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Marine snow derived from abandoned larvacean houses: sinking rates, particle content and mechanisms of aggregate formation

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ABSTRACT: The dynamics and formation mechanisms of marine snow aggregates from abandoned larvacean houses were examined by laboratory experiments and field sampling during a spring diatom bloom in a shallow fjord on the west coast of the USA. Intact aggregates were sampled both from sediment traps and directly from the water column by divers. All aggregates were composed of 1 abandoned house of the larvacean *Oikopleura dioica* to which numerous diatoms, fecal pellets, ciliates, and amorphous detritus were attached. High vertical flux rates (20 000 to 120 000 houses m^{-2} d^{-1}) and settling velocities (average 120 m d^{-1}) imply a rapid turnover of suspended larvacean houses, and concentrations of diatoms and fecal pellets in the aggregates exceeding ambient concentrations by 3 to 5 orders of magnitude suggest their potential importance in driving the vertical flux of particles. Identical particle assemblages were observed in aggregates collected in the water column and in sediment traps. Most of the fecal pellets found in the houses were most likely produced by the larvaceans themselves. Numbers of diatoms per house corresponded with the diatom concentrations in the ambient water and, on average, each aggregate contained diatoms in abundances corresponding to those found in 4.5 ml of ambient water. Laboratory measurements showed that larvacean houses scavenge diatoms and fecal pellets while sinking, and observed scavenging rates were similar to those predicted from theory. However, both predicted and observed scavenging rates in experiments were orders of magnitude too low to account for the particle content observed on aggregates from the water column. Based on models, shear coagulation was also assessed to be insignificant in forming aggregates. It is concluded that most of the particles become attached to the incurrent filters of the larvacean house while it is still inhabited, and that shear and sinking insignificantly contribute to particle collisions and adhesions on the abandoned house.

KEY WORDS: Marine snow · Larvaceans · Scavenging · Mechanisms of aggregation

INTRODUCTION

Macroscopic marine aggregates, known collectively as marine snow, are ubiquitous in the sea (Alldredge & Silver 1988), and sedimentation of organic matter to the sea floor occurs primarily in the form of aggregates (Fowler & Knauer 1986). Marine snow may contain all kinds of particulate material also found suspended in the water column, and the composition of aggregates ranges from pure phytoplankton to aggregates composed of amorphous detritus (Alldredge & Silver 1988).

Members of the zooplankton community such as pteropods, larvaceans and salps produce gelatinous feeding nets and houses which, together with their fecal pellets, also occasionally comprise the aggregates (Alldredge 1972, Pomeroy & Deibel 1980, Morris et al. 1988, Bathmann et al. 1991). Larvaceans, in particular, form a rich source of marine snow as they periodically abandon their old houses (Alldredge 1972, 1976,

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Davoll & Silver 1986, Alldredge & Silver 1988). These spherical or elliptical houses consist of an outer mucopolysaccharide wall and fine mucous nets (Alldredge 1977) which, in themselves, add a small amount of organic material to the aggregates. However, a variety of particles from the surrounding sea water become attached to the houses. The aggregates contain phytoplankton cells, bacteria, flagellates, ciliates, fecal pellets, mineral grains and other particles in concentrations that exceed bulk water concentrations by orders of magnitude (Taguchi 1982, Davoll & Silver 1986, Alldredge & Silver 1988). Due to the accumulation of living organisms these houses may become sites of elevated microbial activity (Davoll & Silver 1986), and their high particle content makes them suitable as food for the zooplankton (Alldredge 1972, Steinberg et al. 1994). On the other hand, sinking rates exceeding 100 m d^{-1} (Silver & Alldredge 1981, Taguchi 1982, Gorsky et al. 1984) also suggest that larvacean houses increase the export of microorganisms out of the photic zone. However, the role of larvaceans in facilitating the sedimentation of organic carbon and microorganisms is largely unknown.

The accumulation of particles on the house is believed to occur as the larvacean pumps water through the house and the larger particles are screened by the incurrent filters, which gradually clog (Alldredge 1977, Davoll & Silver 1986). In addition to this 'biological' process of aggregation, physical mechanisms such as scavenging and shear coagulation may contribute to the particle enrichment of the house after it has been abandoned. For example, Morris et al. (1988) hypothesised that sinking mucous structures of salps collect significant amounts of particulate material in the water column by scavenging; and, based on theoretical considerations, Stoltzenbach (1993) concluded that scavenging of small particles by fast sinking porous aggregates plays an important role in the vertical particle transport in the ocean. However, the significance of particle scavenging by larvacean houses and other marine snow aggregates is poorly understood. Because larvacean houses are coherent structures of uniform shape they provide a unique simple model with which the significance of particle scavenging can be examined.

This study examines the dynamics and formation mechanisms of marine snow aggregates from abandoned larvacean houses during a spring diatom bloom on the U.S west coast with a view to assessing their role in driving the vertical particle

fluxes. We present data on house production rates, sinking velocities, aggregate flux rates, aggregate composition and laboratory measurements of scavenging rates, and we discuss the mechanisms of aggregate formation. We show that shear coagulation and scavenging are both insignificant processes, and that most particles become attached to the larvacean house due to the filtering activity of its inhabitant

MATERIALS AND METHODS

This field study was conducted from April 13 to 24, 1994, in East Sound, Orcas Island, Washington, at a station (48° 40' 02" N, 122° 54' 01" W) with a bottom depth of 28 m (Fig. 1). A diatom bloom, dominated by *Thalassiosira mendiolana*, *Chaetoceros* spp. and *Thalassionema nitschioides*, developed during the study period, with chlorophyll concentrations increasing to about 20 µg l^{-1} (Kiørboe et al. 1996).

