 1972, Lessios 1991, Watts et al. 2001) and that they are more active during the night (P. Kier pers. comm.). If we had conducted our experiments during these times, perhaps we may have observed behavioral responses from the urchins cued by water-borne gametes. However, if gametes do not work in synergy with some other factor such as moon phase or night, they may not be important at all in inducing spawning in *L. variegatus*. Gamete release may occur directly in response to any of the above-mentioned environmental factors. It may also result from a pheromone or tactile stimulus. Chemical communication has been implicated in initiating and maintaining synchronization of gametogenesis in the holothurid *Cucumaria frondosa* (Hamel & Mercier 1996).
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<tr>
<td>Time × Sperm</td>
<td>78.0</td>
<td>5</td>
<td>15.7</td>
<td>0.42</td>
<td>0.829</td>
</tr>
<tr>
<td>Time × Rs × Sperm</td>
<td>182.0</td>
<td>5</td>
<td>36.4</td>
<td>0.98</td>
<td>0.441</td>
</tr>
<tr>
<td>Error</td>
<td>1480.0</td>
<td>40</td>
<td>37.0</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Fig. 6. *Lytechinus variegatus*. Hourly mean (±1.0 SD) distance between urchins with (a) immature and (b) mature gonads in treatments with and without sperm. No significant differences were found among the treatment means.

Fig. 7. *Lytechinus variegatus*. Mean duration of encounters (+1.0 SD) in the laboratory experiment with ripe and non-ripe urchins, in the presence and absence of sperm. No significant differences were found among the treatment means.

**Figures:**

- Fig. 6: Lytechinus variegatus. Hourly mean (±1.0 SD) distance between urchins with (a) immature and (b) mature gonads in treatments with and without sperm. No significant differences were found among the treatment means.

- Fig. 7: Lytechinus variegatus. Mean duration of encounters (+1.0 SD) in the laboratory experiment with ripe and non-ripe urchins, in the presence and absence of sperm. No significant differences were found among the treatment means.

Responses were obscured by urchin reactions to handling or (3) urchins were not given enough time to respond. However, others have obtained positive results using similar techniques.

Unger & Lott (1994) were able to use water-borne gametes to induce spawning by *Sphaerechinus granularis* found in natural aggregations. Urchins were given a short acclimation, yet still spawned in response to water-borne gametes within 20 min of the start of the experiment.

We believe that if water-borne gametes induce spawning or behavior in *Lytechinus variegatus*, they do so in combination with 1 or several exogenous factors that did not exist in our experiments. For instance, water-borne gametes may only be important when present with phytoplankton, moon phase, tide or darkness. It has been suggested that *L. variegatus* spawning may be linked to the lunar phase (Moore et al. 1963, Moore & Lopez 1972, Lessios 1991, Watts et al. 2001) and that they are more active during the night (P. Kier pers. comm.). If we had conducted our experiments during these times, perhaps we may have observed behavioral responses from the urchins cued by water-borne gametes. However, if gametes do not work in synergy with some other factor such as moon phase or night, they may not be important at all in inducing spawning in *L. variegatus*. Gamete release may occur directly in response to any of the above-mentioned environmental factors. It may also result from a pheremone or tactile stimulus. Chemical communication has been implicated in initiating and maintaining synchronization of gametogenesis in the holothurid *Cucumaria frondosa* (Hamel & Mercier 1996).
Table 3. *Lytocinus variegatus*. Results of 2-way ANOVA comparing (a) the number of encounters and (b) duration of encounters of urchins in the laboratory experiment. The independent factors are reproductive state; Rs (2 levels) and the presence or absence of sperm. Means of encounter duration time were log-transformed in order to meet assumptions of ANOVA