Measurements of plankton abundance. Water samples were collected daily between 08:00 and 09:00 h in 30 l Go-Flo bottles at the sub-surface fluorescence maximum (4 to 8 m) and at 21 m depth (deployment depth of sediment traps). Between April 19 and 21, additional water samples were collected at 18:00 h
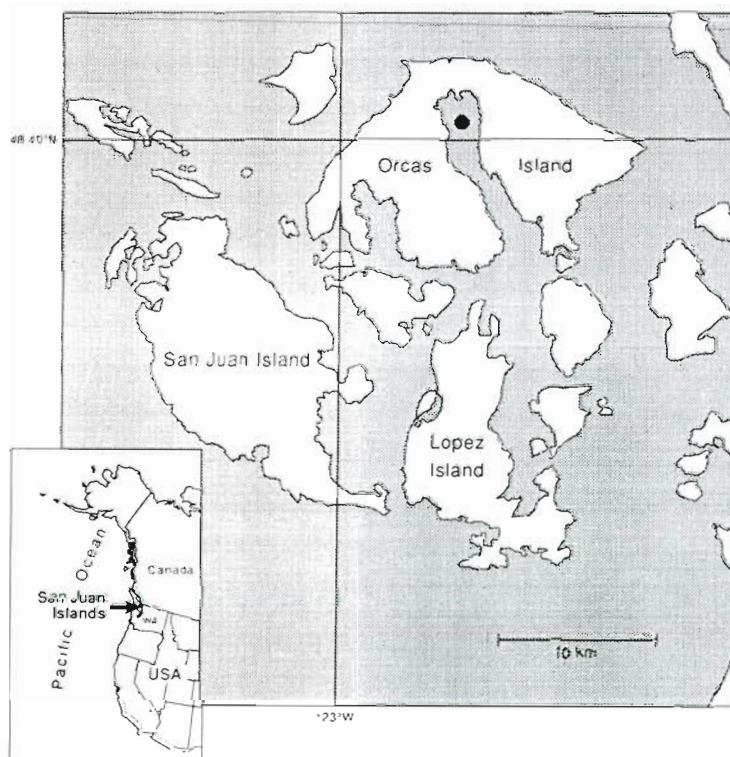

Fig 1 Study site in East Sound, Orcas Island, Washington (48° 40' 02" N, 122° 54' 01" W)

Copepod fecal pellets and phytoplankton were identified and counted as described by Kiørboe et al. (1996). Zooplankton were collected daily between 12:00 and 15:00 h with a 50 cm diameter, 333 μm mesh size net hauled vertically from 20 m to the surface. From April 19 to 24, zooplankton were also collected in 30 l Go-Flo bottles at mean depths of 3, 6, 10, 13, 16 and 20 m. Bottles from the 3 shallowest and 3 deepest depths were drained through a 200 μm mesh and pooled to yield mean abundances above and below 10 m respectively. On April 19 and 20 both types of collection were also repeated at 22:00 h. Larvaceans from all samples were quantified and their trunk lengths measured using microscopy. On April 21 and 23, divers measured the concentration and occupancy rate of visually identifiable larvacean houses passing through a 17.2 cm diameter wire hoop attached to a hand held General Oceanic model 2030 flow meter. Fifty houses were counted on each of 3 transects at 5, 10 and 20 m depths. Occupied houses were easily discernable due to the beating of the animal's tail.

Sampling of larvacean houses. From April 19 to 24, divers collected larvacean houses in syringes at 5, 10 and 20 m depths in batches of 10 to 100 houses per sample as described in Alldredge (1992). Sample volumes were measured, phytoplankton and fecal pellets were counted and numbers of fecal pellets and phytoplankton cells attached to the houses were calculated by subtracting ambient background concentrations measured at 5 and 21 m depth (Kiørboe et al. 1996). Additional larvacean houses were collected and kept unpreserved for 1 to 2 d for measurements of sinking velocity and scavenging efficiency (see below). On April 22, 20 live larvaceans of the species *Oikopleura dioica* were collected at the nearby Friday Harbor Laboratory pier and kept individually in 500 ml beakers containing water from 5 m in East Sound at 12 °C. The number of houses they abandoned over two 12 h periods was monitored. Animals were transferred to fresh sea water at the end of the first period.

Sinking rates. Sinking velocities of larvacean houses collected by divers were measured in 100 ml graduated cylinders filled with filtered seawater from East Sound; about 10 houses were released individually by pipette 2 cm below the surface and sinking times between 6 marks were noted. Only terminal velocities were used. Sinking velocity was also measured for houses being aged in the laboratory (see below).

Scavenging. The potential of settling larvacean houses to scavenge particles from the water column was examined experimentally: larvacean houses were collected in the field on April 19 and incubated in water from the station at the depth of fluorescence maximum (about 5 m) on a rolling table (12 rpm, 2 l bottles with 120 mm radius) or in a standing jar (con-

trol). Houses were sampled at the start, after 12 h and again after 21 h. The attached particles (fecal pellets, ciliates and different phytoplankton species) were enumerated under the inverted microscope. On each sampling occasion the sinking rate of houses was measured as above. We also measured the scavenging efficiency of 1 clean, 1.5 mm larvacean house produced by an *Oikopleura dioica* which had been kept in filtered sea water in the laboratory. The house contained 5 larvacean fecal pellets but was otherwise free of particles. A different procedure to that for the other houses was used: the house was released in an 80 cm transparent tube containing sub-surface sea water from the study site. The tube was closed and then turned upside down every time the house reached the bottom. After 1.8 h the house was collected together with 0.5 ml of water with a wide-mouthed automatic pipette, and the number of *Thalassiosira mendiolana* cells was counted. Four additional 0.5 ml samples were collected for counting of background concentrations of *T. mendiolana*. The sinking velocity was calculated from the number of turns.

Sediment traps. Sediment traps were deployed at 21 m depth. Each individual trap tube measured 65 mm in inner diameter, 675 mm in total height and had a detachable bottom section 85 mm in height. The top 90 mm consisted of 7 baffle tubes, each measuring 17.5 mm in inner diameter (21.5 mm in outer diameter). A varying number of baffle tubes were closed by stoppers to reduce the amount of material in the trap; from the area occluded by the stoppers we assumed that each stopped tube caused a 9 % reduction of the vertical flux. Traps were retrieved every 12 or 24 h.

Two trap tubes were each fitted with a petri dish (60 mm diameter) containing an 8 % polyacrylicamid solution. This highly viscous polymer collects, preserves and retains the 3-dimensional structure of the sedimented material (Lundsgaard 1995). Thus, sedimenting aggregates remain intact. Upon retrieval, most water was siphoned from the tubes, the bottom section was detached, and remaining water was siphoned off using a micro-pipette. Material caught in the polymer was photographed with a Nicon camera equipped with a macro lens (2 to 20× magnification) using films for color slides. At 5× magnification we were able to obtain photographs covering the whole petri dish area. Later, the total number of aggregates was counted from projections of these slides. Aggregates larger than 0.1 mm in diameter were clearly distinguishable. In each sample the volumes of 35 to 100 random aggregates were calculated using close up photographs (20× magnification); from the projected aggregate area we measured 2 perpendicular radii and calculated the volume by assuming an ellipsoid shape for each aggregate. On April 23, larvacean

Table 1 *Oikopleura dioica*. Average abundance between 0 and 20 m and mean trunk length ± SD of animals as determined from bottle casts and net hauls with a net of 330 μm mesh size. The concentrations and occupancy rates of larvacean houses are based on visual observations of indentifiable houses by divers in East Sound. nd = not determined

Date (April)	Abundance m⁻³ Net hauls	Bottle cast	% above 10 m	Trunk length (μm) net hauls	Depth (m)	Houses m⁻³	% Occupied
15	81	30	88	660 ± 160		nd	nd
16	49	–	–	720 ± 150		nd	nd
17	60	–	–	640 ± 170		nd	nd
18	60	–	–	740 ± 180		nd	nd
19	50	191	81	690 ± 130		nd	nd
19.5	117	–	–	660 ± 160		nd	nd
20	38	86	86	630 ± 130		nd	nd
20.5	64	93	93	650 ± 150		nd	nd
21	85	160	96	670 ± 170	5	100 ± 20	63 ± 3
					10	500 ± 200	28 ± 4
					20	600 ± 100	nd
23	80	99	88	670 ± 180	5	800 ± 100	nd
					10	3700 ± 900	nd
					20	4800 ± 900	nd
24	28	80	77	760 ± 140		nd	nd
Mean	65 ± 24	118 ± 42	87 ± 6	680 ± 40			

houses were collected by divers and allowed to settle in a petri dish filled with polymer to be compared with the aggregates from the sediment traps. After approximately 2 h of settling, the size distribution was measured as above. Daily fluxes of larvacean houses were calculated from the number of aggregates >0.3 μl, i.e. aggregates of sizes equivalent to those of the diver-collected houses (see below). Daily aggregate volume fluxes were calculated from the average aggregate size of each sample.

From each petri dish, 2 to 8 aggregates were sampled individually from the polymer with a specially designed pipette mounted on a syringe. Formalin was added to the aggregate samples, which contained 10 to 20 μl polymer, for a final concentration of 2%; samples were then spread on a microscope slide and sealed under a cover slide with fingernail polish. Later, the contents of the aggregates (selected phytoplankton species and fecal pellets) were counted at 200× magnification.

RESULTS

Abundance of larvaceans

The larvacean *Oikopleura dioica* was abundant throughout the study period and occurred in more or less constant densities. Vertical net hauls showed an average water column concentration of 65 larvaceans m⁻³ (Table 1). Sampling with water bottles indicated that net hauls underestimated the densities of larvaceans, because small individuals passed through the 333 μm mesh size net, and also revealed that 87% of the larvaceans occurred in the upper 10 m. The sizes of the net-captured larvaceans did not vary significantly; mean trunk length was 680 μm (Table 1) while the average bottle-collected larvaceans were slightly smaller, 595 ± 63 μm. Abandoned houses of *O. dioica* were always present in the water column. Divers found an average water column concentration of 383 houses m⁻³ on April 21 and 2978 houses m⁻³ on April 23, and the density generally increased with depth. The occupancy rate was 68% at 5 m depth and 28% at 10 m depth on April 21 (Table 1).

Size distribution and flux of larvacean houses

The sediment traps collected aggregates throughout the study period. Microscopical examination of aggregates trapped in the polymer revealed that the majority of settled aggregates were abandoned larvacean houses, or fragments of houses, containing amorphous detritus, fecal pellets and phytoplankton cells. Some aggregates could only be recognized as detritus but their shape was very similar to that of larvacean houses collected by divers and subsequently allowed to settle in the polymer. Larvacean houses collected by divers covered a size range of 0.8 to 1.9 mm equivalent spherical diameter (ESD) or 0.3 to 3.5 μl. The size distribution (volume) of diver-collected houses conformed with