<table>
<thead>
<tr>
<th>Dependent variable</th>
<th>Source</th>
<th>SS</th>
<th>df</th>
<th>MS</th>
<th>F-ratio</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>(a) Encounter number</td>
<td>Reproductive state (Rs)</td>
<td>12.0</td>
<td>1</td>
<td>12.0</td>
<td>0.364</td>
<td>0.563</td>
</tr>
<tr>
<td></td>
<td>Sperm</td>
<td>40.3</td>
<td>1</td>
<td>40.3</td>
<td>1.222</td>
<td>0.301</td>
</tr>
<tr>
<td></td>
<td>Rs x Sperm</td>
<td>56.3</td>
<td>1</td>
<td>56.3</td>
<td>1.707</td>
<td>0.228</td>
</tr>
<tr>
<td></td>
<td>Error</td>
<td>264.0</td>
<td>8</td>
<td>33.0</td>
<td></td>
<td></td>
</tr>
<tr>
<td>(b) Encounter duration</td>
<td>Reproductive state (Rs)</td>
<td>0.153</td>
<td>1</td>
<td>0.153</td>
<td>0.159</td>
<td>0.700</td>
</tr>
<tr>
<td></td>
<td>Sperm</td>
<td>0.022</td>
<td>1</td>
<td>0.022</td>
<td>0.023</td>
<td>0.882</td>
</tr>
<tr>
<td></td>
<td>Rs x Sperm</td>
<td>1.035</td>
<td>1</td>
<td>1.035</td>
<td>1.076</td>
<td>0.330</td>
</tr>
<tr>
<td></td>
<td>Error</td>
<td>7.695</td>
<td>8</td>
<td>0.962</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Movement and aggregation responses to sperm

Sperm did not induce spawning, increase locomotory rates and change mean distance between urchins or change durations of encounters. Behavioral responses did not change with the reproductive state of the urchins nor did they appear related to any of the environmental factors measured in this experiment.

Individual *Lytocinus variegatus* were found to move at heightened speeds in what seemed to be random directions immediately after being handled. They then slowed down after 1 h and maintained much slower locomotory rates. It is likely that this is an evasion response to handling. Similarly, members of aggregations of the echinoid *Diadema* spp. quickly moved in a direction away from a perceived threat when 1 individual of the aggregate was touched (Pearse & Arch 1969).

The lack of significant effects by the independent factors in these experiments could be because of any of the same reasons outlined for the field experiments. However, unlike the field experiments, the presence of gametes in the water had more time (6 h) to induce a response. If urchins respond to water-borne gametes, 6 h is probably a reasonable time to expect a response. Otherwise, a spawning individual would waste an enormous number of gametes before successful reproduction could occur. Further, most KC1-induced *Lytocinus variegatus* only spawn for approximately 10 to 20 min (D. A. McCarthy unpubl. data).

Handling before and during the experiments may have adversely affected urchin behavior. It is possible that some of the urchins spawned a little during handling prior to the experiment. While urchins were generally ripe in our experiments, their gonads may not have been at peak capacity to spawn in response to gametes. Also, prior to each experimental run, the urchins may not have had a sufficient acclimation period with enough natural settings like sand/grass to completely stimulate them to respond as they would in their natural habitat.

**CONCLUSIONS**

Water-borne gametes do not appear to affect spawning or reproductive behavior in *Lytocinus variegatus*. If they do induce such responses, it is most likely that they do so synergistically with either (1) one or several other environmental factors or (2) localized cues from urchins such as pheremones and/or touch. Further, it is likely that such cues will only induce spawning and behavioral responses in small groups within a population. Asynchronous oocyte distributions among ripe females (Beddingfield & McClintock 1998, 2000, McCarthy & Young 2002) suggest urchins may sporadically spawn small cohorts of gametes numerous times during the reproductive season. Considering such a spawning method, a large number of *L. variegatus* might not be expected to simultaneously respond to water-borne gametes like some mass spawning temperate species. Reproductive responses in *L. variegatus* may only occur during very short time periods during the reproductive season, when several urchins are reproductively ready, and the appropriate endogenous or exogenous cues are present.

Table 4. Dissolved oxygen, salinity and temperature means (±1.0 SD) for laboratory experiment with *Lytocinus variegatus* during March and July 1994

<table>
<thead>
<tr>
<th></th>
<th>March 1994</th>
<th>July 1994</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Ripe</td>
<td>Nonripe</td>
</tr>
<tr>
<td></td>
<td>Morning</td>
<td>Afternoon</td>
</tr>
<tr>
<td>Oxygen (g ml⁻¹)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>5.6 (0.45)</td>
<td>5.5 (0.41)</td>
<td>5.8 (0.56)</td>
</tr>
<tr>
<td>Salinity (ppt)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>28.2 (0.47)</td>
<td>29.0 (0.13)</td>
<td>29.2 (0.24)</td>
</tr>
<tr>
<td>Water temp (°C)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>28.3 (0.61)</td>
<td>24.5 (0.14)</td>
<td>25.0 (0.16)</td>
</tr>
</tbody>
</table>
Movement and aggregation responses to sperm

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