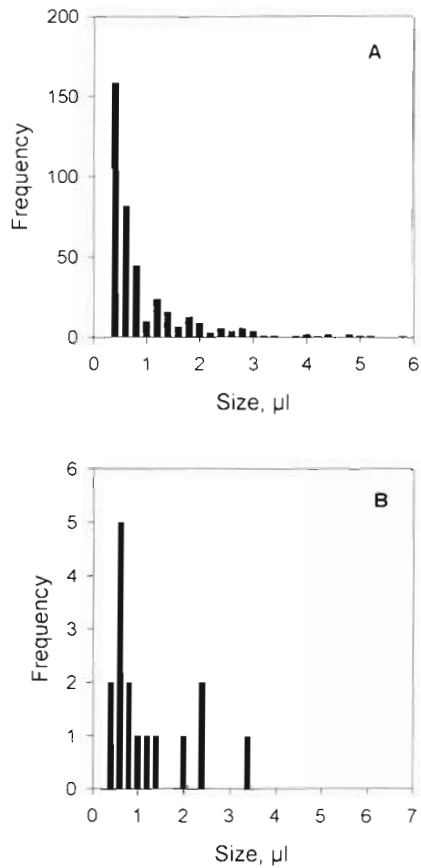Fig. 3. Daily flux of abandoned *Oikopleura dioica* houses (= aggregates larger than 0.3 µl)

Fig. 2. Frequency distribution of aggregates and *Oikopleura dioica* houses obtained from image analysis: (A) aggregates larger than 0.3 µl collected in sediment traps from April 16 to 24, (n = 314), (B) Larvacean houses collected from the water column by divers and subsequently allowed to settle in polymer (n = 16)

the size distribution of all settled aggregates (April 14 to 24) larger than 0.3 µl (Fig. 2) (*t*-test, p > 90%). The average size of larvacean houses was 1.21 mm ESD and average size of aggregates >0.3 µl collected in traps was 1.13 mm ESD. Thus, we considered all aggregates >0.3 µl to be larvacean houses.

The flux of houses was on average 55000 houses m^{-2} d^{-1}, with a maximum value of 116000 houses m^{-2} d^{-1} (Fig. 3). These numbers correspond to average and peak aggregate volume fluxes of 48 ml m^{-2} d^{-1} and 128 ml m^{-2} d^{-1}.

Sinking rates

The grand mean of all sinking velocities measured in the laboratory was 121 m d^{-1} (Table 2). Diver-collected larvacean houses settled at 137 ± 50 m d^{-1} (n = 12). Houses aged for 21 h on the rolling table settled at 69 ± 35 m d^{-1} and the control group at 138 ± 51 m d^{-1}. One clean, 1.5 mm diameter house produced in the labora-

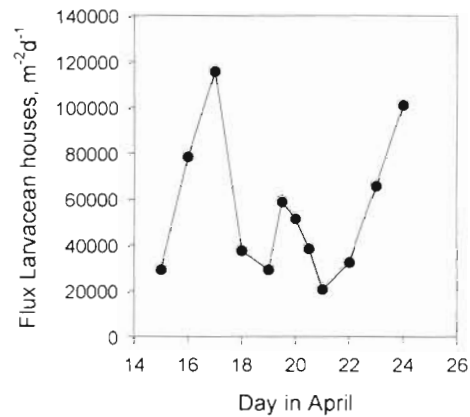tory settled at 115 m d^{-1}. Sinking velocities did not correlate with numbers of attached fecal pellets and diatoms.

House production rate

Oikopleura dioica incubated in the laboratory at 12°C in sea water from East Sound produced 7.2 ± 1.6 houses d^{-1}.

Particle content of larvacean houses

Aggregates in traps

Attached particles could easily be specified and counted on aggregates sampled in the polymer. The aggregates contained high numbers of diatoms, flagellates, ciliates and fecal pellets. The content of *Thalassiosira mendiolana* per aggregate increased from <50 in the beginning of the study to >250 at the end. The abundance of *Thalassionema nitschioides* also increased 5 times during the period to about 600 cells per aggregate. In

Table 2. Sinking velocities of larvacean houses measured in a graduated cylinder or in an 80 cm long transparent tube. Number of houses measured shown in parentheses

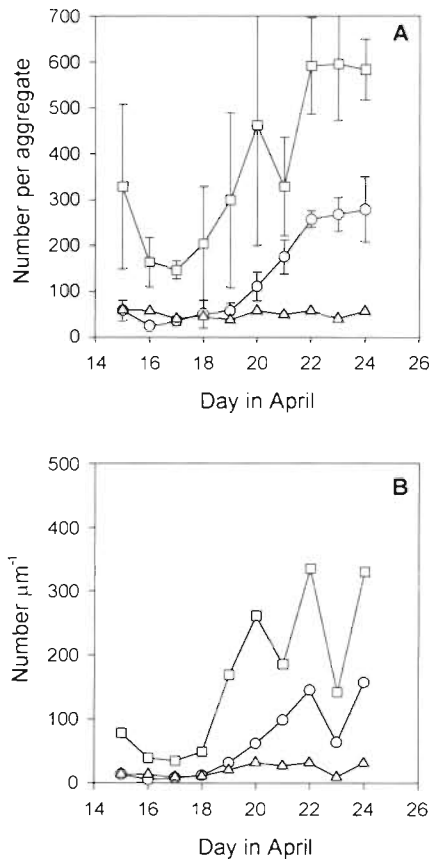| Description of houses	Sinking rate m d^{-1} ± SD
Houses collected *in situ* (12)	137 ± 50
Houses aged for 21 h on rolling table (7)	69 ± 35
Houses in standing jars (11)	138 ± 51
House produced in the laboratory (1)	115
Mean	121

Fig. 4. Contents of diatoms and fecal pellets in aggregates collected in sediment traps. (A) Number per aggregate (n = 2 to 6 per sampling date), (B) number per aggregate volume. (□) *Thalassionema nitschioides*, (O) *Thalassiosira mendiola*, (△) fecal pellets
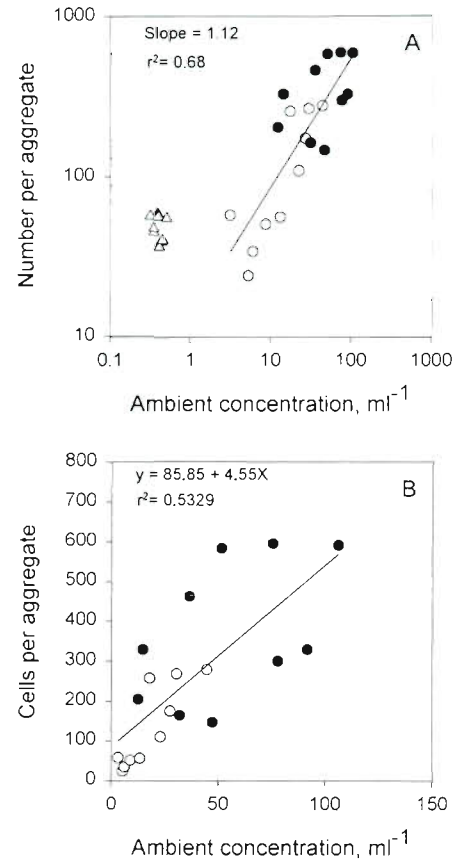

Fig. 5. Trap-collected aggregates: (A) number of particles per aggregate versus water column concentration (mean of concentration at ca 5 and 21 m depth), linear regression (n = 19); (B) as (A) but on a linear scale and without fecal pellets. (●) *Thalassionema nitschioides*, (O) *Thalassiosira mendiola*, (△) fecal pellets

contrast, the content of fecal pellets remained constant at about 50 pellets per aggregate (Fig. 4A). The number of particles per aggregate volume followed the same pattern (Fig. 4B). Increasing numbers of diatoms (*T. mendiolana*, *T. nitschioides*) were closely related to increasing concentrations of these species in the water column, where a diatom bloom was developing. The relationship between bulk water concentration and diatom content of aggregates was given by a first order function (Fig. 5A). On average for the whole study period each aggregate contained these diatoms in abundances corresponding to those found in 4.55 ml of ambient water (Fig. 5B). Inside the aggregates, diatom concentrations were 1000 times higher than in ambient water. Aggregates were even more enriched in fecal pellets, with concentrations 100 000 times higher than in the surrounding water, corresponding to a 'cleared volume' of 100 ml for each aggregate, which refers to the volume of ambient water containing the same number of particles as 1 aggregate.

Diver-collected aggregates

Suspended larvacean houses carried particle assemblages similar to those carried by houses collected in the traps. A large number of diatom species, fecal pellets and ciliates were found on the houses (Table 3). There was no consistent temporal pattern in aggregate enrichment from April 19 to 24 (Table 4). Larvacean houses sampled at 10 m depth seemed to be more enriched than those from 5 and 20 m but the background concentration was obtained from interpolation of concentrations at 5 and 21 m and thus the pattern was not conclusive. The number of attached diatoms averaged 2140 ± 1086 cells house^{-1}, equal to the content in 4.47 ml of ambient water; this is similar to the above estimate (4.55 ml house^{-1}) based on trap-collected houses. Total content of diatoms in houses followed the water column concentration; a plot (Fig. 6) based on abundances of individual diatom species yields a significant relation (linear regression: n = 212,

Table 3. Average number of diatoms, flagellates, ciliates and fecal pellets attached to larvacean houses collected from 5 to 20 m by divers between April 19 and 24. Total sample size is 14, each containing 10 to 100 houses [n (total) = 316]. Data from Kiørboe et al. (1996)

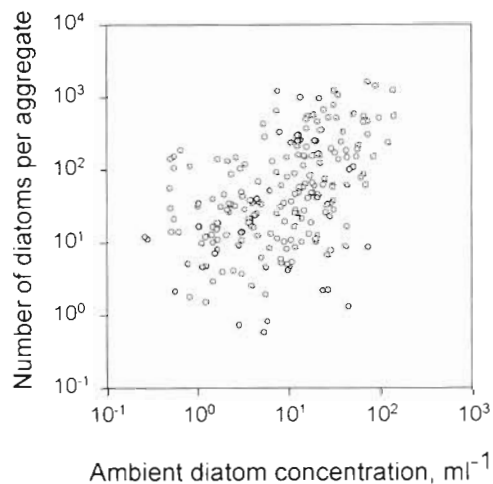| Type of particle	Number of species	Cells per house
Diatoms		
Chaetoceros sp.	10	63
Ditylum brightwellii	–	6.3
Laudaria annulata	–	28
Leptocylindrus danicus	–	32
Navicula sp.	1	15
Nitzschia sp.	2	312
Pseudonitzschia sp.	2	81
Rhizosolenia sp.	6	296
Skeletonema costatum	–	15
Thalassionema nitschioides	–	800
Thalassiosira sp.	4	474
Dictyocha speculum	–	23
Phaeocystis pouchetii	–	1211
Fecal pellets	–	42
Oligotrich ciliates	–	12

Fig. 6. Diver-collected aggregates: number of diatoms per aggregate versus concentration in ambient water. Each data point represents 1 diatom species on 1 sampling day

$t = 3.34$, $p < 0.1\%$) in spite of a considerable scatter ($r^2 = 0.053$), and the slope of the relation, 5.32 ± 0.68 ml house^{-1}, is again similar to the above estimates. Again houses were more enriched with fecal pellets than with diatoms, with an average of 47 ± 56 pellets per house corresponding to the fecal pellet content of 42 ± 43 ml of ambient water; this is of a similar order of magnitude as the 100 ml house^{-1} estimated for the trap samples.

The considerable variation in the numbers of attached particles in diver-collected houses may be caused by differences in their age. For example, some houses were still occupied when collected, while others obviously had been abandoned for some time.

Particle scavenging by settling larvacean houses

Larvacean houses incubated on the rolling table contained more of all groups of particles than did the controls (Table 5). Thus, larvacean houses do scavenge

Table 4. Amounts of fecal pellets and diatoms in larvacean houses collected at various depths. 'Volume cleared' is the calculated volume of ambient water containing the same number of particles as 1 larvacean house. Estimates of volume cleared at 10 m depth were based on the average of ambient concentrations at 5 and 21 m. Data on ambient concentrations of fecal pellets and diatoms are from Kiørboe et al. (1996)

Date (April)	Depth (m)	Diatoms house^{-1}	Volume cleared (diatoms, ml house^{-1})	Pellets house^{-1}	Volume cleared (pellets, ml house^{-1})
19	5	2351	4.45	176	85.1
	10	2378	3.66	21	3.6
	20	2887	3.74	13	5.8
20	5	2624	6.28	25	31.8
	10	3855	9.26	17	59.8
	20	2257	5.44	28	–
21	5	1424	2.56	19	16.2
	10	4106	8.81	190	138.0
	20	1701	4.51	16	22.8
23	5	1149	1.70	38	11.5
	10	1182	2.64	37	9.2
	20	398	1.81	14	3.9
24	5	603	0.8	36	38.9
	10	3042	7.0	26	115.0
Mean ± SD		2140 ± 1086	4.47 ± 2.5	46.7 ± 56.3	41.7 ± 43.1

Table 5. Particle scavenging by sinking larvacean houses. Diver-collected larvacean houses from 10 m (n = 20) incubated in water from 5 m on a rolling table. Another 20 houses were incubated in a standing jar (control). Aggregates were sampled after 12 and 21 h and attached particles were counted and identified. Table shows average number per aggregate and number of houses sampled

| | After 12 h | | After 21h | | Conc. in ambient water |
	Rolling table	Control	Rolling table	Control	(ml^{-1})
Number of houses	1	8	5	6	–
Fecal pellets	16.1	3.9 ± 2.0	25 ± 23	9.3 ± 2.9	0.44
Ciliates	10.0	2.2 ± 1.0	23 ± 20	4.0 ± 7.5	–
Thalassionema sp. (chains)	42.6	8.7 ± 4.4	52 ± 34	9.7 ± 6.5	–
Chaetoceros sp. (chains)	40.9	16.2 ± 14	–	–	–
Thalassiosira sp. (chains)	9.8	5.5 ± 4.3	–	–	–
Phaeocystis (colonies)	2.4	1.5 ± 1.9	–	–	–
Thalassiosira mendiolana	14.1	06.7 ± 13.7	32 ± 17	12.5 ± 7.0	12.3

particles as they settle. We quantified scavenging as a clearance rate, i.e. as the volume of ambient water cleared for particles per unit time = (difference in particles per house in the control and the experimental container)/incubation time/ambient concentration. From Table 5 we estimate that *Thalassiosira mendiolana* cells were cleared at a rate of 0.06 ± 0.01 ml house^{-1} h^{-1} while fecal pellets were cleared at a rate of 2.0 ± 0.4 ml house^{-1} h^{-1} in the rolling table. The larvacean house settling for 1.8 h in the reverting tube at an ambient concentration of 58 ± 14 *T. mendiolana* cells ml^{-1} collected 4 cells; this yields a clearance rate of 0.06 ± 0.08 ml h^{-1}, consistent with the above.

The relatively higher uptake rate of fecal pellets than diatoms in the experiments agrees well with observations of larvacean houses from both the water column and the sediment traps. However, scavenging rates obtained in the laboratory cannot explain the observed particle content of the houses; if larvacean houses settle at 100 m d^{-1} they will on average spend 2 to 3 h in the water column above the trap depth after being abandoned. Thus, expected diatom and fecal pellet abundances in houses due to this mechanism would correspond to the contents in 0.15 and 5 ml of ambient water, respectively. This is about 20 times less than observed.

DISCUSSION

The fragile nature of marine aggregates normally causes them to disintegrate in sediment trap samples. In this study, the polymer in the traps enabled us to demonstrate, for the first time, that marine aggregates found suspended in the water column were similar to aggregates simultaneously collected in sediment traps. Not only were the size distributions similar (Fig. 2), but the quantities of diatoms and fecal pellets found in aggregates were almost identical between those col-

lected in the water column and those collected in the traps. In the present case most aggregates turned out to be discarded larvacean houses. Dense populations of the larvacean *Oikopleura dioica* have earlier been observed to cause high concentrations of suspended aggregates (Alldredge 1979), and there are several previous observations that the gelatinous structures of the house form the matrices of aggregates carrying numerous phytoplankton cells, ciliates, fecal pellets and amorphous detritus (Alldredge 1972, 1976, Taguchi 1982, Davoll & Silver 1986). Alldredge & Silver (1988) concluded that whenever larvaceans are present they form an important source of marine snow. In East Sound we observed that the fate of these aggregates was sedimentation, as they totally dominated the flux of aggregates.

Dynamics and settling of discarded larvacean houses

The net hauls probably undersampled the smaller *Oikopleura dioica* and, consequently, the population estimates based on net hauls are smaller than those estimated by water bottles or divers (Table 1) and do not compare with house fluxes, as observed in the sediment traps. On April 21, divers counted 300 to 400 houses m^{-3} above the depth of the traps. Between 16 and 31% of these houses were occupied (Table 1; assuming an occupation ratio between 0 and 28% at 20 m depth), meaning that the abundance of abandoned houses was about 300 houses m^{-3}. That particular day the flux of houses was 27 000 m^{-2} d^{-1} (average for April 21 and 22). At steady state, these numbers suggest an average *in situ* sinking velocity of (27 000 houses m^{-2} d^{-1}/300 houses m^{-3} =) 90 m d^{-1}. A similar calculation for April 23 and 24 yields 41 m d^{-1} (assuming an occupation ratio of 31%). These numbers agree reasonably well with our laboratory measurements (69 to 138 m d^{-1}) and with Gorsky et al. (1984), who

found settling velocities of 26 to 157 m d^{-1} of *O. dioica* houses. These sinking rates imply that, after being abandoned, the houses are exported out of the euphotic zone within 2 to 3 h and the standing stock in the water column turns over 4 to 5 times d^{-1} Because the houses turn over rapidly in the water column we can calculate the average *in situ* house production rate from house flux divided by the population size of animals above the trap depth; i.e. 27 000 houses m^{-2} d^{-1}/160 animals m^{-3}/21 m = 8 houses animal^{-1} d^{-1}. The same calculation gives a house production rate of 25.7 houses animal^{-1} d^{-1} on April 23. Although a bit high, these values are comparable to our laboratory measurement of 7.2 houses animal^{-1} d^{-1}. Literature values of house production rates range from a few houses per day to >16 houses animal^{-1} d^{-1} (Gorsky et al. 1984, Fenaux 1985). At a comparable temperature (15°C) Fenaux (1985) found a production rate of 7.5 houses animal^{-1} d^{-1} in *O. dioica*.

Mechanisms of aggregation

Our data consistently suggest that: (1) composition and abundance of particles attached to larvacean houses are related to ambient particle composition and abundance, (2) diatom abundance in the aggregates corresponds to a 'cleared' ambient volume of 4 to 5 ml while (3) fecal pellet abundance in aggregates corresponds to 50 to 100 ml of ambient water. There are a number of mechanisms by which suspended particles can become attached to larvacean houses. The larvacean itself produces fecal pellets while inhabiting the house and these remain inside the discarded house (Taguchi 1982). Larvacean pellets resemble copepod fecal pellets, and could not be distinguished. In fact, the number of fecal pellets per house found here (ca 50) is similar to the number of pellets left in discarded houses by *Oikopleura longicauda* (Taguchi 1982), and are most likely due to the animal itself. For the rest of the attached particles there are several possible mechanisms involved in bringing them into physical contact with the house. Obviously, self-motile cells can actively colonize the houses and may explain the association of ciliates with the houses. Large diatoms may be captured on the coarse incurrent filter while the house is still inhabited (Paffenhöffer 1973, Alldredge 1977, Davoll & Silver 1986) thereby forming an aggregate prior to abandonment. In fact, clogging of filters has been invoked as the cause of house renewal (Alldredge 1977). After being discarded, these relatively fast settling houses may scavenge suspended particles on their way through the water column as hypothesised by Morris et al. (1988). Finally abandoned houses may

be brought into contact with suspended particles by shear in the water column.

The efficiency of shear coagulation can be examined theoretically. Assume that an abandoned house can be considered a solid sphere. Then a house collides with suspended particles occurring at concentration C, according to Hill (1992):

$$\text{Collision rate} = \beta_{\text{Shear}} E_{\text{Shear}} C$$

where β_{Shear} is the collision kernel for shear coagulation:

$$\beta_{\text{Shear}} = 0.163 \gamma (d_i + d_j)^3$$

where γ is the shear rate; and E_{Shear} is the contact efficiency of unlike sized solid particles:

$$E_{\text{Shear}} = 7.5 (d_i/d_j)^2 (1 + 2 d_i/d_j)^{-2} \qquad \text{for } d_i < d_j,$$

where d_j is the diameter of the house and d_i is the diameter of the suspended particles. The volume cleared (V) by each house is:

$$V = E_{\text{Shear}} \beta_{\text{Shear}} t \tag{1}$$

where t is the time needed for each house to travel through the water column. The mean vertical current shear rate in East Sound ranged from about 0.01 s^{-1} to slightly more than 0.02 s^{-1} during the study period, (Kiørboe et al. 1996). Since there was hardly any wind during the study period, wind generated turbulence would be insignificant. Assuming $\gamma = 0.02$ s^{-1}, then a 1 mm house sinking 100 m d^{-1} in a suspension of 100 µm diatoms will theoretically clear 0.004 ml of ambient water before reaching the sediment trap. Even if we ignore the hydrodynamic effect (i.e. set $E_{\text{Shear}} = 1$), which is equivalent to assuming that the house is infinitely permeable (Stoltzenbach 1993), then the estimated volume cleared would increase to 0.08 ml. This is small compared to the 4 to 5 ml actually observed and we conclude that shear coagulation contributes insignificantly to aggregate enlargement.

The efficiency of scavenging can be examined in a similar way by replacing the expressions for β and E in Eq. (1) as follows (Hill 1992):

$$V = \beta_{\text{Settling}} E_{\text{Settling}} t \tag{2}$$

where β_{Settling} is the coagulation kernel for differential settling of particles having settling velocities v_i for the suspended particles and v_j for the house and E_{Settling} is the contact efficiency of solid settling particles:

$$\beta_{\text{Settling}} = (\pi/4)(d_i + d_j)^2 |v_i - v_j|$$
$$E_{\text{Settling}} = (d_i/d_j)^2/2[1 + (d_i/d_j)]^2 \qquad \text{for } d_i < d_j$$

Eq. (2) then reduces to:

$$V = (\pi/8) d_i^2 |v_i - v_j| t$$

If the large particle sinks much faster than the small one ($v_i \ll v_j$) then $|v_i - v_j| \approx v_j$. Because $v_j \times t$ is the depth

(z) of the water column, the cleared volume per house is calculated from:

$$V = (\pi/8)\, d_i{}^2 z \qquad \text{(for solid particles)} \qquad (3)$$
$$V = (\pi/4)(d_i + d_j)^2 z \quad \text{(for permeable particles, } E = 1) \quad (4)$$

Eq. (3) shows that the scavenging efficiency does not depend on the size or settling velocity of the sinking particle (i.e. the larvacean house), whereas the size of the smaller particles is critical. For example, a larvacean house sinking a distance of 10 m through a suspension of 100 µm diatoms (e.g. *Thalassiosira mendiola*) would clear 0.04 ml of ambient water before reaching the sediment trap. This is about 2 orders of magnitude less than the 4 to 5 ml house⁻¹ estimated for field-collected houses but similar to the scavenging rates actually observed in the laboratory experiments (Table 6). By ignoring hydrodynamic effects ($E = 1$) this estimate increases to 9.5 ml house⁻¹ for a 1 mm house. However, our scavenging experiments yielded rates which suggest that settling houses behave more like solid than permeable particles (Table 6). The experiments also showed that the larger fecal pellets were scavenged about 30 times faster than diatoms, which is predicted from Eq. (3) if the fecal pellets are about 5 times larger than the diatoms. In contrast, a permeable house would scavenge diatoms and fecal pellets at about the same rate as long as the house is much larger than the particles being scavenged (Eq. 4). These findings are consistent with the non-porous mucous membrane surrounding the house. Trap-collected houses did not carry more particles than those collected in the water column, and particle loading did not increase with collection depth as expected if scavenging was important. Thus, observations, experiments, and theory suggest particle scavenging to be of secondary importance.

By filtering alone, each animal should filter about 35 ml d⁻¹ in order to fill 7.2 houses with the observed number of diatoms. According to Paffenhöfer (1975) an animal with a trunk length of 680 µm filters about 65 ml d⁻¹ at 13 °C (the surface temperature at the study site was 11 to 12 °C). The length-grazing rate regression of *Oikopleura dioica* from King et al. (1980) predicts a filtering rate of 21 ml d⁻¹ at 13.5°C, and according to Alldredge (1981) individuals of this size filter about 121 ml d⁻¹ at 23 °C. Thus, filtering rates reported in the literature appear sufficient to account for the observed abundance of diatoms attached to discarded larvacean houses.

We conclude that the feeding activity of *Oikopleura dioica* itself is the most important mechanism for aggregation of particles onto the houses. Once these impermeable houses become abandoned further collection of particles by scavenging and shear coagulation is insignificant, and their presence in the water column has no quantitative effect on the overall aggregation processes. In fact, the daily flux of houses in East Sound would theoretically scavenge less than 0.1 % of the 100 µm diatom population per day and only 0.001 % of the 10 µm size fraction.

By comparison, if the aggregate flux in East Sound had been composed of diatom aggregates, which are known to be highly permeable (Logan & Alldredge 1989), the sinking aggregates would on average scavenge 5.2 % d⁻¹ [average flux multiplied by cleared volume per aggregate (Eq. 4): 0.95 ml house⁻¹ m⁻¹ × 55 000 houses m⁻² d⁻¹ = 52300 ml m⁻³] of the particles in the entire water column including the smallest size fraction. This last example serves to highlight the general importance of the structure (permeability) of marine snow and, thus, of the origin of the marine snow particles, especially in a deep water column.

Significance of larvacean houses in the pelagic food web

Marine snow particles are recognized as sites of elevated microbial activity (Alldredge & Cox 1982, Alldredge & Silver 1988), and Davoll & Silver (1986) described how the developing microbial community on a discarded larvacean house causes its disintegration within about a week. However, in shallow waters like

Table 6. Volumes cleared of diatoms by larvacean houses as estimated by different approaches: field observations, experiments, and theoretically due to various coagulation mechanisms. The experimental value (0.06 ml h⁻¹) has been converted to the volume cleared during the expected average residence time of an abandoned larvacean house in the upper 20 m of the water column (3 h). Theoretical values are calculated for a 1 mm diameter larvacean house settling at 100 m d⁻¹ in a suspension of 100 µm diameter diatom cells. The houses are assumed to be either porous or non-porous

Observations/mechanisms	Equation	Cleared volume (ml house⁻¹)
Observed *in situ*		
Houses in traps	–	4.55
Houses in suspension	–	4.47
Experiment		
Scavenging of diatoms	–	0.07
Theoretical		
Scavenging		
Porous	$\beta_{Settling}\, t$	9.50
Non-porous	$\beta_{Settling}\, E_{Settling}\, t$	0.04
Shear coagulation ($\gamma = 0.02$ s⁻¹)		
Porous	$\beta_{Shear}\, t$	0.08
Non-porous	$\beta_{Shear}\, E_{Shear}\, t$	0.004

East Sound, discarded houses reach the sea floor in a few hours, and in such areas the most important role of abandoned larvacean houses appears to be their contribution to the vertical flux of organic material. Since scavenging and shear coagulation are unimportant as mechanisms of aggregation, the house production rate of *Oikopleura dioica* affects only the volume flux of marine snow, whereas the flux of attached particles depends solely on the filtration rate of the larvacean. By its feeding current each larvacean daily collects an amount of particulate material corresponding to the pumping rate multiplied by the ambient concentration of particles. This material is transformed into fast sinking aggregates due to fecal pellet production (Pomeroy & Deibel 1980) and collection on the house. Thus, functionally the entire abandoned house resembles sinking fecal pellets of other zooplankters. However, as the larvaceans consume only a certain size spectrum, they may cause much more organic material to settle undigested from the euphotic zone than they actually consume and this material includes live phytoplankton.

Although the abundance of larvaceans in this study suggests a rather limited effect on the sedimentation rate of the phytoplankton (Kiørboe et al. 1996), the observed flux of live diatoms was mainly due to cells attached to larvacean houses (Kiørboe et al. 1996). Since larvaceans may occur in dense swarms exceeding $10\,000$ m^{-3} (Seki 1973), abandoned larvacean houses may at times account for a significant fractional removal rate of large particles from the euphotic zone, and they appear to be one of the several mechanisms behind the rapid fallout of live phytoplankton sometimes seen during diatom blooms.

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